



National Health Data Dictionary volume 1

Data elements A to C (by short name)
Generated on 05/01/2007

© Australian Institute of Health and Welfare 2006

This work is copyright. Apart from any use as permitted under the *Copyright Act 1968*, no part may be reproduced without prior written permission from the Australian Institute of Health and Welfare. Requests and enquiries concerning reproduction and rights should be directed to the Head, Business Promotion and Media, Australian Institute of Health and Welfare, GPO Box 570, Canberra ACT 2601.

Any enquiries about or comments on this publication should be directed to:

National Data Development and Standards Unit
Australian Institute of Health and Welfare
GPO Box 570
Canberra ACT 2601
Email: datadevelopment@aihw.gov.au
Phone: (02) 6244 1222 Fax: (02) 6244 1166

List of metadata items

Data Elements	10
Accrued mental health care days	11
Activity and participation life area	14
Activity when injured	17
Activity when injured (non-admitted patient)	19
Actual place of birth	21
Acute coronary syndrome procedure type	23
Acute coronary syndrome stratum	25
Additional diagnosis	28
Address line (person)	31
Address line (service provider organisation)	34
Address type (person)	37
Address type (service provider organisation)	39
Address – country identifier (person)	41
Admission date	43
Admission time	46
Admitted patient election status	47
Age	49
Age range	51
Alcohol consumption frequency (self reported)	53
Alcohol consumption in standard drinks per day (self reported)	58
Anaesthesia administered for operative delivery of the baby	61
Analgesia administered for labour	63
Angiotensin converting enzyme (ACE) inhibitors therapy status	65
Anticipated patient election status	67
Apgar score at 1 minute	69
Apgar score at 5 minutes	70
Area of usual residence	72
Aspirin therapy status	75
Assistance with activities	77
Australian State/Territory identifier (establishment)	80
Australian state/territory identifier	83
Australian state/territory identifier (service provider organisation)	85
Behaviour-related risk factor intervention - purpose	87
Behaviour-related risk factor intervention purpose	91
Beta-blocker therapy status	93
Birth order	95
Birth plurality	97
Bleeding episode using TIMI criteria (status)	99
Blindness (diabetes complication)	101
Blood pressure – diastolic (measured)	105
Blood pressure – systolic (measured)	110
Bodily location of main injury	115
Body function	118
Body mass index – adult (measured)	121
Body mass index – adult (self-reported)	125
Body mass index – child (measured)	129
Body mass index – child (self-reported)	132
Body mass index – classification	136

Body structure.....	141
Building/complex sub-unit number (person).....	144
Building/complex sub-unit number (service provider organisation)	146
Building/complex sub-unit type – abbreviation (person).....	147
Building/complex sub-unit type – abbreviation (service provider organisation)	147
Building/property name (person).....	147
Building/property name (service provider organisation).....	147
CVD drug therapy – condition	147
Caesarean section indicator, last previous birth	147
Cancer initial treatment completion date.....	147
Cancer initial treatment starting date	147
Cancer staging – M stage code.....	147
Cancer staging – N stage code	147
Cancer staging – T stage code.....	147
Cancer staging – TNM stage grouping code	147
Cancer treatment type.....	147
Cancer treatment – target site (ICD-10-AM).....	147
Cancer treatment – target site (ICDO-3).....	147
Cardiovascular medication (current).....	147
Care type.....	147
Carer participation arrangements – carer consultants employed.....	147
Carer participation arrangements – carer satisfaction surveys.....	147
Carer participation arrangements – formal complaints mechanism	147
Carer participation arrangements – formal participation policy	147
Carer participation arrangements – regular discussion groups.....	147
Cataract - history	147
Category reassignment date.....	147
Census date	147
Centrelink customer reference number	147
Cerebral stroke due to vascular disease (history)	147
Change to body structure.....	147
Chest pain pattern category	147
Cholesterol – HDL (measured).....	147
Cholesterol – LDL (calculated)	147
Cholesterol – total (measured).....	147
Classification of health labour force job	147
Client type (alcohol and other drug treatment services).....	147
Clinical evidence of chronic lung disease (status)	147
Clinical evidence of heart failure (status).....	147
Clinical evidence of peripheral arterial disease (status).....	147
Clinical evidence of sleep apnoea syndrome (status).....	147
Clinical evidence of stroke (status)	147
Clinical procedure timing (status).....	147
Clinical urgency	147
Clopidogrel therapy status.....	147
Co-location status of mental health service	147
Compensable status	147
Complication of labour and delivery.....	147
Complications of pregnancy	147
Concurrent clinical condition (on presentation)	147

Congenital malformations.....	147
Congenital malformations – BPA code.....	147
Consumer committee representation arrangements	147
Consumer participation arrangements – consumer consultants employed.....	147
Consumer participation arrangements – consumer satisfaction surveys.....	147
Consumer participation arrangements – formal complaints mechanism	147
Consumer participation arrangements – formal participation policy.....	147
Consumer participation arrangements – regular discussion groups	147
Contract establishment identifier	147
Contract procedure flag.....	147
Contract role.....	147
Contract type.....	147
Contracted care commencement date.....	147
Contracted care completion date.....	147
Coronary artery disease – history of intervention or procedure.....	147
Country of birth.....	147
Creatine kinase MB isoenzyme level (index code).....	147
Creatine kinase MB isoenzyme level (international units)	147
Creatine kinase MB isoenzyme level (kCat per litre)	147
Creatine kinase MB isoenzyme level (micrograms per litre).....	147
Creatine kinase MB isoenzyme level (nanograms per decilitre).....	147
Creatine kinase MB isoenzyme level (percentage)	147
Creatine kinase MB isoenzyme – upper limit of normal range (index code).....	147
Creatine kinase MB isoenzyme – upper limit of normal range (international units).....	147
Creatine kinase MB isoenzyme – upper limit of normal range (kCat per litre).....	147
Creatine kinase MB isoenzyme – upper limit of normal range (micrograms per litre).....	147
Creatine kinase MB isoenzyme – upper limit of normal range (nanograms per decilitre)	147
Creatine kinase MB isoenzyme – upper limit of normal range (percentage).....	147
Creatinine serum level (measured)	147

Data Element Technical Names

Adult – body mass index (measured), ratio NN[N].N[N]	121
Adult – body mass index (self-reported), ratio NN[N].N[N]	125
Birth event – anaesthesia administered, code N	61
Birth event – analgesia administered, code N	63
Birth event – birth plurality, code N	97
Birth event – complication, code (ICD-10-AM 5th edn) ANN{.N[N]}	147
Birth event – setting of birth (actual), code N	21
Birth – Apgar score (at 1 minute), code NN	69
Birth – Apgar score (at 5 minutes), code NN	70
Birth – birth order, code N	95
Cancer treatment – cancer treatment type, code N	147
Cancer treatment – non-surgical cancer treatment completion date, DDMMYYYY	147
Cancer treatment – non-surgical cancer treatment start date, DDMMYYYY	147
Cancer treatment – target site for cancer treatment, code (ICD-10-AM 5th edn) ANN{.N[N]}	147
Cancer treatment – target site for cancer treatment, code (ICDO-3) ANN	147
Child – body mass index (measured), ratio NN[N].N[N]	129
Child – body mass index (self-reported), ratio NN[N].N[N]	132
Contracted hospital care – contracted care commencement date, DDMMYYYY	147
Contracted hospital care – contracted care completed date, DDMMYYYY	147
Contracted hospital care – organisation identifier, NNX[X]NNNNN	147
Elective care waiting list episode – category reassignment date, DDMMYYYY	147
Elective surgery waiting list episode – anticipated accommodation status, code N	67
Elective surgery waiting list episode – clinical urgency, code N	147
Episode of admitted patient care – admission date, DDMMYYYY	43
Episode of admitted patient care – admission time, hhmm	46
Episode of admitted patient care – patient election status, code N	47
Episode of care (procedure) – contracted procedure flag, code N	147
Episode of care – additional diagnosis, code (ICD-10-AM 5th edn) ANN{.N[N]}	28
Episode of care – behaviour-related risk factor intervention purpose, code N	87
Episode of care – behaviour-related risk factor intervention, code NN	91
Episode of treatment for alcohol and other drugs – client type, code N	147
Establishment – accrued mental health care days, total N[N(7)]	11
Establishment – Australian state/territory identifier, code N	80
Female – caesarean section indicator (last previous birth) code N	147
Health professional – occupation, code ANN	147
Hospital census (of elective surgery waitlist patients) – census date, DDMMYYYY	147
Hospital service – care type, code N[N].N	147
Hospital – contract role, code A	147
Hospital – contract type, code N	147
Injury event – activity type, code (ICD-10-AM 5th edn) ANNNN	17
Injury event – activity type, non-admitted patient code N[N]	19
Laboratory standard – upper limit of normal range for creatine kinase myocardial band isoenzyme, index code X[XXX]	147
Laboratory standard – upper limit of normal range for creatine kinase myocardial band isoenzyme, total international units N[NNN]	147
Laboratory standard – upper limit of normal range for creatine kinase myocardial band isoenzyme, total kCat per litre N[NNN]	147

Laboratory standard – upper limit of normal range for creatine kinase myocardial band isoenzyme, total micrograms per litre N[NNN]	147
Laboratory standard – upper limit of normal range for creatine kinase myocardial band isoenzyme, total nanograms per decilitre N[NNN]	147
Laboratory standard – upper limit of normal range for creatine kinase myocardial band isoenzyme, percentage N[NNN].....	147
Patient – compensable status, code N.....	147
Person (address) – address line, text [X(180)].....	31
Person (address) – address type, code N	37
Person (address) – building/complex sub-unit identifier, [X(7)]	144
Person (address) – building/complex sub-unit type, code A[AAA]	147
Person (address) – building/property name, text [X(30)]	147
Person (address) – country identifier, code (SACC 1998) NNNN.....	41
Person with cancer – distant metastasis status, M stage (UICC TNM Classification of Malignant Tumours 5th ed) code XX.....	147
Person with cancer – extent of primary cancer, TNM stage (UICC TNM Classification of Malignant Tumours 5th ed) code XXXX{[X]XX}	147
Person with cancer – primary tumour status, T stage (UICC TNM Classification of Malignant Tumours 5th ed) code XX[X].....	147
Person with cancer – regional lymph node metastasis status, N stage (UICC TNM Classification of Malignant Tumours 5th ed) code XX.....	147
Person – activity and participation life area, code (ICF 2001) AN[NNN]	14
Person – acute coronary syndrome concurrent clinical condition, code NN.....	147
Person – acute coronary syndrome procedure type, code NN	23
Person – acute coronary syndrome risk stratum, code N	25
Person – age range, code NN.....	51
Person – age, total years N[NN].....	49
Person – alcohol consumption amount (self-reported), total standard drinks NN.....	58
Person – alcohol consumption frequency (self-reported), code NN	53
Person – angiotensin converting enzyme inhibitors therapy status, code NN.....	65
Person – area of usual residence, geographical location code (ASGC 2006) NNNNN.....	72
Person – aspirin therapy status, code NN.....	75
Person – Australian state/territory identifier, code N	83
Person – beta-blocker therapy status, code NN	93
Person – bleeding episode status, code N	99
Person – blindness, code N	101
Person – blood pressure (diastolic) (measured), millimetres of mercury NN[N]	105
Person – blood pressure (systolic) (measured), millimetres of mercury NN[N]	110
Person – bodily location of main injury, code NN.....	115
Person – body function, code (ICF 2001) AN[NNNN].....	118
Person – body mass index (classification), code N[.N]	136
Person – body structure, code (ICF 2001) AN[NNNN]	141
Person – cardiovascular disease condition targeted by drug therapy, code NN	147
Person – cardiovascular medication taken (current), code N.....	147
Person – cataract status, code N	147
Person – cerebral stroke due to vascular disease (history), code N.....	147
Person – chest pain pattern, code N.....	147
Person – cholesterol level (measured), total millimoles per litre N[N].N	147
Person – clinical evidence status (chronic lung disease), code N	147
Person – clinical evidence status (heart failure), code N.....	147
Person – clinical evidence status (peripheral arterial disease), code N.....	147

Person—clinical evidence status (sleep apnoea syndrome), code N.....	147
Person—clinical evidence status (stroke), code N	147
Person—clinical procedure timing, code N	147
Person—clopidogrel therapy status, code NN.....	147
Person—congenital malformation, code (BPA 1979) ANN.N[N].....	147
Person—congenital malformation, code (ICD-10-AM 5th edn) ANN{.N[N]}	147
Person—coronary artery disease intervention (history), code N.....	147
Person—country of birth, code (SACC 1998) NNNN	147
Person—creatine kinase myocardial band isoenzyme level (measured), index code X[XXX]	147
Person—creatine kinase myocardial band isoenzyme level (measured), total kCat per litre N[NNN]	147
.....	
Person—creatine kinase myocardial band isoenzyme level (measured), total nanograms per decilitre N[NNN].....	147
Person—creatine kinase myocardial band isoenzyme level (measured), percentage N[NNN].....	147
Person—creatine kinase-myocardial band isoenzyme level (measured), total international units N[NNN].....	147
Person—creatine kinase-myocardial band isoenzyme level (measured), total micrograms per litre N[NNNN].....	147
Person—creatinine serum level, micromoles per litre NN[NN].....	147
Person—government funding identifier, Centrelink customer reference number {N(9)A}.....	147
Person—high-density lipoprotein cholesterol level (measured), total millimoles per litre [N].NN.	147
Person—low-density lipoprotein cholesterol level (calculated), total millimoles per litre N[N].N.	147
Person—nature of impairment of body structure, code (ICF 2001) N	147
Person—need for assistance with activities in a life area, code N	77
Pregnancy (current)—complication, code (ICD-10-AM 5th edn) ANN{.N[N]}	147
Service provider organisation (address)—address line, text [X(180)].....	34
Service provider organisation (address)—address type, code N	39
Service provider organisation (address)—building/complex sub-unit identifier, [X(7)]	146
Service provider organisation (address)—building/complex sub-unit type, code A[AAA]	147
Service provider organisation (address)—building/property name, text [X(30)]	147
Service provider organisation—Australian state/territory identifier, code N	85
Specialised mental health service organisation—carer participation arrangements status (carer consultants employed), code N	147
Specialised mental health service organisation—carer participation arrangements status (carer satisfaction surveys), code N	147
Specialised mental health service organisation—carer participation arrangements status (formal complaints mechanism), code N	147
Specialised mental health service organisation—carer participation arrangements status (formal participation policy), code N	147
Specialised mental health service organisation—carer participation arrangements status (regular discussion groups), code N	147
Specialised mental health service organisation—consumer committee representation arrangements, code N	147
Specialised mental health service organisation—consumer participation arrangements (consumer consultants employed), code N	147
Specialised mental health service organisation—consumer participation arrangements (consumer satisfaction surveys), code N	147
Specialised mental health service organisation—consumer participation arrangements (formal complaints mechanism), code N	147
Specialised mental health service organisation—consumer participation arrangements (formal participation policy), code N	147

Specialised mental health service organisation – consumer participation arrangements (regular discussion groups), code N	147
Specialised mental health service – co-location with acute care hospital, code N	147

Data Elements

Accrued mental health care days

Identifying and definitional attributes

<i>Metadata item type:</i>	Data Element
<i>Technical name:</i>	Establishment – accrued mental health care days, total N[N(7)]
<i>METeOR identifier:</i>	286770
<i>Registration status:</i>	NHIG, Standard 08/12/2004
<i>Definition:</i>	The total number of accrued mental health care days provided by admitted patient care services and residential mental health care services within the reference period (from 1 July to 30 June inclusive).

Data element concept attributes

<i>Data element concept:</i>	Establishment – accrued mental health care days
<i>Definition:</i>	The accrued number of mental health care days provided by admitted patient care services and residential mental health care services within the reference period (from 1 July to 30 June inclusive).
<i>Context:</i>	Specialised mental health services
<i>Object class:</i>	Establishment
<i>Property:</i>	Accrued mental health care days

Value domain attributes

Representational attributes

<i>Representation class:</i>	Total
<i>Data type:</i>	Number
<i>Format:</i>	N[N(7)]
<i>Maximum character length:</i>	8
<i>Unit of measure:</i>	Day

Data element attributes

Collection and usage attributes

<i>Guide for use:</i>	<p>The days to be counted are only those days occurring within the reference period, i.e. from 1 July to the following 30 June for the relevant period, even if the patient/resident was admitted prior to the reference period or discharged after the reference period. A day is measured from midnight to 2359 hours.</p> <p>The following basic rules are used to calculate the number of accrued mental health care days:</p> <ul style="list-style-type: none">• Admission and discharge on the same day is equal to one mental health care day.• For a patient/resident admitted and discharged on different days all days are counted as mental health care days, except the day of discharge and any leave days.• If the patient/resident remains in hospital or residential care facility from midnight to 2359 hours count as a mental health care day.
-----------------------	--

- The day a patient/resident goes on leave is not counted as a mental health care day, unless this was also the admission day.
- The day the patient/resident returns from leave is counted as a mental health care day, unless the patient/resident goes on leave again on the same day of return or is discharged.
- Leave days involving an overnight absence are not counted as mental health care days.
- If a patient/resident goes on leave the day they are admitted and does not return from leave until the day they are discharged, count as one mental health care day.
- If the patient/resident remains in a hospital or residential care facility from 1 July to 30 June (the whole of the reference period) count as 365 days (or 366 days in a leap year).
- If the patient/resident remains in a hospital or residential care facility after the end of the reference period (i.e. after 30 June) do not count any days after the end of the reference period.

The following additional rules cover special circumstances and in such cases, override the basic rules:

When calculating accrued mental health care days for the reference period:

- Count the mental health care days of those patients/residents separated during the reference period. Exclude any days that may have occurred before the beginning of the reference period.
- Count the mental health care days of those patients/residents admitted during the reference period who did not separate until the following reference period. Exclude the days after the end of the reference period.
- For patients/residents admitted before the reference period and who remain in after the reference period (i.e. after 30 June), count the mental health care days within the reference period only. Exclude all days before and after the reference period.

Examples of mental health care day counting for a reference period 1 July 2004 to 30 June 2005:

Patient/resident A was admitted to hospital on 4 June 2004 and separated on 6 July 2004. If no leave or transfer occurred counting starts on 1 July. Count would be 5 days as day of discharge is not counted.

Patient/resident B was admitted to hospital on 1 August 2004 and separated on 8 August 2004. If no leave or transfer occurred counting starts on 1 August. Count would be 7 days as day of discharge is not counted.

Patient/resident C was admitted to hospital on 1 June 2005 and separated on 6 July 2005. If no leave or transfer occurred counting starts on 1 June. Count would be 30 days as patient/resident was not discharged on 30 June, so every day up to and including 30 June would be counted.

Patient/resident D was admitted to hospital on 1 August 2003 and has remained continuously in hospital to the present time. If no leave or transfer occurred counting starts on 1 July 2004

and concludes on 30 June 2005. Count would be 365 days as there is no day of discharge.

Collection methods:

To be reported for admitted patient care services, including services that are staffed for less than 24 hours, and non-government organisation services where included.

NOTE: These data need to be disaggregated by Specialised mental health service setting (excluding Ambulatory care settings). For admitted patient care settings these counts also need to be disaggregated by Specialised mental health service program type and Specialised mental health service target population.

Relational attributes

Implementation in Data Set Specifications:

Mental health establishments NMDS 2005-2006 NHIG, Superseded 07/12/2005

Implementation start date: 01/07/2005

Mental health establishments NMDS 2005-2006 NHIG, Superseded 21/03/2006

Implementation start date: 01/07/2005

Implementation end date: 30/06/2006

Mental health establishments NMDS 2006-2007 NHIG, Superseded 23/10/2006

Implementation start date: 01/07/2006

Implementation end date: 30/06/2007

Mental health establishments NMDS 2007-2008 NHIG, Standard 23/10/2006

Implementation start date: 01/07/2007

Activity and participation life area

Identifying and definitional attributes

<i>Metadata item type:</i>	Data Element
<i>Technical name:</i>	Person – activity and participation life area, code (ICF 2001) AN[NNN]
<i>METeOR identifier:</i>	320125
<i>Registration status:</i>	NHIG, Standard 29/11/2006 NCSIMG, Standard 16/10/2006
<i>Definition:</i>	The life area in which a person participates or undertakes activities, as represented by a code.
<i>Context:</i>	Human functioning and disability

Data element concept attributes

<i>Data element concept:</i>	Person – activity and participation life area
<i>Definition:</i>	The life area in which a person may participate or undertake activities.
<i>Object class:</i>	Person
<i>Property:</i>	Activity and participation life area

Value domain attributes

Representational attributes

<i>Classification scheme:</i>	International Classification of Functioning, Disability and Health 2001
<i>Representation class:</i>	Code
<i>Data type:</i>	String
<i>Format:</i>	AN[NNN]
<i>Maximum character length:</i>	5

Collection and usage attributes

<i>Guide for use:</i>	<p>This metadata item contributes to the definition of the concept 'Disability' and gives an indication of the experience of disability for a person.</p> <p>The activities and participation codes are a neutral list that covers the full range of life areas in which a person can be involved. The domains can be used to record positive or neutral experience of functioning as well as limitations and restrictions. Data can be collected at the three digit level in one chapter and at the chapter level in another. However it is only possible to collect data at a single level of the hierarchy in a single chapter to maintain mutual exclusivity. For example, it is not permitted to collect both 'Self care' (chapter level) and 'Looking after one's health' (3 digit level) as the former includes the latter.</p> <p>The value domain below refers to the highest hierarchical level (ICF chapter level). Data collected at this level, in association with respective qualifiers (Activity difficulty level, Activity Need for assistance, Participation extent and Participation satisfaction level) will use the codes as indicated.</p> <p>CODE d1 Learning and applying knowledge</p>
-----------------------	--

CODE d2 General tasks and demands
CODE d3 Communication
CODE d4 Mobility
CODE d5 Self-care
CODE d6 Domestic life
CODE d7 Interpersonal interactions and relationships
CODE d8 Major life areas
CODE d9 Community, social and civic life

Data collected at this level will provide a general description of functioning for the person and can only be compared with data collected at the same level.

Each chapter contains categories at different levels ordered from general to detailed. For specific more detailed information the user should follow the structure of the ICF; the codes should be drawn from the same hierarchical level within any particular chapter. The full range of permissible values is listed in the **Activities and Participation** component of the ICF.

An example of a value domain at the 3 digit level from the Self-care chapter may include:

CODE d510 Washing oneself
CODE d520 Caring for body parts
CODE d530 Toileting
CODE d540 Dressing
CODE d550 Eating
CODE d560 Drinking
CODE d570 Looking after one's health

An example of value domains at the 4 digit level from the Mobility chapter may include:

CODE d4600 Moving around within the home
CODE d4601 Moving around within buildings other than home
CODE d4602 Moving around outside the home and other buildings
CODE d4701 Using private motorized transportation
CODE d4702 Using public motorized transportation

The prefix *d* denotes the domains within the component of *Activities and Participation*. At the user's discretion, the prefix *d* can be replaced by *a* or *p*, to denote activities or participation respectively.

Source and reference attributes

<i>Submitting organisation:</i>	Australian Institute of Health and Welfare (AIHW) which is the Australian Collaborating Centre for the World Health Organization Family of International Classifications.
<i>Origin:</i>	WHO 2001. ICF: International Classification of Functioning, Disability and Health. Geneva: WHO AIHW 2003. ICF Australian User Guide Version 1.0. Canberra: AIHW
<i>Reference documents:</i>	Further information on the ICF, including more detailed codes, can be found in the ICF itself and the ICF Australian User Guide (AIHW 2003), at the following websites: <ul style="list-style-type: none">• WHO ICF website http://www.who.int/classifications/icf/en/• Australian Collaborating Centre ICF website

Data element attributes

Collection and usage attributes

Guide for use:

This metadata item, in conjunction with *Activity difficulty level code N*, enables the provision of information about the presence and extent of activity limitation for any given life area; with *Activity need for assistance code N*, the provision of information about the need for assistance with the given life area.

The extent of, and level of satisfaction with, participation in a given area are indicated by the use of this metadata item with the qualifiers *Participation extent code N* and *Participation satisfaction level code N*.

Source and reference attributes

Submitting organisation:

Australian Institute of Health and Welfare (AIHW) which is the Australian Collaborating Centre for the World Health Organization Family of International Classifications.

Relational attributes

Implementation in Data Set Specifications:

Activities and Participation cluster NHIG, Standard 29/11/2006
NCSIMG, Standard 16/10/2006

Activity when injured

Identifying and definitional attributes

<i>Metadata item type:</i>	Data Element
<i>Technical name:</i>	Injury event – activity type, code (ICD-10-AM 5th edn) ANNNN
<i>METeOR identifier:</i>	333849
<i>Registration status:</i>	NHIG, Standard 07/12/2005
<i>Definition:</i>	The type of activity being undertaken by the person when injured, for admitted patients, as represented by a code.

Data element concept attributes

<i>Data element concept:</i>	Injury event – activity type
<i>Definition:</i>	The type of activity being undertaken by the person when injured.
<i>Context:</i>	Injury surveillance
<i>Object class:</i>	Injury event
<i>Property:</i>	Activity type

Value domain attributes

Representational attributes

<i>Classification scheme:</i>	International Statistical Classification of Diseases and Related Health Problems, Tenth Revision, Australian Modification 5th edition
<i>Representation class:</i>	Code
<i>Data type:</i>	String
<i>Format:</i>	ANNNN
<i>Maximum character length:</i>	5

Data element attributes

Collection and usage attributes

<i>Guide for use:</i>	Use the appropriate External Causes of Morbidity and Mortality Activity codes from the current edition of ICD-10-AM. Used with ICD-10-AM external cause codes and assigned according to the Australian Coding Standards. External cause codes in the range W00 to Y34, except Y06 and Y07 must be accompanied by a place of occurrence code. External cause codes V01 to Y34 must be accompanied by an activity code.
<i>Comments:</i>	Enables categorisation of injury and poisoning according to factors important for injury control. Necessary for defining and monitoring injury control targets, injury costing and identifying cases for in-depth research. This term is the basis for identifying work-related and sport-related injuries.

Source and reference attributes

Origin:

National Centre for Classification in Health
National Injury Surveillance Unit

Relational attributes

Related metadata references:

Supersedes Injury event – activity type, code (ICD-10-AM 4th edn) ANNNN NHIG, Superseded 07/12/2005

Implementation in Data Set Specifications:

Admitted patient care NMDS 2007-2008 NHIG, Standard 22/11/2006

Implementation start date: 01/07/2007

Information specific to this data set:

To be used with ICD-10-AM external cause codes.

Effective for collection from 01/07/2007

Injury surveillance DSS NHIG, Standard 03/05/2006

Information specific to this data set:

Effective for collection from 01/07/2006

Injury surveillance NMDS NHIG, Superseded 03/05/2006

Implementation start date: 01/07/2005

Implementation end date: 30/06/2006

Information specific to this data set:

Effective for collection from 01/07/2006

Activity when injured (non-admitted patient)

Identifying and definitional attributes

<i>Metadata item type:</i>	Data Element
<i>Technical name:</i>	Injury event – activity type, non-admitted patient code N[N]
<i>METeOR identifier:</i>	268942
<i>Registration status:</i>	NHIG, Standard 01/03/2005
<i>Definition:</i>	The type of activity undertaken by the non-admitted patient when injured, as represented by a code.

Data element concept attributes

<i>Data element concept:</i>	Injury event – activity type
<i>Definition:</i>	The type of activity being undertaken by the person when injured.
<i>Context:</i>	Injury surveillance
<i>Object class:</i>	Injury event
<i>Property:</i>	Activity type

Value domain attributes

Representational attributes

<i>Representation class:</i>	Code																																				
<i>Data type:</i>	String																																				
<i>Format:</i>	N[N]																																				
<i>Maximum character length:</i>	2																																				
<i>Permissible values:</i>	<table><thead><tr><th>Value</th><th>Meaning</th></tr></thead><tbody><tr><td>0</td><td>Sports activity</td></tr><tr><td>00</td><td>Football, rugby</td></tr><tr><td>01</td><td>Football, Australian</td></tr><tr><td>02</td><td>Football, soccer</td></tr><tr><td>03</td><td>Hockey</td></tr><tr><td>04</td><td>Squash</td></tr><tr><td>05</td><td>Basketball</td></tr><tr><td>06</td><td>Netball</td></tr><tr><td>07</td><td>Cricket</td></tr><tr><td>08</td><td>Roller blading</td></tr><tr><td>09</td><td>Other and unspecified sporting activity</td></tr><tr><td>1</td><td>Leisure activity (excluding sporting activity)</td></tr><tr><td>2</td><td>Working for income</td></tr><tr><td>3</td><td>Other types of work</td></tr><tr><td>4</td><td>Resting, sleeping, eating or engaging in other vital activities</td></tr><tr><td>5</td><td>Other specified activities</td></tr><tr><td>6</td><td>Unspecified activities</td></tr></tbody></table>	Value	Meaning	0	Sports activity	00	Football, rugby	01	Football, Australian	02	Football, soccer	03	Hockey	04	Squash	05	Basketball	06	Netball	07	Cricket	08	Roller blading	09	Other and unspecified sporting activity	1	Leisure activity (excluding sporting activity)	2	Working for income	3	Other types of work	4	Resting, sleeping, eating or engaging in other vital activities	5	Other specified activities	6	Unspecified activities
Value	Meaning																																				
0	Sports activity																																				
00	Football, rugby																																				
01	Football, Australian																																				
02	Football, soccer																																				
03	Hockey																																				
04	Squash																																				
05	Basketball																																				
06	Netball																																				
07	Cricket																																				
08	Roller blading																																				
09	Other and unspecified sporting activity																																				
1	Leisure activity (excluding sporting activity)																																				
2	Working for income																																				
3	Other types of work																																				
4	Resting, sleeping, eating or engaging in other vital activities																																				
5	Other specified activities																																				
6	Unspecified activities																																				

Collection and usage attributes

Guide for use:

To be used for injury surveillance purposes for non-admitted patients when it is not possible to use ICD-10-AM codes. Select the code which best characterises the type of activity being undertaken by the person when injured, on the basis of the information available at the time it is recorded. If two or more categories are judged to be equally appropriate, select the one that comes first in the code list.

Data element attributes

Collection and usage attributes

Comments:

Enables categorisation of injury and poisoning according to factors important for injury control. Necessary for defining and monitoring injury control targets, injury costing and identifying cases for in-depth research. This item is the basis for identifying work-related and sport-related injuries.

Source and reference attributes

Origin:

National Centre for Classification in Health
National Injury Surveillance Unit

Relational attributes

Related metadata references:

Supersedes Activity when injured, version 3, DE, NHDD, NHIMG, Superseded 01/03/2005

Implementation in Data Set Specifications:

Injury surveillance DSS NHIG, Standard 03/05/2006
Injury surveillance NMDS NHIG, Superseded 03/05/2006

Implementation start date: 01/07/2005

Implementation end date: 30/06/2006

Injury surveillance NMDS NHIG, Superseded 07/12/2005

Actual place of birth

Identifying and definitional attributes

<i>Metadata item type:</i>	Data Element
<i>Technical name:</i>	Birth event – setting of birth (actual), code N
<i>METeOR identifier:</i>	269937
<i>Registration status:</i>	NHIG, Standard 01/03/2005
<i>Definition:</i>	The actual place where the birth occurred, as represented by a code.
<i>Context:</i>	Perinatal statistics

Data element concept attributes

<i>Data element concept:</i>	Birth event – setting of birth
<i>Definition:</i>	The place where the birth occurred.
<i>Object class:</i>	Birth event
<i>Property:</i>	Setting of birth

Value domain attributes

Representational attributes

<i>Representation class:</i>	Code														
<i>Data type:</i>	Number														
<i>Format:</i>	N														
<i>Maximum character length:</i>	1														
<i>Permissible values:</i>	<table><thead><tr><th>Value</th><th>Meaning</th></tr></thead><tbody><tr><td>1</td><td>Hospital, excluding birth centre</td></tr><tr><td>2</td><td>Birth centre, attached to hospital</td></tr><tr><td>3</td><td>Birth centre, free standing</td></tr><tr><td>4</td><td>Home</td></tr><tr><td>8</td><td>Other</td></tr><tr><td>9</td><td>Not stated</td></tr></tbody></table>	Value	Meaning	1	Hospital, excluding birth centre	2	Birth centre, attached to hospital	3	Birth centre, free standing	4	Home	8	Other	9	Not stated
Value	Meaning														
1	Hospital, excluding birth centre														
2	Birth centre, attached to hospital														
3	Birth centre, free standing														
4	Home														
8	Other														
9	Not stated														
<i>Supplementary values:</i>															

Collection and usage attributes

<i>Comments:</i>	The development of a definition of a birth centre is currently under consideration by the Commonwealth in conjunction with the states and territories.
------------------	--

Data element attributes

Collection and usage attributes

<i>Guide for use:</i>	This is to be recorded for each baby the mother delivers from this pregnancy. CODE 4 Home Should be reserved for those births that occur at the home intended. CODE 8 Other Used when birth occurs at a home other than that intended.
-----------------------	--

May also include a community health centre or be used for babies 'born before arrival'.

Source and reference attributes

Submitting organisation: National Perinatal Data Development Committee

Relational attributes

Related metadata references: Supersedes Actual place of birth, version 2, DE, NHDD, NHIMG, Superseded 01/03/2005

Implementation in Data Set Specifications: Perinatal NMDS NHIG, Superseded 07/12/2005

Implementation start date: 01/07/2005

Implementation end date: 30/06/2006

Perinatal NMDS NHIG, Superseded 06/09/2006

Implementation start date: 01/07/2006

Implementation end date: 30/06/2007

Information specific to this data set:

Used to analyse the risk factors and outcomes by place of birth. While most deliveries occur within hospitals, an increasing number of births now occur in other settings. It is important to monitor the births occurring outside hospitals and to ascertain whether or not the actual place of delivery was planned.

Perinatal NMDS 2007-2008 NHIG, Standard 06/09/2006

Implementation start date: 01/07/2007

Information specific to this data set:

Used to analyse the risk factors and outcomes by place of birth. While most deliveries occur within hospitals, an increasing number of births now occur in other settings. It is important to monitor the births occurring outside hospitals and to ascertain whether or not the actual place of delivery was planned.

Acute coronary syndrome procedure type

Identifying and definitional attributes

<i>Metadata item type:</i>	Data Element
<i>Technical name:</i>	Person – acute coronary syndrome procedure type, code NN
<i>METeOR identifier:</i>	284660
<i>Registration status:</i>	NHIG, Standard 04/06/2004
<i>Definition:</i>	The type of procedure performed, that is pertinent to the treatment of acute coronary syndrome, as represented by a code.

Data element concept attributes

<i>Data element concept:</i>	Person – acute coronary syndrome procedure type
<i>Definition:</i>	The type of procedure performed, that is pertinent to the treatment of acute coronary syndrome.
<i>Context:</i>	Acute coronary syndrome treatment settings.
<i>Object class:</i>	Person
<i>Property:</i>	Acute coronary syndrome procedure type

Value domain attributes

Representational attributes

<i>Representation class:</i>	Code																																
<i>Data type:</i>	String																																
<i>Format:</i>	NN																																
<i>Maximum character length:</i>	2																																
<i>Permissible values:</i>	<table><thead><tr><th>Value</th><th>Meaning</th></tr></thead><tbody><tr><td>01</td><td>Coronary artery bypass graft (CABG)</td></tr><tr><td>02</td><td>Coronary stent (bare metal)</td></tr><tr><td>03</td><td>Coronary stent (drug eluding)</td></tr><tr><td>04</td><td>Angioplasty</td></tr><tr><td>05</td><td>Reperfusion fibrinolytic therapy</td></tr><tr><td>06</td><td>Reperfusion primary percutaneous coronary intervention (PCI)</td></tr><tr><td>07</td><td>Rescue angioplasty/stenting</td></tr><tr><td>08</td><td>Vascular reconstruction, bypass surgery, or percutaneous intervention to the extremities or for aortic aneurysm</td></tr><tr><td>09</td><td>Amputation for arterial vascular insufficiency</td></tr><tr><td>10</td><td>Diagnostic cardiac catheterisation/angiography</td></tr><tr><td>11</td><td>Blood transfusion</td></tr><tr><td>12</td><td>Insertion of pacemaker</td></tr><tr><td>13</td><td>Implantable cardiac defibrillator</td></tr><tr><td>14</td><td>Intra-aortic balloon pump (IABP)</td></tr><tr><td>15</td><td>Non-invasive ventilation (CPAP)</td></tr></tbody></table>	Value	Meaning	01	Coronary artery bypass graft (CABG)	02	Coronary stent (bare metal)	03	Coronary stent (drug eluding)	04	Angioplasty	05	Reperfusion fibrinolytic therapy	06	Reperfusion primary percutaneous coronary intervention (PCI)	07	Rescue angioplasty/stenting	08	Vascular reconstruction, bypass surgery, or percutaneous intervention to the extremities or for aortic aneurysm	09	Amputation for arterial vascular insufficiency	10	Diagnostic cardiac catheterisation/angiography	11	Blood transfusion	12	Insertion of pacemaker	13	Implantable cardiac defibrillator	14	Intra-aortic balloon pump (IABP)	15	Non-invasive ventilation (CPAP)
Value	Meaning																																
01	Coronary artery bypass graft (CABG)																																
02	Coronary stent (bare metal)																																
03	Coronary stent (drug eluding)																																
04	Angioplasty																																
05	Reperfusion fibrinolytic therapy																																
06	Reperfusion primary percutaneous coronary intervention (PCI)																																
07	Rescue angioplasty/stenting																																
08	Vascular reconstruction, bypass surgery, or percutaneous intervention to the extremities or for aortic aneurysm																																
09	Amputation for arterial vascular insufficiency																																
10	Diagnostic cardiac catheterisation/angiography																																
11	Blood transfusion																																
12	Insertion of pacemaker																																
13	Implantable cardiac defibrillator																																
14	Intra-aortic balloon pump (IABP)																																
15	Non-invasive ventilation (CPAP)																																

	16	Invasive ventilation
	17	Defibrillation
	88	Other
<i>Supplementary values:</i>	99	Not stated/inadequately described

Source and reference attributes

Submitting organisation: Australian Institute of Health and Welfare

Data element attributes

Collection and usage attributes

Guide for use: More than one procedure can be recorded. Record all codes that apply.
Codes '88' and '99' in combination cannot be used in multiple entries.
When read in conjunction with Person—clinical procedure timing, code N, this metadata item provides information on the procedure(s) provided to a patient prior to or during admission.
When read in conjunction with Person—acute coronary syndrome risk stratum, code N, codes 01 to 10 of this metadata item provide information for risk stratification.

Source and reference attributes

Submitting organisation: Acute coronary syndrome data working group

Steward: The National Heart Foundation of Australia and The Cardiac Society of Australia and New Zealand

Relational attributes

Related metadata references: Supersedes Acute coronary syndrome procedure type, version 1, DE, NHDD, NHIMG, Superseded 01/03/2005

Implementation in Data Set Specifications: Acute coronary syndrome (clinical) DSS NHIG, Standard 07/12/2005

Implementation start date: 07/12/2005

Acute coronary syndrome (clinical) DSS NHIG, Superseded 07/12/2005

Acute coronary syndrome stratum

Identifying and definitional attributes

<i>Metadata item type:</i>	Data Element
<i>Technical name:</i>	Person – acute coronary syndrome risk stratum, code N
<i>METeOR identifier:</i>	284656
<i>Registration status:</i>	NHIG, Standard 04/06/2004
<i>Definition:</i>	Risk stratum of the patient presenting with clinical features consistent with an acute coronary syndrome defined by accompanying clinical, electrocardiogram (ECG) and biochemical features, as represented by a code.

Data element concept attributes

<i>Data element concept:</i>	Person – acute coronary syndrome risk stratum
<i>Definition:</i>	Risk stratum of the patient presenting with clinical features consistent with an acute coronary syndrome (chest pain or overwhelming shortness of breath (SOB)) defined by accompanying clinical, electrocardiogram (ECG) and biochemical features.
<i>Context:</i>	Health care and clinical settings. The clinical, electrocardiogram and biochemical characteristics are important to enable early risk stratification.
<i>Object class:</i>	Person
<i>Property:</i>	Acute coronary syndrome risk stratum

Value domain attributes

Representational attributes

<i>Representation class:</i>	Code										
<i>Data type:</i>	Number										
<i>Format:</i>	N										
<i>Maximum character length:</i>	1										
<i>Permissible values:</i>	<table><thead><tr><th>Value</th><th>Meaning</th></tr></thead><tbody><tr><td>1</td><td>with ST elevation (myocardial infarction)</td></tr><tr><td>2</td><td>with non-ST elevation ACS with high-risk features</td></tr><tr><td>3</td><td>with non-ST elevation ACS with intermediate-risk features</td></tr><tr><td>4</td><td>with non-ST elevation ACS with low-risk features</td></tr></tbody></table>	Value	Meaning	1	with ST elevation (myocardial infarction)	2	with non-ST elevation ACS with high-risk features	3	with non-ST elevation ACS with intermediate-risk features	4	with non-ST elevation ACS with low-risk features
Value	Meaning										
1	with ST elevation (myocardial infarction)										
2	with non-ST elevation ACS with high-risk features										
3	with non-ST elevation ACS with intermediate-risk features										
4	with non-ST elevation ACS with low-risk features										
<i>Supplementary values:</i>	<table><tbody><tr><td>9</td><td>Not reported</td></tr></tbody></table>	9	Not reported								
9	Not reported										

Collection and usage attributes

<i>Guide for use:</i>	CODE 1 With ST elevation (myocardial infarction) This code is used where persistent ST elevation of ≥ 1 mm in two contiguous limb leads, or ST elevation of ≥ 2 mm in two contiguous chest leads, or with left bundle branch block (BBB) pattern on the ECG.
-----------------------	---

This classification is intended for identification of patients potentially eligible for reperfusion therapy, either pharmacologic or catheter-based. Other considerations such as the time to presentation and the clinical appropriateness of instituting reperfusion are not reflected in this metadata item.

CODE 2 With non-ST elevation ACS with high-risk features

This code is used when presentation with clinical features consistent with an acute coronary syndrome (chest pain or overwhelming SOB) with high-risk features which include either:

- classical rise and fall of at least one cardiac biomarker (troponin or CK-MB),
- persistent or dynamic ECG changes of ST segment depression ≥ 0.5 mm or new T wave inversion in three or more contiguous leads,
- transient ($= 0.5$ mm) in more than 2 contiguous leads,
- haemodynamic compromise: Blood pressure ≥ 1 , and/or new onset mitral regurgitation, and/or syncope, or
- presence of known diabetes without persistent ST elevation of > 1 mm in two or more contiguous leads or new or presumed new bundle branch block (BBB) pattern on the initial ECG, i.e. not meeting the definition for ST elevation MI.

This classification is intended for identification of patients potentially eligible for early invasive management and the use of intravenous glycoprotein IIb/IIIa inhibition.

CODE 3 With non-ST elevation ACS with intermediate-risk features

This code is used when presentation with clinical features consistent with an acute coronary syndrome (chest pain or overwhelming SOB) with intermediate-risk features which include either:

- prolonged but resolved chest pain/discomfort at rest age greater than 65yrs,
- known coronary heart disease: prior MI, prior revascularisation, known coronary lesion $> 50\%$,
- pathological Q waves or ECG changes of ST deviation nocturnal pain,
- two or more risk factors of known hypertension, family history, active smoking or hyperlipidaemia, or
- prior aspirin use and not meeting the definition for ST elevation MI or Non-ST elevation with high-risk features.

This classification is intended for identification of patients potentially eligible for admission and in-hospital investigation that may or may not include angiography.

CODE 4 With non-ST elevation ACS with low-risk features

This code is used when presentation with clinical features consistent with an acute coronary syndrome (chest pain or overwhelming SOB) without features of ST elevation MI or Non-ST elevation ACS with intermediate or high-risk features.

This classification is intended for identification of patients potentially eligible for outpatient investigation.

Source and reference attributes

Data element attributes

Collection and usage attributes

<i>Guide for use:</i>	Other clinical considerations influencing the decision to admit and investigate are not reflected in this metadata item. This metadata item is intended to simply provide a diagnostic classification at the time of, or within hours of clinical presentation.
<i>Collection methods:</i>	Collected at time of presentation. Only one code should be recorded. Must be collected in conjunction with Person – acute coronary syndrome procedure type, code NN and Person – clinical procedure timing, code N.

Source and reference attributes

<i>Submitting organisation:</i>	Acute coronary syndrome data working group
<i>Steward:</i>	The National Heart Foundation of Australia and The Cardiac Society of Australia and New Zealand
<i>Origin:</i>	Management of Unstable Angina Guidelines - 2000, The National Heart Foundation of Australia, The Cardiac Society of Australia and New Zealand MJA, 173 (Supplement) S65-S88 Antman, MD; et al. The TIMI Risk Score for Unstable Angina/Non-ST Elevation MI JAMA. 2000; 284:835-842.

Relational attributes

<i>Related metadata references:</i>	Supersedes Acute coronary syndrome stratum, version 1, DE, NHDD, NHIMG, Superseded 01/03/2005
<i>Implementation in Data Set Specifications:</i>	Acute coronary syndrome (clinical) DSS NHIG, Standard 07/12/2005 <i>Implementation start date:</i> 07/12/2005 Acute coronary syndrome (clinical) DSS NHIG, Superseded 07/12/2005

Additional diagnosis

Identifying and definitional attributes

<i>Metadata item type:</i>	Data Element
<i>Technical name:</i>	Episode of care – additional diagnosis, code (ICD-10-AM 5th edn) ANN{.N[N]}
<i>METeOR identifier:</i>	333832
<i>Registration status:</i>	NHIG, Standard 07/12/2005
<i>Definition:</i>	A condition or complaint either coexisting with the principal diagnosis or arising during the episode of admitted patient care, episode of residential care or attendance at a health care establishment, as represented by a code.

Data element concept attributes

<i>Data element concept:</i>	Episode of care – additional diagnosis
<i>Definition:</i>	A condition or complaint either coexisting with the principal diagnosis or arising during the episode of admitted patient care, episode of residential care or attendance at a health care establishment.
<i>Context:</i>	Additional diagnoses give information on factors which result in increased length of stay, more intensive treatment or the use of greater resources. They are used for casemix analyses relating to severity of illness and for correct classification of patients into Australian Refined Diagnosis Related Groups (AR-DRGs).
<i>Object class:</i>	Episode of care
<i>Property:</i>	Additional diagnosis

Value domain attributes

Representational attributes

<i>Classification scheme:</i>	International Statistical Classification of Diseases and Related Health Problems, Tenth Revision, Australian Modification 5th edition
<i>Representation class:</i>	Code
<i>Data type:</i>	String
<i>Format:</i>	ANN{.N[N]}
<i>Maximum character length:</i>	6

Data element attributes

Collection and usage attributes

<i>Guide for use:</i>	Record each additional diagnosis relevant to the episode of care in accordance with the ICD-10-AM Australian Coding Standards. Generally, external cause, place of occurrence and activity codes will be included in the string of additional diagnosis codes. In some data collections these codes may also be copied into specific fields. The diagnosis can include a disease, condition, injury,
-----------------------	---

poisoning, sign, symptom, abnormal finding, complaint, or other factor influencing health status.

Collection methods:

An additional diagnosis should be recorded and coded where appropriate upon separation of an episode of admitted patient care or the end of an episode of residential care. The additional diagnosis is derived from and must be substantiated by clinical documentation.

Comments:

Additional diagnoses are significant for the allocation of Australian Refined Diagnosis Related Groups. The allocation of patient to major problem or complication and co-morbidity Diagnosis Related Groups is made on the basis of the presence of certain specified additional diagnoses. Additional diagnoses should be recorded when relevant to the patient's episode of care and not restricted by the number of fields on the morbidity form or computer screen.

External cause codes, although not diagnosis of condition codes, should be sequenced together with the additional diagnosis codes so that meaning is given to the data for use in injury surveillance and other monitoring activities.

Source and reference attributes

Origin:

National Centre for Classification in Health

Relational attributes

Related metadata references:

Supersedes Episode of care—additional diagnosis, code (ICD-10-AM 4th edn) ANN{.N[N]} NHIG, Superseded 07/12/2005

Implementation in Data Set Specifications:

Admitted patient care NMDS 2007-2008 NHIG, Standard 29/11/2006

Implementation start date: 01/07/2007

Information specific to this data set:

An unlimited number of diagnosis and procedure codes should be able to be collected in hospital morbidity systems. Where this is not possible, a minimum of 20 codes should be able to be collected.

Effective for collection from 01/07/2007

Admitted patient mental health care NMDS NHIG, Superseded 23/10/2006

Implementation start date: 01/07/2006

Implementation end date: 30/06/2007

Information specific to this data set:

An unlimited number of diagnosis and procedure codes should be able to be collected in hospital morbidity systems. Where this is not possible, a minimum of 20 codes should be able to be collected.

Admitted patient mental health care NMDS 2007-2008 NHIG, Standard 23/10/2006

Implementation start date: 01/07/2007

Information specific to this data set:

An unlimited number of diagnosis and procedure codes should be able to be collected in hospital morbidity systems. Where this is not possible, a minimum of 20 codes should be able to be collected.

Admitted patient palliative care NMDS 2006-2007 NHIG,
Superseded 23/10/2006

Implementation start date: 01/07/2006

Implementation end date: 30/06/2007

Information specific to this data set:

An unlimited number of diagnosis and procedure codes should be able to be collected in hospital morbidity systems. Where this is not possible, a minimum of 20 codes should be able to be collected.

Effective for collection from 01/07/2006

Admitted patient palliative care NMDS 2007-08 NHIG,
Standard 23/10/2006

Implementation start date: 01/07/2007

Information specific to this data set:

An unlimited number of diagnosis and procedure codes should be able to be collected in hospital morbidity systems. Where this is not possible, a minimum of 20 codes should be able to be collected.

Effective for collection from 01/07/2006

Residential mental health care NMDS 2006-2007 NHIG,
Superseded 23/10/2006

Implementation start date: 01/07/2006

Implementation end date: 30/06/2007

Information specific to this data set:

Effective for collection from 01/07/2006

Residential mental health care NMDS 2007-2008 NHIG,
Standard 23/10/2006

Implementation start date: 01/07/2007

Information specific to this data set:

Effective for collection from 01/07/2006

Address line (person)

Identifying and definitional attributes

<i>Metadata item type:</i>	Data Element
<i>Technical name:</i>	Person (address)— address line, text [X(180)]
<i>METeOR identifier:</i>	286620
<i>Registration status:</i>	NHIG, Standard 04/05/2005 NCSIMG, Standard 30/09/2005
<i>Definition:</i>	A composite of one or more standard address components that describes a low level of geographical/physical description of a location, as represented by a text. Used in conjunction with the other high-level address components i.e. Suburb/town/locality, Postcode— Australian, Australian state/territory, and Country, forms a complete geographical/physical address of a person.

Data element concept attributes

<i>Data element concept:</i>	Person (address)— address line
<i>Definition:</i>	A composite of one or more standard address components that describes a low level of geographical/physical description of a location that, used in conjunction with the other high-level address components i.e. Suburb/town/locality, Postcode— Australian, Australian state/territory, and Country, forms a complete geographical/physical address of a person.
<i>Object class:</i>	Person
<i>Property:</i>	Address line

Value domain attributes

Representational attributes

<i>Representation class:</i>	Text
<i>Data type:</i>	String
<i>Format:</i>	[X(180)]
<i>Maximum character length:</i>	180

Data element attributes

Collection and usage attributes

<i>Guide for use:</i>	<p>A high-level address component is defined as a broad geographical area that is capable of containing more than one specific physical location. Some examples of a broad geographical area are:</p> <ul style="list-style-type: none">- Suburb, town or locality- Postcode— Australian or international- State, Territory, local government area, electorate, statistical local area- Postal delivery point identifier- Countries, provinces, etc other than in Australia <p>These components of a complete address do not form part of</p>
-----------------------	---

the Address line.

When addressing an Australian location, following are the standard address data elements that may be concatenated in the Address line:

- Building/complex sub-unit type
- Building/complex sub-unit number
- Building/property name
- Floor/level number
- Floor/level type
- House/property number
- Lot/section number
- Street name
- Street type code
- Street suffix code

One complete identification/description of a location/site of an address can comprise one or more than one instance of address line.

Instances of address lines are commonly identified in electronic information systems as Address-line 1, Address-line 2, etc.

The format of data collection is less important than consistent use of conventions in the recording of address data. Hence, address may be collected in an unstructured manner but should ideally be stored in a structured format.

Where Address line is collected as a stand-alone item, software may be used to parse the Address line details to separate the sub-components.

Multiple Address lines may be recorded as required.

Collection methods:

The following concatenation rules should be observed when collecting address lines addressing an Australian location.

- Building/complex sub-unit type is to be collected in conjunction with Building/complex sub-unit number and vice versa.
- Floor/level type is to be collected in conjunction with Floor/level number and vice versa.
- Street name is to be used in conjunction with Street type code and Street suffix code.
- Street type code is to be used in conjunction with Street name and Street suffix code.
- Street suffix code is to be used in conjunction with Street name and Street type code.
- House/property number is to be used in conjunction with Street name.

Source and reference attributes

<i>Submitting organisation:</i>	Standards Australia
<i>Origin:</i>	Health Data Standards Committee AS5017 Health Care Client Identification, 2002, Sydney: Standards Australia.
<i>Reference documents:</i>	AS4846 Health Care Provider Identification, 2004, Sydney: Standards Australia

Relational attributes

<i>Related metadata references:</i>	Supersedes Person (address) – health address line, text [X(180)]
-------------------------------------	--

NHIG, Superseded 04/05/2005

Is formed using Person (address) – building/property name, text [X(30)] NHIG, Standard 01/03/2005, NCSIMG, Standard 30/09/2005

Is formed using Person (address) – building/complex sub-unit identifier, [X(7)] NHIG, Standard 01/03/2005, NCSIMG, Standard 30/09/2005

Is formed using Person (address) – building/complex sub-unit type, code A[AAA] NHIG, Standard 01/03/2005, NCSIMG, Standard 30/09/2005

Is formed using Person (address) – floor/level identifier, [NNNA] NHIG, Standard 01/03/2005, NCSIMG, Standard 30/09/2005

Is formed using Person (address) – floor/level type, code A[A] NHIG, Standard 01/03/2005, NCSIMG, Standard 30/09/2005

Is formed using Person (address) – house/property identifier, text [X(12)] NHIG, Standard 01/03/2005, NCSIMG, Standard 30/09/2005

Is formed using Person (address) – lot/section identifier, N[X(14)] NHIG, Standard 01/03/2005, NCSIMG, Standard 30/09/2005

Is formed using Person (address) – street name, text [A(30)] NHIG, Standard 01/03/2005, NCSIMG, Standard 30/09/2005

Is formed using Person (address) – street type, code A[AAA] NHIG, Standard 01/03/2005, NCSIMG, Standard 30/09/2005

Is formed using Person (address) – street suffix, code A[A] NHIG, Standard 01/03/2005, NCSIMG, Standard 30/09/2005

Implementation in Data Set Specifications:

Cancer (clinical) DSS NHIG, Standard 07/12/2005

Cancer (clinical) DSS NHIG, Superseded 07/12/2005

Health care client identification DSS NHIG, Standard 04/05/2005

NCSIMG, Standard 03/10/2006

Health care provider identification DSS NHIG, Standard 04/05/2005

Address line (service provider organisation)

Identifying and definitional attributes

<i>Metadata item type:</i>	Data Element
<i>Technical name:</i>	Service provider organisation (address) – address line, text [X(180)]
<i>METeOR identifier:</i>	290315
<i>Registration status:</i>	NHIG, Standard 04/05/2005 NCSIMG, Standard 30/09/2005
<i>Definition:</i>	A composite of one or more standard address components, as represented by text.

Data element concept attributes

<i>Data element concept:</i>	Service provider organisation (address) – address line
<i>Definition:</i>	A composite of one or more standard address components that describes a low level of geographical/physical description of a location that, used in conjunction with the other high-level address components i.e. Suburb/town/locality, Postcode – Australian, Australian state/territory, and Country, forms a complete geographical/physical address of an organisation.
<i>Object class:</i>	Service provider organisation
<i>Property:</i>	Address line

Value domain attributes

Representational attributes

<i>Representation class:</i>	Text
<i>Data type:</i>	String
<i>Format:</i>	[X(180)]
<i>Maximum character length:</i>	180

Data element attributes

Collection and usage attributes

<i>Guide for use:</i>	<p>A high-level address component is defined as a broad geographical area that is capable of containing more than one specific physical location. Some examples of a broad geographical area are:</p> <ul style="list-style-type: none">• Suburb, town or locality• Postcode• Australian or international• State, Territory, local government area, electorate, statistical local area• Postal delivery point identifier• Countries, provinces, etc. other than in Australia <p>These components of a complete address do not form part of the Address line.</p> <p>When addressing an Australian location, following are the standard address data elements that may be concatenated in the</p>
-----------------------	---

Address line:

- Building/complex sub-unit type
- Building/complex sub-unit number
- Building/property name
- Floor/level number
- Floor/level type
- House/property number
- Lot/section number
- Street name
- Street type code
- Street suffix code

One complete identification/description of a location/site of an address can comprise one or more than one instance of address line. Instances of address lines are commonly identified in electronic information systems as Address-line 1, Address-line 2, etc. The format of data collection is less important than consistent use of conventions in the recording of address data. Hence, address may be collected in an unstructured manner but should ideally be stored in a structured format. Where Address line is collected as a stand-alone item, software may be used to parse the Address line details to separate the sub-components. Multiple Address lines may be recorded as required.

Collection methods:

The following concatenation rules should be observed when collecting address lines addressing an Australian location.

- Building/complex sub-unit type is to be collected in conjunction with Building/complex sub-unit number and vice versa.
- Floor/level type is to be collected in conjunction with Floor/level number and vice versa.
- Street name is to be used in conjunction with Street type code and Street suffix code.
- Street type code is to be used in conjunction with Street name and Street suffix code.
- Street suffix code is to be used in conjunction with Street name and Street type code.
- House/property number is to be used in conjunction with Street name.

Source and reference attributes

Submitting organisation:

Standards Australia

Origin:

Health Data Standards Committee

AS5017 Health Care Client Identification, 2002, Sydney:
Standards Australia.

Reference documents:

AS4846 Health Care Provider Identification, 2004, Sydney:
Standards Australia

Relational attributes

Related metadata references:

Is formed using Service provider organisation (address) – street suffix, code A[A] NHIG, Standard 04/05/2005, NCSIMG, Standard 30/09/2005

Is formed using Service provider organisation (address) – street type, code A[AAA] NHIG, Standard 04/05/2005, NCSIMG,

Standard 30/09/2005

Is formed using Service provider organisation (address) – street name, text [A(30)] NHIG, Standard 04/05/2005, NCSIMG, Standard 30/09/2005

Is formed using Service provider organisation (address) – lot/section identifier, N[X(14)] NHIG, Standard 04/05/2005, NCSIMG, Standard 30/09/2005

Is formed using Service provider organisation (address) – house/property identifier, text [X(12)] NHIG, Standard 04/05/2005, NCSIMG, Standard 30/09/2005

Is formed using Service provider organisation (address) – floor/level type, code A[A] NHIG, Standard 04/05/2005, NCSIMG, Standard 30/09/2005

Is formed using Service provider organisation (address) – floor/level identifier, [NNNA] NHIG, Standard 04/05/2005, NCSIMG, Standard 30/09/2005

Is formed using Service provider organisation (address) – building/complex sub-unit type, code A[AAA] NHIG, Standard 04/05/2005, NCSIMG, Standard 30/09/2005

Is formed using Service provider organisation (address) – building/complex sub-unit identifier, [X(7)] NHIG, Standard 04/05/2005, NCSIMG, Standard 30/09/2005

Is formed using Service provider organisation (address) – building/property name, text [X(30)] NHIG, Standard 04/05/2005, NCSIMG, Standard 30/09/2005

Implementation in Data Set Specifications:

Health care provider identification DSS NHIG, Standard 04/05/2005

Address type (person)

Identifying and definitional attributes

<i>Metadata item type:</i>	Data Element
<i>Technical name:</i>	Person (address) – address type, code N
<i>METeOR identifier:</i>	286728
<i>Registration status:</i>	NHIG, Standard 04/05/2005 NCSIMG, Standard 30/09/2005
<i>Definition:</i>	A code set representing a type of address, as represented by a code.

Data element concept attributes

<i>Data element concept:</i>	Person (address) – address type
<i>Definition:</i>	The type of geographical/physical location where a person can be located.
<i>Object class:</i>	Person
<i>Property:</i>	Address type

Value domain attributes

Representational attributes

<i>Representation class:</i>	Code												
<i>Data type:</i>	Number												
<i>Format:</i>	N												
<i>Maximum character length:</i>	1												
<i>Permissible values:</i>	<table><thead><tr><th>Value</th><th>Meaning</th></tr></thead><tbody><tr><td>1</td><td>Business</td></tr><tr><td>2</td><td>Mailing or postal</td></tr><tr><td>3</td><td>Residential</td></tr><tr><td>4</td><td>Temporary residential</td></tr><tr><td>9</td><td>Unknown/Not stated/inadequately described</td></tr></tbody></table>	Value	Meaning	1	Business	2	Mailing or postal	3	Residential	4	Temporary residential	9	Unknown/Not stated/inadequately described
Value	Meaning												
1	Business												
2	Mailing or postal												
3	Residential												
4	Temporary residential												
9	Unknown/Not stated/inadequately described												
<i>Supplementary values:</i>													

Collection and usage attributes

<i>Guide for use:</i>	<p>CODE 1 Business This code is used to indicate an address that is the physical location of a business, an office or from where a service is delivered.</p> <p>CODE 2 Mailing or postal This code is used to indicate an address that is only for correspondence purposes.</p> <p>CODE 3 Residential This code is used to indicate where a person is living. Note that this code is not valid for organisations.</p> <p>CODE 4 Temporary residential Temporary accommodation address (such as for a person from rural Australia who is visiting an oncology centre for a course of treatment, or a person who usually resides overseas). Note that this is not valid for organisations.</p>
-----------------------	--

CODE 9 Unknown/Not stated/inadequately described
This code may also be used where the person has no fixed address or does not wish to have their residential or a correspondence address recorded.

Data element attributes

Collection and usage attributes

<i>Guide for use:</i>	A single address may have multiple address types associated with it. Record as many as required.
<i>Collection methods:</i>	At least one address must be recorded (this may be an unknown Address type). Health care establishments should always attempt to collect the residential address of a person who is a health care client when a service is provided. When recording the address for a health care provider or organisation, the business address should always be collected. In addition, other addresses may also need to be recorded for individuals and organisations. Overseas address: For individuals record the overseas address as the residential address and record a temporary accommodation address as their contact address in Australia.
<i>Comments:</i>	'No fixed address' is coded as unknown because it (the concept) is not a type of address for a person but is an attribute of the person only i.e. it is not a location for which an address may be derived. It is not recommended that an implementation collects this attribute as an address type. A person not having a fixed address constrains the number of address types that can be collected i.e. temporary accommodation and residential address types cannot be collected. However, if it is imperative that this occurs, it is suggested that code 9 be used.

Source and reference attributes

<i>Submitting organisation:</i>	Standards Australia Australian Institute of Health and Welfare
<i>Origin:</i>	AS5017 Health Care Client Identification, 2002, Sydney: Standards Australia
<i>Reference documents:</i>	AS4846 Health Care Provider Identification, 2004, Sydney: Standards Australia In AS4846 and AS5017 alternative alphabetic codes are presented. Refer to the current standard for more details.

Relational attributes

<i>Related metadata references:</i>	Supersedes Person (address) – address type, code A NHIG, Superseded 04/05/2005
<i>Implementation in Data Set Specifications:</i>	Health care client identification DSS NHIG, Standard 04/05/2005 NCSIMG, Standard 03/10/2006 Health care provider identification DSS NHIG, Standard 04/05/2005

Address type (service provider organisation)

Identifying and definitional attributes

<i>Metadata item type:</i>	Data Element
<i>Technical name:</i>	Service provider organisation (address) – address type, code N
<i>METeOR identifier:</i>	286792
<i>Registration status:</i>	NHIG, Standard 04/05/2005 NCSIMG, Standard 30/09/2005
<i>Definition:</i>	The type of geographical/physical location where an organisation can be located, as represented by a code.

Data element concept attributes

<i>Data element concept:</i>	Service provider organisation (address) – address type
<i>Definition:</i>	The type of geographical/physical location where an organisation can be located.
<i>Object class:</i>	Service provider organisation
<i>Property:</i>	Address type

Value domain attributes

Representational attributes

<i>Representation class:</i>	Code								
<i>Data type:</i>	Number								
<i>Format:</i>	N								
<i>Maximum character length:</i>	1								
<i>Permissible values:</i>	<table><thead><tr><th>Value</th><th>Meaning</th></tr></thead><tbody><tr><td>1</td><td>Business</td></tr><tr><td>2</td><td>Mailing or postal</td></tr><tr><td>9</td><td>Unknown/Not stated/inadequately described</td></tr></tbody></table>	Value	Meaning	1	Business	2	Mailing or postal	9	Unknown/Not stated/inadequately described
Value	Meaning								
1	Business								
2	Mailing or postal								
9	Unknown/Not stated/inadequately described								
<i>Supplementary values:</i>									

Collection and usage attributes

<i>Guide for use:</i>	<p>CODE 1 Business This code is used to indicate an address that is the physical location of a business, an office or from where a service is delivered.</p> <p>CODE 2 Mailing or postal This code is used to indicate an address that is only for correspondence purposes.</p> <p>CODE 9 Unknown/Not stated/inadequately described This code may also be used where the person has no fixed address or does not wish to have their residential or a correspondence address recorded</p>
-----------------------	--

Data element attributes

Collection and usage attributes

<i>Guide for use:</i>	A single address may have multiple address types associated
-----------------------	---

with it. Record as many as required.

Collection methods:

At least one address must be recorded (this may be an unknown Address type). When recording the address for a health care provider or organisation, the business address should always be collected. In addition, other addresses may also need to be recorded for individuals and organisations.

Source and reference attributes

Origin:

AS5017 Health Care Client Identification, 2002, Sydney: Standards Australia

Reference documents:

AS4846 Health Care Provider Identification, 2004, Sydney: Standards Australia

In AS4846 and AS5017 alternative alphabetic codes are presented. Refer to the current standard for more details.

Relational attributes

Implementation in Data Set Specifications:

Health care provider identification DSS NHIG, Standard 04/05/2005

Address—country identifier (person)

Identifying and definitional attributes

<i>Metadata item type:</i>	Data Element
<i>Technical name:</i>	Person (address)—country identifier, code (SACC 1998) NNNN
<i>METeOR identifier:</i>	288091
<i>Registration status:</i>	NHIG, Standard 04/05/2005 NCSIMG, Standard 30/09/2005
<i>Definition:</i>	The country component of the address of a person, as represented by a code.

Data element concept attributes

<i>Data element concept:</i>	Person (address)—country identifier
<i>Definition:</i>	The country component of the address of a person.
<i>Object class:</i>	Person
<i>Property:</i>	Country identifier

Value domain attributes

Representational attributes

<i>Classification scheme:</i>	Standard Australian Classification of Countries 1998
<i>Representation class:</i>	Code
<i>Data type:</i>	Number
<i>Format:</i>	NNNN
<i>Maximum character length:</i>	4

Collection and usage attributes

<i>Guide for use:</i>	<p>The Standard Australian Classification of Countries 1998 (SACC) is a four-digit, three-level hierarchical structure specifying major group, minor group and country.</p> <p>A country, even if it comprises other discrete political entities such as states, is treated as a single unit for all data domain purposes. Parts of a political entity are not included in different groups. Thus, Hawaii is included in Northern America (as part of the identified country United States of America), despite being geographically close to and having similar social and cultural characteristics as the units classified to Polynesia.</p>
-----------------------	--

Data element attributes

Collection and usage attributes

<i>Collection methods:</i>	Collect the data at the 4-digit level.
<i>Comments:</i>	Note that the Standard Australian Classification of Countries (SACC) is mappable to but not identical to Australian Standard Classification of Countries for Social Statistics (ASCCSS).

Source and reference attributes

<i>Reference documents:</i>	Standard Australian Classification of Countries, Catalogue
-----------------------------	--

number 1269.0, 1998, Canberra: Australian Bureau of Statistics
Standard Australian Classification of Countries, Revision 2.01,
Canberra 1999, Australian Bureau of Statistics. Catalogue
Number 1269.0
Standard Australian Classification of Countries, Revision 2.02,
Canberra 2004, Australian Bureau of Statistics. Catalogue
Number 1269.0

Relational attributes

*Implementation in Data Set
Specifications:*

Health care client identification DSS NHIG, Standard
04/05/2005
NCSIMG, Standard 03/10/2006
Health care provider identification DSS NHIG, Standard
04/05/2005

Admission date

Identifying and definitional attributes

<i>Metadata item type:</i>	Data Element
<i>Technical name:</i>	Episode of admitted patient care – admission date, DDMMYYYY
<i>METeOR identifier:</i>	269967
<i>Registration status:</i>	NHIG, Standard 01/03/2005
<i>Definition:</i>	Date on which an admitted patient commences an episode of care.

Data element concept attributes

<i>Data element concept:</i>	Episode of admitted patient care – admission date
<i>Definition:</i>	Date on which an admitted patient commences an episode of care.
<i>Context:</i>	Required to identify the period in which the admitted patient episode and hospital stay occurred and for derivation of length of stay.
<i>Object class:</i>	Episode of admitted patient care
<i>Property:</i>	Admission date

Value domain attributes

Representational attributes

<i>Representation class:</i>	Date
<i>Data type:</i>	Date/Time
<i>Format:</i>	DDMMYYYY
<i>Maximum character length:</i>	8

Data element attributes

Source and reference attributes

<i>Origin:</i>	National Health Data Committee
----------------	--------------------------------

Relational attributes

<i>Related metadata references:</i>	Supersedes Admission date, version 4, DE, NHDD, NHIMG, Superseded 01/03/2005
	Is used in the formation of Episode of admitted patient care – length of stay (excluding leave days), total N[NN] NHIG, Standard 01/03/2005
	Is used in the formation of Episode of care – number of psychiatric care days, total N[NNNN] NHIG, Standard 01/03/2005
	Is used in the formation of Episode of admitted patient care – major diagnostic category, code (AR-DRG v5.1) NN NHIG, Standard 01/03/2005
	Is used in the formation of Episode of admitted patient care – length of stay (including leave days), total N[NN] NHIG, Standard 01/03/2005

Is used in the formation of Episode of admitted patient care – diagnosis related group, code (AR-DRG v5.1) ANNA NHIG, Standard 01/03/2005

Is used in the formation of Episode of admitted patient care (antenatal) – length of stay (including leave days), total N[NN] NHIG, Standard 01/03/2005

Is used in the formation of Non-admitted patient emergency department service episode – waiting time (to hospital admission), total hours and minutes NNNN NHIG, Standard 01/03/2005

Is used in the formation of Elective surgery waiting list episode – waiting time (at removal), total days N[NNN] NHIG, Standard 01/03/2005

Implementation in Data Set Specifications:

Admitted patient care NMDS NHIG, Superseded 07/12/2005

Implementation start date: 01/07/2005

Implementation end date: 30/06/2006

Information specific to this data set:

Right justified and zero filled.

admission date ≤ separation date

admission date ≥ date of birth

Admitted patient care NMDS 2007-2008 NHIG, Standard 29/11/2006

Implementation start date: 01/07/2007

Information specific to this data set:

Right justified and zero filled.

admission date ≤ separation date

admission date ≥ date of birth

Admitted patient mental health care NMDS NHIG, Superseded 07/12/2005

Implementation start date: 01/07/2005

Implementation end date: 30/06/2006

Information specific to this data set:

Right justified and zero filled.

admission date ≤ separation date

admission date ≥ date of birth

Admitted patient mental health care NMDS NHIG, Superseded 23/10/2006

Implementation start date: 01/07/2006

Implementation end date: 30/06/2007

Information specific to this data set:

Right justified and zero filled.

admission date ≤ separation date

admission date ≥ date of birth

Admitted patient mental health care NMDS 2007-2008 NHIG, Standard 23/10/2006

Implementation start date: 01/07/2007

Information specific to this data set:

Right justified and zero filled.
admission date ≤ separation date
admission date ≥ date of birth

Admitted patient palliative care NMDS NHIG, Superseded
07/12/2005

Implementation start date: 01/07/2005

Implementation end date: 30/06/2006

Information specific to this data set:

Right justified and zero filled.
admission date ≤ separation date
admission date ≥ date of birth

Admitted patient palliative care NMDS 2006-2007 NHIG,
Superseded 23/10/2006

Implementation start date: 01/07/2006

Implementation end date: 30/06/2007

Information specific to this data set:

Right justified and zero filled.
admission date ≤ separation date
admission date ≥ date of birth

Admitted patient palliative care NMDS 2007-08 NHIG,
Standard 23/10/2006

Implementation start date: 01/07/2007

Information specific to this data set:

Right justified and zero filled.
admission date ≤ separation date
admission date ≥ date of birth

Admission time

Identifying and definitional attributes

<i>Metadata item type:</i>	Data Element
<i>Technical name:</i>	Episode of admitted patient care – admission time, hhmm
<i>METeOR identifier:</i>	269972
<i>Registration status:</i>	NHIG, Standard 01/03/2005
<i>Definition:</i>	Time at which an admitted patient commences an episode of care.

Data element concept attributes

<i>Data element concept:</i>	Episode of admitted patient care – admission time
<i>Definition:</i>	Time at which an admitted patient commences an episode of care.
<i>Context:</i>	Admitted patient care
<i>Object class:</i>	Episode of admitted patient care
<i>Property:</i>	Admission time

Value domain attributes

Representational attributes

<i>Representation class:</i>	Time
<i>Data type:</i>	Date/Time
<i>Format:</i>	hhmm
<i>Maximum character length:</i>	4

Source and reference attributes

<i>Reference documents:</i>	ISO 8601:2000 : Data elements and interchange formats - Information interchange - Representation of dates and times
-----------------------------	---

Data element attributes

Collection and usage attributes

<i>Comments:</i>	Required to identify the time of commencement of the episode or hospital stay, for calculation of waiting times and length of stay.
------------------	---

Source and reference attributes

<i>Origin:</i>	National Health Data Committee
----------------	--------------------------------

Relational attributes

<i>Related metadata references:</i>	Supersedes Admission time, version 2, DE, NHDD, NHIMG, Superseded 01/03/2005 Is used in the formation of Non-admitted patient emergency department service episode – waiting time (to hospital admission), total hours and minutes NNNN NHIG, Standard 01/03/2005
-------------------------------------	--

Admitted patient election status

Identifying and definitional attributes

<i>Metadata item type:</i>	Data Element
<i>Technical name:</i>	Episode of admitted patient care – patient election status, code N
<i>METeOR identifier:</i>	326619
<i>Registration status:</i>	NHIG, Standard 29/11/2006
<i>Definition:</i>	Accommodation chargeable status elected by a patient on admission , as represented by a code.

Data element concept attributes

<i>Data element concept:</i>	Episode of admitted patient care – patient election status
<i>Definition:</i>	Accommodation chargeable status elected by a patient on admission .
<i>Context:</i>	Admitted patient care.
<i>Object class:</i>	Episode of admitted patient care
<i>Property:</i>	Patient election status

Value domain attributes

Representational attributes

<i>Representation class:</i>	Code						
<i>Data type:</i>	Number						
<i>Format:</i>	N						
<i>Maximum character length:</i>	1						
<i>Permissible values:</i>	<table><thead><tr><th>Value</th><th>Meaning</th></tr></thead><tbody><tr><td>1</td><td>Public</td></tr><tr><td>2</td><td>Private</td></tr></tbody></table>	Value	Meaning	1	Public	2	Private
Value	Meaning						
1	Public						
2	Private						

Collection and usage attributes

<i>Guide for use:</i>	<p>Public patient: A person, eligible for Medicare, who receives or elects to receive a public hospital service free of charge. Includes: patients in public psychiatric hospitals who do not have the choice to be treated as a private patient. Also includes overseas visitors who are covered by a reciprocal health care agreement, and who elect to be treated as public patients.</p> <p>Private patient: A person who elects to be treated as a private patient and elects to be responsible for paying fees for the type referred to in clause 49 of the Australian Health Care Agreements (2003–2008). Clause 49 states that: Private patients, compensable patients and ineligible persons may be charged an amount for public hospital services as determined by (the state or territory). All patients in private hospitals (other than those receiving</p>
-----------------------	--

public hospital services and electing to be treated as a public patient) are private patients.

Includes: all patients who are charged (regardless of the level of the charge) or for whom a charge is raised for a third party payer (for example, Department of Veterans' Affairs and Compensable patients). Also includes patients who are Medicare ineligible and receive public hospital services free of charge at the discretion of the hospital, and prisoners, who are Medicare ineligible while incarcerated.

Data element attributes

Collection and usage attributes

Guide for use:

Australian Health Care Agreements 2003–08 state that eligible persons are to be given the choice to receive, free of charge as public patients, health and emergency services.

At the time of, or as soon as practicable after, admission for a public hospital service, the patient must elect in writing to be treated as either

- a public patient or
- a private patient

This item is independent of the patient's hospital insurance status and room type.

Notes:

Inability to sign: In cases where the patient is unable to complete the patient election form, the patient should be assumed to be a public patient.

Compensation funding decisions: A patient may be recorded as a public patient as an interim patient election status while the patient's compensable status is being decided.

Inter-hospital contracted care: If the patient receives inter-hospital contracted care the following guidelines can be used if no further information is available:

- If the patient received contracted care that was purchased by a public hospital then it will be assumed that they elected to be treated as a public patient.
- If the patient received contracted care that was purchased by a private hospital then it will be assumed that they elected to be treated as a private patient.

Source and reference attributes

Submitting organisation:

Admitted patient care NMDS Technical Reference Group

Relational attributes

Related metadata references:

Supersedes Episode of admitted patient care – elected accommodation status, code N NHIG, Superseded 29/11/2006

Implementation in Data Set Specifications:

Admitted patient care NMDS 2007-2008 NHIG, Standard 29/11/2006

Implementation start date: 01/07/2007

Age

Identifying and definitional attributes

<i>Metadata item type:</i>	Data Element
<i>Technical name:</i>	Person – age, total years N[NN]
<i>METeOR identifier:</i>	303794
<i>Registration status:</i>	NHIG, Standard 08/02/2006 NCSIMG, Standard 29/04/2006 NHDAMG, Standard 10/02/2006
<i>Definition:</i>	The age of the person in (completed) years at a specific point in time.
<i>Context:</i>	Age is a core data element in a wide range of social, labour and demographic statistics. It is used in the analyses of service utilisation by age group and can be used as an assistance eligibility criterion.

Data element concept attributes

<i>Data element concept:</i>	Person – age
<i>Definition:</i>	The age of the person.
<i>Context:</i>	Age is a core data element in a wide range of social, labour and demographic statistics.
<i>Object class:</i>	Person
<i>Property:</i>	Age

Value domain attributes

Representational attributes

<i>Representation class:</i>	Total
<i>Data type:</i>	Number
<i>Format:</i>	N[NN]
<i>Maximum character length:</i>	3
<i>Supplementary values:</i>	Value Meaning 999 Unknown/not stated
<i>Unit of measure:</i>	Year

Data element attributes

Collection and usage attributes

<i>Guide for use:</i>	Age in single years (if aged under one year, record as zero). If age (or date of birth) is unknown or not stated, and cannot be estimated, use Code 999. National community services and housing assistance data dictionary specific: If year of birth is known (but date of birth is not) use the date, 0101YYYY of the birth year to estimate age (where YYYY is the year of birth). National housing assistance data dictionary specific:
-----------------------	--

In the housing assistance data collections age is calculated at 30 June for the corresponding year.

Collection methods:

Although collection of date of birth allows more precise calculation of age, this may not be feasible in some data collections, and alternative questions are: Age last birthday?
What was age last birthday?
What is age in complete years?

Comments:

National community services data dictionary specific:
Different rules for reporting data may apply when estimating the Date of birth of children aged under 2 years since the rapid growth and development of children within this age group means that a child's development can vary considerably over the course of a year. Thus, more specific reporting of estimated age is recommended.
Those who need to conduct data collections for children where age is collected in months, weeks, or days should do so in a manner that allows for aggregation of those results to this standard.

Source and reference attributes

Submitting organisation:

National Public Health Information Working Group

Origin:

Australian Bureau of Statistics, *Standards for Social, Labour and Demographic Variables*. Reference through:
www.abs.gov.au/Ausstats/abs@.nsf/StatsLibrary

Relational attributes

Related metadata references:

Supersedes Person – age, total years N[NN] NHIG, Superseded 08/02/2006

Implementation in Data Set Specifications:

Computer Assisted Telephone Interview demographic module DSS NHIG, Standard 04/05/2005

Information specific to this data set:

In CATI surveys, age refers to completed age of respondent on day of interview.

If collecting age in single years is not possible, age may be collected as a range. Refer to the data element Person – age range, code NN.

Age range

Identifying and definitional attributes

<i>Metadata item type:</i>	Data Element
<i>Technical name:</i>	Person – age range, code NN
<i>METeOR identifier:</i>	290540
<i>Registration status:</i>	NHIG, Standard 04/05/2005
<i>Definition:</i>	The age range that best accommodates a person's completed age in years, at the time of data collection, as represented by a code.

Data element concept attributes

<i>Data element concept:</i>	Person – age range
<i>Definition:</i>	The age range that best accommodates a person's completed age in years
<i>Context:</i>	
<i>Object class:</i>	Person
<i>Property:</i>	Age range

Value domain attributes

Representational attributes

<i>Representation class:</i>	Code																						
<i>Data type:</i>	Number																						
<i>Format:</i>	NN																						
<i>Maximum character length:</i>	2																						
<i>Permissible values:</i>	<table><thead><tr><th>Value</th><th>Meaning</th></tr></thead><tbody><tr><td>01</td><td>0-4</td></tr><tr><td>02</td><td>5-14</td></tr><tr><td>03</td><td>15-24</td></tr><tr><td>04</td><td>25-34</td></tr><tr><td>05</td><td>35-44</td></tr><tr><td>06</td><td>45-54</td></tr><tr><td>07</td><td>55-64</td></tr><tr><td>08</td><td>65-74</td></tr><tr><td>09</td><td>75 years or older</td></tr><tr><td>99</td><td>Not stated</td></tr></tbody></table>	Value	Meaning	01	0-4	02	5-14	03	15-24	04	25-34	05	35-44	06	45-54	07	55-64	08	65-74	09	75 years or older	99	Not stated
Value	Meaning																						
01	0-4																						
02	5-14																						
03	15-24																						
04	25-34																						
05	35-44																						
06	45-54																						
07	55-64																						
08	65-74																						
09	75 years or older																						
99	Not stated																						
<i>Supplementary values:</i>	99 Not stated																						

Data element attributes

Collection and usage attributes

<i>Guide for use:</i>	Used in computer assisted telephone interview (CATI) surveys in cases where the specific age is not available. Depending on the collection a different starting age may be used, but should map back to the standard output. Information at a finer level can be collected as long as it maps
-----------------------	---

back to the proposed data domain, e.g. 75+ age group can be split into 75-84 and 85 years or older.

Collection methods:

Although collection of date of birth allows more precise calculation of age, as does the collection of a single age, this may not always be feasible. Age range should be derived from a question on date of birth or age at last birthday.

Comments:

In cases where an exact age is not known or not stated, age may be reported as an age range. The age ranges are consistent with the standard 10 year ranges recommended by the ABS.

Source and reference attributes

Submitting organisation:

National Public Health Information Working Group

Origin:

ABS, Statistical Concepts Library, Standards for Social, Labour and Demographic Variables. Age.

Reference documents:

Reference through:

<http://www.abs.gov.au/Ausstats/abs@.nsf/StatsLibrary> and choose, Other ABS Statistical Standards, Standards for Social, Labour and Demographic Variables, Demographic Variables, Age.

Relational attributes

Implementation in Data Set Specifications:

Computer Assisted Telephone Interview demographic module
DSS NHIG, Standard 04/05/2005

Information specific to this data set:

For some data collection settings, using Computer Assisted Telephone Interviewing (CATI), the suggested question is :

Which age group are you in? Would it be....

0-4

5-14

15-24

25-34

35-44

45-54

55-64

65-74

75 years or older

Refused

Alcohol consumption frequency (self reported)

Identifying and definitional attributes

<i>Metadata item type:</i>	Data Element
<i>Technical name:</i>	Person – alcohol consumption frequency (self-reported), code NN
<i>METeOR identifier:</i>	270247
<i>Registration status:</i>	NHIG, Standard 01/03/2005
<i>Definition:</i>	A person's self-reported frequency of alcohol consumption, as represented by a code.

Data element concept attributes

<i>Data element concept:</i>	Person – alcohol consumption frequency
<i>Definition:</i>	An indicator of how frequently alcohol is consumed by a person.
<i>Context:</i>	Public health, health care and clinical settings.
<i>Object class:</i>	Person
<i>Property:</i>	Alcohol consumption frequency

Value domain attributes

Representational attributes

<i>Representation class:</i>	Code																										
<i>Data type:</i>	String																										
<i>Format:</i>	NN																										
<i>Maximum character length:</i>	2																										
<i>Permissible values:</i>	<table><thead><tr><th>Value</th><th>Meaning</th></tr></thead><tbody><tr><td>01</td><td>Every day/7 days per week</td></tr><tr><td>02</td><td>5 to 6 days per week</td></tr><tr><td>03</td><td>3 to 4 days per week</td></tr><tr><td>04</td><td>1 to 2 days per week</td></tr><tr><td>05</td><td>2 to 3 days per month</td></tr><tr><td>06</td><td>Once per month</td></tr><tr><td>07</td><td>7 to 11 days in the past year</td></tr><tr><td>08</td><td>4 to 6 days in the past year</td></tr><tr><td>09</td><td>2 to 3 days in the past year</td></tr><tr><td>10</td><td>Once in the past year</td></tr><tr><td>11</td><td>Never drank any alcoholic beverage in the past year</td></tr><tr><td>12</td><td>Never in my life</td></tr></tbody></table>	Value	Meaning	01	Every day/7 days per week	02	5 to 6 days per week	03	3 to 4 days per week	04	1 to 2 days per week	05	2 to 3 days per month	06	Once per month	07	7 to 11 days in the past year	08	4 to 6 days in the past year	09	2 to 3 days in the past year	10	Once in the past year	11	Never drank any alcoholic beverage in the past year	12	Never in my life
Value	Meaning																										
01	Every day/7 days per week																										
02	5 to 6 days per week																										
03	3 to 4 days per week																										
04	1 to 2 days per week																										
05	2 to 3 days per month																										
06	Once per month																										
07	7 to 11 days in the past year																										
08	4 to 6 days in the past year																										
09	2 to 3 days in the past year																										
10	Once in the past year																										
11	Never drank any alcoholic beverage in the past year																										
12	Never in my life																										
<i>Supplementary values:</i>	99 Not reported																										

Data element attributes

Collection and usage attributes

Collection methods:

The World Health Organisation, in its 2000 International Guide for Monitoring Alcohol Consumption and Related Harm document, suggests that in assessing alcohol consumption patterns a 'Graduated Quantity Frequency' method is preferred. This method requires that questions about the quantity and frequency of alcohol consumption should be asked to help determine short-term and long-term health consequences. This information can be collected (but not confined to) the following ways:

- in a clinical setting with questions asked by a primary healthcare professional
- as a self-completed questionnaire in a clinical setting
- as part of a health survey
- as part of a computer aided telephone interview.

It should be noted that, particularly in telephone interviews, the question(s) asked may not be a direct repetition of the Value domain; yet they may still yield a response that could be coded to the full Value domain or a collapsed version of the Value domain.

Source and reference attributes

Submitting organisation:

Cardiovascular Data Working Group

Origin:

Australian Alcohol Guidelines: Health Risks and Benefits, National Health & Medical Research Council, October 2001

Relational attributes

Related metadata references:

Supersedes Alcohol consumption frequency- self report, version 1, DE, NHDD, NHIMG, Superseded 01/03/2005

Implementation in Data Set Specifications:

Cardiovascular disease (clinical) DSS NHIG, Superseded 15/02/2006

Information specific to this data set:

These data can be used to help determine the overall health profile of an individual or of a population. Certain patterns of alcohol consumption can be associated with a range of social and health problems. These problems include:

- social problems such as domestic violence, unsafe sex,
- financial and relationship problems,
- physical conditions such as high blood pressure, gastrointestinal problems, pancreatitis,
- an increased risk of physical injury.

Alcohol can also be a contributor to acute health problems.

Evidence from prospective studies indicates that heavy alcohol consumption is associated with increased mortality and morbidity from coronary heart disease and stroke (Hanna et al 1992). However, there is some evidence to suggest that alcohol appears to provide some protection against heart disease (both illness and death) for both men and women from middle age onwards. Most, if not all, of this benefit is achieved with 1-2 standard drinks per day for men and less than 1 standard drink for women (the National Health and Medical Research Council's Australian Alcohol Guidelines, October 2001).

Where this information is collected by survey and the sample permits, population estimates should be presented by sex and 5-year age groups. Summary statistics may need to be adjusted for age and other relevant variables. It is recommended that, in surveys of alcohol consumption, data on age, sex, and other socio-demographic variables also be collected where it is possible and desirable to do so. It is also recommended that, when alcohol consumption is investigated in relation to health, data on other risk factors including overweight and obesity, smoking, high blood pressure and physical inactivity should be collected. The Australian Alcohol Guidelines: Health Risk and Benefits endorsed by the National Health and Medical Research Council in October 2001 have defined risk of harm in the short term and long term based on patterns of drinking.

The table below outlines those patterns.

Alcohol consumption shown in the tables is not recommended for people who: - have a condition made worse by drinking,

- are on medication,
- are under 18 years of age,
- are pregnant,
- are about to engage in activities involving risk or a degree of skill (e.g. driving, flying, water sports, skiing, operating machinery).

Risk of harm in the short-term			
	Low risk (standard drinks)	Risky (standard drinks)	High risk (standard drinks)
Males (on a single occasion)	Up to 6	7 to 10	11 or more
Females (on a single occasion)	Up to 4	5 to 6	7 or more

Source: NH&MRC Australian Alcohol Guidelines: Health Risk and Benefits 2001.

<i>Risk of harm in the long-term</i>			
	<i>Low risk (standard drinks)</i>	<i>Risky (standard drinks)</i>	<i>High risk (standard drinks)</i>
<i>Males (on an average day)</i>	<i>Up to 4</i>	<i>5 to 6</i>	<i>7 or more</i>
<i>Overall weekly level</i>	<i>Up to 28 Per week</i>	<i>29 to 42 Per week</i>	<i>43 or more Per week</i>
<i>Females (on an average day)</i>	<i>Up to 2</i>	<i>3 to 4</i>	<i>5 or more</i>

<i>Overall weekly level</i>	<i>Up to 14 Per week</i>	<i>15 to 28 Per week</i>	<i>29 or more Per week</i>
-----------------------------	--------------------------	--------------------------	----------------------------

Source: NH&MRC Australian Alcohol Guidelines: Health Risk and Benefits 2001.

Cardiovascular disease (clinical) DSS NHIG, Standard
15/02/2006

Information specific to this data set:

These data can be used to help determine the overall health profile of an individual or of a population. Certain patterns of alcohol consumption can be associated with a range of social and health problems. These problems include:

- social problems such as domestic violence, unsafe sex,
- financial and relationship problems,
- physical conditions such as high blood pressure, gastrointestinal problems, pancreatitis,
- an increased risk of physical injury.

Alcohol can also be a contributor to acute health problems. Evidence from prospective studies indicates that heavy alcohol consumption is associated with increased mortality and morbidity from coronary heart disease and stroke (Hanna et al 1992). However, there is some evidence to suggest that alcohol appears to provide some protection against heart disease (both illness and death) for both men and women from middle age onwards. Most, if not all, of this benefit is achieved with 1-2 standard drinks per day for men and less than 1 standard drink for women (the National Health and Medical Research Council's Australian Alcohol Guidelines, October 2001).

Where this information is collected by survey and the sample permits, population estimates should be presented by sex and 5-year age groups. Summary statistics may need to be adjusted for age and other relevant variables. It is recommended that, in surveys of alcohol consumption, data on age, sex, and other socio-demographic variables also be collected where it is possible and desirable to do so. It is also recommended that, when alcohol consumption is investigated in relation to health, data on other risk factors including overweight and obesity, smoking, high blood pressure and physical inactivity should be collected. The Australian Alcohol Guidelines: Health Risk and Benefits endorsed by the National Health and Medical Research Council in October 2001 have defined risk of harm in the short term and long term based on patterns of drinking.

The table below outlines those patterns.

Alcohol consumption shown in the tables is not recommended for people who: - have a condition made worse by drinking,

- are on medication,
- are under 18 years of age,
- are pregnant,

- are about to engage in activities involving risk or a degree of skill (e.g. driving, flying, water sports, skiing, operating machinery).

Risk of harm in the short-term			
	Low risk (standard drinks)	Risky (standard drinks)	High risk (standard drinks)
Males (on a single occasion)	Up to 6	7 to 10	11 or more
Females (on a single occasion)	Up to 4	5 to 6	7 or more

Source: NH&MRC Australian Alcohol Guidelines: Health Risk and Benefits 2001.

Risk of harm in the long-term			
	Low risk (standard drinks)	Risky (standard drinks)	High risk (standard drinks)
Males (on an average day)	Up to 4	5 to 6	7 or more
Overall weekly level	Up to 28 Per week	29 to 42 Per week	43 or more Per week
Females (on an average day)	Up to 2	3 to 4	5 or more
Overall weekly level	Up to 14 Per week	15 to 28 Per week	29 or more Per week

Source: NH&MRC Australian Alcohol Guidelines: Health Risk and Benefits 2001.

Alcohol consumption in standard drinks per day (self reported)

Identifying and definitional attributes

<i>Metadata item type:</i>	Data Element
<i>Technical name:</i>	Person – alcohol consumption amount (self-reported), total standard drinks NN
<i>METeOR identifier:</i>	270249
<i>Registration status:</i>	NHIG, Standard 01/03/2005
<i>Definition:</i>	A person's self-reported usual number of alcohol-containing standard drinks on a day when they consume alcohol.

Data element concept attributes

<i>Data element concept:</i>	Person – alcohol consumption amount
<i>Definition:</i>	The ethyl alcohol (ethanol) consumed by a person in alcoholic beverages such as beer, cider, wine, spirits and mixed drinks.
<i>Context:</i>	Public health, health care and clinical settings.
<i>Object class:</i>	Person
<i>Property:</i>	Alcohol consumption amount

Value domain attributes

Representational attributes

<i>Representation class:</i>	Total
<i>Data type:</i>	Number
<i>Format:</i>	NN
<i>Maximum character length:</i>	2
<i>Supplementary values:</i>	Value Meaning
	99 Consumption not reported
<i>Unit of measure:</i>	Standard drink

Collection and usage attributes

<i>Guide for use:</i>	Alcohol consumption is usually measured in standard drinks. An Australian standard drink contains 10 grams of alcohol, which is equivalent to 12.5 millilitres of alcohol.
-----------------------	--

Data element attributes

Collection and usage attributes

<i>Guide for use:</i>	This estimation is based on the person's description of the type (spirits, beer, wine, other) and number of standard drinks, as defined by the National Health and Medical Research Council (NH&MRC), consumed per day. One standard drink contains 10 grams of alcohol. The following gives the NH&MRC examples of a standard drink: <ul style="list-style-type: none">• Light beer (2.7%):
-----------------------	---

- 1 can or stubbie = 0.8 a standard drink
- Medium light beer (3.5%):
 - 1 can or stubbie = 1 standard drink
- Regular Beer - (4.9% alcohol):
 - 1 can = 1.5 standard drinks
 - 1 jug = 4 standard drinks
 - 1 slab (cans or stubbies) = about 36 standard drinks
- Wine (9.5% - 13% alcohol):
 - 750-ml bottle = about 7 to 8 standard drinks
 - 4-litre cask = about 30 to 40 standard drinks
- Spirits:
 - 1 nip = 1 standard drink
 - Pre-mixed spirits (around 5% alcohol) = 1.5 standard drinks

When calculating consumption in standard drinks per day, the total should be reported with part drinks recorded to the next whole standard drink (e.g. 2.4 = 3).

Collection methods:

The *World Health Organisation's 2000 International Guide for Monitoring Alcohol Consumption and Related Harm* document suggests that in assessing alcohol consumption patterns a 'Graduated Quantity Frequency' method is preferred. This method requires that questions about the quantity and frequency of alcohol consumption should be asked to help determine short-term and long-term health consequences.

Source and reference attributes

Submitting organisation:

Cardiovascular Data Working Group

Origin:

The World Health Organisation's 2000 International Guide for Monitoring Alcohol Consumption and Related Harm document -National Health and Medical Research Council's Australian Alcohol Guidelines, October 2001.

Relational attributes

Related metadata references:

Supersedes Alcohol consumption in standard drinks per day - self report, version 1, DE, NHDD, NHIMG, Superseded 01/03/2005

Implementation in Data Set Specifications:

Cardiovascular disease (clinical) DSS NHIG, Superseded 15/02/2006

Information specific to this data set:

These data are used to help determine the overall health profile of an individual. Certain patterns of alcohol consumption can be associated with a range of social and health problems. These problems include:

- social problems such as domestic violence, unsafe sex,
- financial and relationship problems,
- physical conditions such as high blood pressure, gastrointestinal problems, pancreatitis,
- an increased risk of physical injury.
- Alcohol can also be a contributor to acute health problems.

Evidence from prospective studies indicates that heavy alcohol consumption is associated with increased

mortality and morbidity from coronary heart disease and stroke (Hanna et al. 1992). However, there is some evidence to suggest that alcohol appears to provide some protection against heart disease (both illness and death) for both men and women from middle age onwards. Most if not all of this benefit is achieved with 1-2 standard drinks per day for men and less than 1 standard drink for women (the National Health and Medical Research Council's Australian Alcohol Guidelines, October 2001).

Cardiovascular disease (clinical) DSS NHIG, Standard
15/02/2006

Information specific to this data set:

These data are used to help determine the overall health profile of an individual. Certain patterns of alcohol consumption can be associated with a range of social and health problems. These problems include:

- social problems such as domestic violence, unsafe sex,
- financial and relationship problems,
- physical conditions such as high blood pressure, gastrointestinal problems, pancreatitis,
- an increased risk of physical injury.
- Alcohol can also be a contributor to acute health problems.

Evidence from prospective studies indicates that heavy alcohol consumption is associated with increased mortality and morbidity from coronary heart disease and stroke (Hanna et al. 1992). However, there is some evidence to suggest that alcohol appears to provide some protection against heart disease (both illness and death) for both men and women from middle age onwards. Most if not all of this benefit is achieved with 1-2 standard drinks per day for men and less than 1 standard drink for women (the National Health and Medical Research Council's Australian Alcohol Guidelines, October 2001).

Anaesthesia administered for operative delivery of the baby

Identifying and definitional attributes

<i>Metadata item type:</i>	Data Element
<i>Technical name:</i>	Birth event – anaesthesia administered, code N
<i>METeOR identifier:</i>	292044
<i>Registration status:</i>	NHIG, Standard 07/12/2005
<i>Definition:</i>	Anaesthesia administered to the woman for the operative delivery of the baby, as represented by a code.

Data element concept attributes

<i>Data element concept:</i>	Birth event – anaesthesia administered
<i>Definition:</i>	Anaesthesia administered to the woman for the operative delivery of the baby.
<i>Context:</i>	Perinatal statistics
<i>Object class:</i>	Birth event
<i>Property:</i>	Anaesthesia administered

Value domain attributes

Representational attributes

<i>Representation class:</i>	Code																		
<i>Data type:</i>	Number																		
<i>Format:</i>	N																		
<i>Maximum character length:</i>	1																		
<i>Permissible values:</i>	<table><thead><tr><th>Value</th><th>Meaning</th></tr></thead><tbody><tr><td>1</td><td>None</td></tr><tr><td>2</td><td>Local anaesthetic to perineum</td></tr><tr><td>3</td><td>Pudendal</td></tr><tr><td>4</td><td>Epidural or caudal</td></tr><tr><td>5</td><td>Spinal</td></tr><tr><td>6</td><td>General anaesthetic</td></tr><tr><td>7</td><td>Combined spinal-epidural</td></tr><tr><td>8</td><td>Other</td></tr></tbody></table>	Value	Meaning	1	None	2	Local anaesthetic to perineum	3	Pudendal	4	Epidural or caudal	5	Spinal	6	General anaesthetic	7	Combined spinal-epidural	8	Other
Value	Meaning																		
1	None																		
2	Local anaesthetic to perineum																		
3	Pudendal																		
4	Epidural or caudal																		
5	Spinal																		
6	General anaesthetic																		
7	Combined spinal-epidural																		
8	Other																		
<i>Supplementary values:</i>	9 Not stated/inadequately described																		

Data element attributes

Collection and usage attributes

<i>Guide for use:</i>	Operative delivery includes caesarean section, forceps and vacuum extraction. Code 7: this code is used when this technique has been selected for the administration of anaesthesia for the operative delivery of the baby.
<i>Collection methods:</i>	More than one agent or technique can be recorded, except

where 1=none applies.

This item should only be recorded for the operative delivery of the baby and not third stage labour e.g. removal of placenta.

Comments:

Anaesthetic use may influence the duration of labour, may affect the health status of the baby at birth and is an indicator of obstetric intervention.

Source and reference attributes

Submitting organisation:

National Perinatal Data Development Committee

Relational attributes

Related metadata references:

Supersedes Birth event – anaesthesia administered, code N NHIG, Superseded 07/12/2005

Analgesia administered for labour

Identifying and definitional attributes

<i>Metadata item type:</i>	Data Element
<i>Technical name:</i>	Birth event – analgesia administered, code N
<i>METeOR identifier:</i>	292546
<i>Registration status:</i>	NHIG, Standard 07/12/2005
<i>Definition:</i>	Analgesia administered to the woman to relieve pain for labour, as represented by a code.

Data element concept attributes

<i>Data element concept:</i>	Birth event – analgesia administered
<i>Definition:</i>	Analgesia administered to the woman to relieve pain for labour.
<i>Context:</i>	Perinatal statistics
<i>Object class:</i>	Birth event
<i>Property:</i>	Analgesia administered

Value domain attributes

Representational attributes

<i>Representation class:</i>	Code																		
<i>Data type:</i>	Number																		
<i>Format:</i>	N																		
<i>Maximum character length:</i>	1																		
<i>Permissible values:</i>	<table><thead><tr><th>Value</th><th>Meaning</th></tr></thead><tbody><tr><td>1</td><td>None</td></tr><tr><td>2</td><td>Nitrous oxide</td></tr><tr><td>4</td><td>Epidural or caudal</td></tr><tr><td>5</td><td>Spinal</td></tr><tr><td>6</td><td>Systemic opioids</td></tr><tr><td>7</td><td>Combined spinal-epidural</td></tr><tr><td>8</td><td>Other</td></tr><tr><td>9</td><td>Not stated/inadequately described</td></tr></tbody></table>	Value	Meaning	1	None	2	Nitrous oxide	4	Epidural or caudal	5	Spinal	6	Systemic opioids	7	Combined spinal-epidural	8	Other	9	Not stated/inadequately described
Value	Meaning																		
1	None																		
2	Nitrous oxide																		
4	Epidural or caudal																		
5	Spinal																		
6	Systemic opioids																		
7	Combined spinal-epidural																		
8	Other																		
9	Not stated/inadequately described																		
<i>Supplementary values:</i>																			

Collection and usage attributes

<i>Guide for use:</i>	
<i>Comments:</i>	Note: Code 3, which had a meaning in previous versions of the data standard is no longer used. As is good practice, the code will not be reused.

Data element attributes

Collection and usage attributes

<i>Guide for use:</i>	Systemic opioids include both intra-muscular and intravenous opioids. Code 7: this code is used when this technique has been selected
-----------------------	--

for the administration of analgesia for labour.

Collection methods: More than one agent or technique can be recorded, except where 1=none applies.
This item is to be recorded for first and second stage labour, but not third stage labour e.g. removal of placenta.

Comments: Analgesia use may influence the duration of labour, may affect the health status of the baby at birth and is an indicator of obstetric intervention.

Source and reference attributes

Submitting organisation: National Perinatal Data Development Committee

Relational attributes

Related metadata references: Supersedes Birth event – analgesia administered, code N NHIG, Superseded 07/12/2005

Angiotensin converting enzyme (ACE) inhibitors therapy status

Identifying and definitional attributes

<i>Metadata item type:</i>	Data Element
<i>Technical name:</i>	Person – angiotensin converting enzyme inhibitors therapy status, code NN
<i>METeOR identifier:</i>	284751
<i>Registration status:</i>	NHIG, Standard 04/06/2004
<i>Definition:</i>	The person's ACE inhibitor therapy status, as represented by a code.

Data element concept attributes

<i>Data element concept:</i>	Person – angiotensin converting enzyme inhibitors therapy status
<i>Definition:</i>	Identifies the person's ACE inhibitor therapy status.
<i>Context:</i>	Health care and clinical settings.
<i>Object class:</i>	Person
<i>Property:</i>	Angiotensin converting enzyme inhibitors therapy status

Value domain attributes

Representational attributes

<i>Representation class:</i>	Code																						
<i>Data type:</i>	Number																						
<i>Format:</i>	NN																						
<i>Maximum character length:</i>	2																						
<i>Permissible values:</i>	<table><thead><tr><th>Value</th><th>Meaning</th></tr></thead><tbody><tr><td>10</td><td>Given</td></tr><tr><td>21</td><td>Not given - patient refusal</td></tr><tr><td>22</td><td>Not given - allergy or intolerance (e.g. cough) to ACE inhibitors</td></tr><tr><td>23</td><td>Not given - moderate to severe aortic stenosis</td></tr><tr><td>24</td><td>Not given - bilateral renal artery stenosis</td></tr><tr><td>25</td><td>Not given - history of angio-oedema, hives, or rash in response to ACE inhibitors</td></tr><tr><td>26</td><td>Not given - hyperkalaemia</td></tr><tr><td>27</td><td>Not given - symptomatic hypotension</td></tr><tr><td>28</td><td>Not given - severe renal dysfunction</td></tr><tr><td>29</td><td>Not given - other</td></tr></tbody></table>	Value	Meaning	10	Given	21	Not given - patient refusal	22	Not given - allergy or intolerance (e.g. cough) to ACE inhibitors	23	Not given - moderate to severe aortic stenosis	24	Not given - bilateral renal artery stenosis	25	Not given - history of angio-oedema, hives, or rash in response to ACE inhibitors	26	Not given - hyperkalaemia	27	Not given - symptomatic hypotension	28	Not given - severe renal dysfunction	29	Not given - other
Value	Meaning																						
10	Given																						
21	Not given - patient refusal																						
22	Not given - allergy or intolerance (e.g. cough) to ACE inhibitors																						
23	Not given - moderate to severe aortic stenosis																						
24	Not given - bilateral renal artery stenosis																						
25	Not given - history of angio-oedema, hives, or rash in response to ACE inhibitors																						
26	Not given - hyperkalaemia																						
27	Not given - symptomatic hypotension																						
28	Not given - severe renal dysfunction																						
29	Not given - other																						
<i>Supplementary values:</i>	90 Not stated/inadequately described																						

Collection and usage attributes

<i>Guide for use:</i>	CODES 21 - 29 Not given If recording 'Not given', record the principal reason if more than one code applies.
-----------------------	---

Source and reference attributes

Submitting organisation: Australian Institute of Health and Welfare

Data element attributes

Source and reference attributes

Submitting organisation: Acute coronary syndrome data working group

Steward: The National Heart Foundation of Australia and The Cardiac Society of Australia and New Zealand

Relational attributes

Related metadata references: Supersedes Angiotensin converting enzyme (ACE) inhibitors therapy status, version 1, DE, NHDD, NHIMG, Superseded 01/03/2005

Implementation in Data Set Specifications: Acute coronary syndrome (clinical) DSS NHIG, Standard 07/12/2005

Implementation start date: 07/12/2005

Information specific to this data set:

For Acute coronary syndrome (ACS) reporting, can be collected at any time point during the management of the current event (i.e. at the time of triage, at times during the admission, or at the time of discharge).

Acute coronary syndrome (clinical) DSS NHIG, Superseded 07/12/2005

Information specific to this data set:

For Acute coronary syndrome (ACS) reporting, can be collected at any time point during the management of the current event (i.e. at the time of triage, at times during the admission, or at the time of discharge).

Anticipated patient election status

Identifying and definitional attributes

<i>Metadata item type:</i>	Data Element
<i>Technical name:</i>	Elective surgery waiting list episode – anticipated accommodation status, code N
<i>METeOR identifier:</i>	270074
<i>Registration status:</i>	NHIG, Standard 01/03/2005
<i>Definition:</i>	Accommodation chargeable status nominated by the patient when placed on an elective surgery waiting list, as represented by a code.

Data element concept attributes

<i>Data element concept:</i>	Elective surgery waiting list episode – anticipated accommodation status
<i>Definition:</i>	Accommodation chargeable status nominated by the patient when placed on an elective surgery waiting list.
<i>Context:</i>	Elective surgery waiting times
<i>Object class:</i>	Elective surgery waiting list episode

Value domain attributes

Representational attributes

<i>Representation class:</i>	Code						
<i>Data type:</i>	Number						
<i>Format:</i>	N						
<i>Maximum character length:</i>	1						
<i>Permissible values:</i>	<table><thead><tr><th>Value</th><th>Meaning</th></tr></thead><tbody><tr><td>1</td><td>Public</td></tr><tr><td>2</td><td>Private</td></tr></tbody></table>	Value	Meaning	1	Public	2	Private
Value	Meaning						
1	Public						
2	Private						

Collection and usage attributes

<i>Guide for use:</i>	CODE 1 Public patient: An eligible person who receives or elects to receive a public hospital service free of charge. CODE 2 Private patient: An eligible person who elects to be treated as a private patient; and elects to be responsible for paying fees of the type referred to in clause 57 (clause 58 of the Northern Territory Agreement) of the Australian Health Care Agreements. Clause 57 states that 'Private patients and ineligible persons may be charged an amount for public hospital services as determined by the State'.
-----------------------	---

Data element attributes

Collection and usage attributes

<i>Guide for use:</i>	The election status nominated by the patient at the time of
-----------------------	---

being placed on an elective surgery waiting list, to be treated as either:

- a public patient; or
- a private patient

This item is independent of patient's hospital insurance status. The definitions of a public and private patient are those in the 1998-2003 Australian Health Care Agreements

Patients whose charges are to be met by the Department of Veterans' Affairs are regarded as private patients.

Comments:

Anticipated election status may be used for the management of elective surgery waiting lists, but the term is not defined under the 1998-2003 Australian Health Care Agreements. Under the Australian Health Care Agreements, patients are required to elect to be treated as a public or private patient, at the time of, or as soon as practicable after admission. Therefore, the anticipated patient election status is not binding on the patient and may vary from the election the patient makes on admission.

Relational attributes

Related metadata references:

Supersedes Anticipated patient election status, version 1, DE, NHDD, NHIMG, Superseded 01/03/2005

Apgar score at 1 minute

Identifying and definitional attributes

<i>Metadata item type:</i>	Data Element
<i>Technical name:</i>	Birth – Apgar score (at 1 minute), code NN
<i>METeOR identifier:</i>	289345
<i>Registration status:</i>	NHIG, Standard 07/12/2005
<i>Definition:</i>	Numerical score used to indicate the baby's condition at 1 minute after birth.

Data element concept attributes

<i>Data element concept:</i>	Birth – Apgar score
<i>Definition:</i>	Numerical score used to indicate the baby's condition after birth.
<i>Context:</i>	Perinatal statistics
<i>Object class:</i>	Birth
<i>Property:</i>	Apgar score

Value domain attributes

Representational attributes

<i>Representation class:</i>	Code						
<i>Data type:</i>	String						
<i>Format:</i>	NN						
<i>Maximum character length:</i>	2						
<i>Permissible values:</i>	<table><thead><tr><th>Value</th><th>Meaning</th></tr></thead><tbody><tr><td>00-10</td><td>Apgar score</td></tr><tr><td>99</td><td>Not stated/inadequately described</td></tr></tbody></table>	Value	Meaning	00-10	Apgar score	99	Not stated/inadequately described
Value	Meaning						
00-10	Apgar score						
99	Not stated/inadequately described						
<i>Supplementary values:</i>	99 Not stated/inadequately described						

Collection and usage attributes

<i>Guide for use:</i>	The score is based on the five characteristics of heart rate, respiratory condition, muscle tone, reflexes and colour. The maximum or best score being 10.
-----------------------	--

Data element attributes

Collection and usage attributes

<i>Comments:</i>	Required to analyse pregnancy outcome, particularly after complications of pregnancy, labour and birth. The Apgar score is an indicator of the health of a baby.
------------------	--

Source and reference attributes

<i>Submitting organisation:</i>	National Perinatal Data Development Committee
---------------------------------	---

Relational attributes

<i>Related metadata references:</i>	Supersedes Birth – Apgar score (at 1 minute), code NN NHIG, Superseded 07/12/2005
-------------------------------------	---

Apgar score at 5 minutes

Identifying and definitional attributes

<i>Metadata item type:</i>	Data Element
<i>Technical name:</i>	Birth – Apgar score (at 5 minutes), code NN
<i>METeOR identifier:</i>	289360
<i>Registration status:</i>	NHIG, Standard 07/12/2005
<i>Definition:</i>	Numerical score used to indicate the baby's condition at 5 minutes after birth.

Data element concept attributes

<i>Data element concept:</i>	Birth – Apgar score
<i>Definition:</i>	Numerical score used to indicate the baby's condition after birth.
<i>Context:</i>	Perinatal statistics
<i>Object class:</i>	Birth
<i>Property:</i>	Apgar score

Value domain attributes

Representational attributes

<i>Representation class:</i>	Code						
<i>Data type:</i>	String						
<i>Format:</i>	NN						
<i>Maximum character length:</i>	2						
<i>Permissible values:</i>	<table><thead><tr><th>Value</th><th>Meaning</th></tr></thead><tbody><tr><td>00-10</td><td>Apgar score</td></tr><tr><td>99</td><td>Not stated/inadequately described</td></tr></tbody></table>	Value	Meaning	00-10	Apgar score	99	Not stated/inadequately described
Value	Meaning						
00-10	Apgar score						
99	Not stated/inadequately described						
<i>Supplementary values:</i>	99 Not stated/inadequately described						

Collection and usage attributes

<i>Guide for use:</i>	The score is based on the five characteristics of heart rate, respiratory condition, muscle tone, reflexes and colour. The maximum or best score being 10.
-----------------------	--

Data element attributes

Collection and usage attributes

<i>Comments:</i>	Required to analyse pregnancy outcome, particularly after complications of pregnancy, labour and birth. The Apgar score is an indicator of the health of a baby.
------------------	--

Source and reference attributes

<i>Submitting organisation:</i>	National Perinatal Data Development Committee
---------------------------------	---

Relational attributes

<i>Related metadata references:</i>	Supersedes Birth – Apgar score (at 5 minutes), code NN NHIG, Superseded 07/12/2005
-------------------------------------	--

Implementation in Data Set Specifications:

Perinatal NMDS NHIG, Superseded 06/09/2006

Implementation start date: 01/07/2006

Implementation end date: 30/06/2007

Information specific to this data set:

Required to analyse pregnancy outcome, particularly after complications of pregnancy, labour and birth. The Apgar score is an indicator of the health of a baby.

Perinatal NMDS 2007-2008 NHIG, Standard 06/09/2006

Implementation start date: 01/07/2007

Information specific to this data set:

Required to analyse pregnancy outcome, particularly after complications of pregnancy, labour and birth. The Apgar score is an indicator of the health of a baby.

Area of usual residence

Identifying and definitional attributes

<i>Metadata item type:</i>	Data Element
<i>Technical name:</i>	Person – area of usual residence, geographical location code (ASGC 2006) NNNNN
<i>METeOR identifier:</i>	341800
<i>Registration status:</i>	NHIG, Standard 14/09/2006
<i>Definition:</i>	Geographical location of usual residence of the person, as represented by a code.

Data element concept attributes

<i>Data element concept:</i>	Person – area of usual residence
<i>Definition:</i>	Geographical location of usual residence of the person.
<i>Object class:</i>	Person
<i>Property:</i>	Area of usual residence

Value domain attributes

Representational attributes

<i>Classification scheme:</i>	Australian Standard Geographical Classification 2006
<i>Representation class:</i>	Code
<i>Data type:</i>	Number
<i>Format:</i>	NNNNN
<i>Maximum character length:</i>	5

Data element attributes

Collection and usage attributes

<i>Guide for use:</i>	<p>The geographical location is reported using a five digit numerical code. The first digit is the single-digit code to indicate State or Territory. The remaining four digits are the numerical code for the Statistical Local Area (SLA) within the State or Territory.</p> <p>The single digit codes for the states and territories and the four digit codes for the SLAs are as defined in the Australian Standard Geographical Classification (ASGC).</p> <p>The ASGC is updated on an annual basis with a date of effect of 1 July each year. Therefore, the edition effective for the data collection reference year should be used.</p> <p>The codes for SLA are unique within each State and Territory, but not within the whole country. Thus, to define a unique location, the code of the State or Territory is required in addition to the code for the SLA.</p> <p>The Australian Bureau of Statistics '(ABS) National Localities Index (NLI) (ABS Catalogue number 1252.0) can be used to assign each locality or address in Australia to a SLA. The NLI is a comprehensive list of localities in Australia with their full code (including State or Territory and SLA) from the main</p>
-----------------------	--

structure of the ASGC.

For the majority of localities, the locality name (suburb or town, for example) is sufficient to assign a SLA. However, some localities have the same name. For most of these, limited additional information such as the postcode or State can be used with the locality name to assign the SLA. In addition, other localities cross one or more SLA boundaries and are referred to as split localities. For these, the more detailed information of the number and street of the person's residence is used with the Streets Sub-index of the NLI to assign the SLA. If the information available on the person's address indicates that it is in a split locality but is insufficient to assign an SLA, the code for the SLA which includes most of the split locality should be reported. This is in accordance with the NLI assignment of SLA when a split locality is identified and further detail about the address is not available.

The NLI does not assign a SLA code if the information about the address is insufficient to identify a locality, or is not an Australian locality. In these cases, the appropriate codes for undefined SLA within Australia (State or Territory unstated), undefined SLA within a stated State or Territory, no fixed place of abode (within Australia or within a stated State or Territory) or overseas should be used.

Collection methods:

When collecting the geographical location of a person's usual place of residence, the Australian Bureau of Statistics (ABS) recommends that 'usual' be defined as: 'the place where the person has or intends to live for 6 months or more, or the place that the person regards as their main residence, or where the person has no other residence, the place they currently reside.' Apart from collecting a person's usual place of residence there is also a need in some collections to collect area of residence immediately prior to or after assistance is provided, or at some other point in time.

Comments:

Geographical location is reported using Statistical Local Area (SLA) to enable accurate aggregation of information to larger areas within the Australian Standard Geographical Classification (ASGC) (such as Statistical Subdivisions and Statistical Divisions) as well as detailed analysis at the SLA level. The use of SLA also allows analysis relating the data to information compiled by the Australian Bureau of Statistics on the demographic and other characteristics of the population of each SLA. Analyses facilitated by the inclusion of SLA information include:

- comparison of the use of services by persons residing in different geographical areas,
- characterisation of catchment areas and populations for establishments for planning purposes, and
- documentation of the provision of services to residents of States or Territories other than the State or Territory of the provider.

Source and reference attributes

Origin:

Health Data Standards Committee

Relational attributes

Related metadata references:

Supersedes Person – area of usual residence, geographical location code (ASGC 2005) NNNNN NHIG, Superseded 14/09/2006

Implementation in Data Set Specifications:

Admitted patient care NMDS 2007-2008 NHIG, Standard 29/11/2006

Implementation start date: 01/07/2007

Admitted patient mental health care NMDS 2007-2008 NHIG, Standard 23/10/2006

Implementation start date: 01/07/2007

Admitted patient palliative care NMDS 2007-08 NHIG, Standard 23/10/2006

Implementation start date: 01/07/2007

Community mental health care NMDS 2007-2008 NHIG, Standard 23/10/2006

Implementation start date: 01/07/2007

Perinatal NMDS 2007-2008 NHIG, Standard 06/09/2006

Implementation start date: 01/07/2007

Residential mental health care NMDS 2007-2008 NHIG, Standard 23/10/2006

Implementation start date: 01/07/2007

Aspirin therapy status

Identifying and definitional attributes

<i>Metadata item type:</i>	Data Element
<i>Technical name:</i>	Person – aspirin therapy status, code NN
<i>METeOR identifier:</i>	284785
<i>Registration status:</i>	NHIG, Standard 04/06/2004
<i>Definition:</i>	The person's aspirin therapy status, as represented by a code.

Data element concept attributes

<i>Data element concept:</i>	Person – aspirin therapy status
<i>Definition:</i>	Identifies the person's aspirin therapy status.
<i>Context:</i>	Health care and clinical settings.
<i>Object class:</i>	Person
<i>Property:</i>	Aspirin therapy status

Value domain attributes

Representational attributes

<i>Representation class:</i>	Code																
<i>Data type:</i>	Number																
<i>Format:</i>	NN																
<i>Maximum character length:</i>	2																
<i>Permissible values:</i>	<table><thead><tr><th>Value</th><th>Meaning</th></tr></thead><tbody><tr><td>10</td><td>Given</td></tr><tr><td>21</td><td>Not given - patient refusal</td></tr><tr><td>22</td><td>Not given - true allergy to aspirin</td></tr><tr><td>23</td><td>Not given - active bleeding</td></tr><tr><td>24</td><td>Not given - bleeding risk</td></tr><tr><td>29</td><td>Not given - other</td></tr><tr><td>90</td><td>Not stated/inadequately described</td></tr></tbody></table>	Value	Meaning	10	Given	21	Not given - patient refusal	22	Not given - true allergy to aspirin	23	Not given - active bleeding	24	Not given - bleeding risk	29	Not given - other	90	Not stated/inadequately described
Value	Meaning																
10	Given																
21	Not given - patient refusal																
22	Not given - true allergy to aspirin																
23	Not given - active bleeding																
24	Not given - bleeding risk																
29	Not given - other																
90	Not stated/inadequately described																
<i>Supplementary values:</i>	90																

Collection and usage attributes

<i>Guide for use:</i>	CODES 21 - 29 Not given If recording 'Not given', record the principal reason if more than one code applies.
-----------------------	---

Source and reference attributes

<i>Submitting organisation:</i>	Australian Institute of Health and Welfare
---------------------------------	--

Data element attributes

Source and reference attributes

<i>Submitting organisation:</i>	Acute coronary syndrome data working group
<i>Steward:</i>	The National Heart Foundation of Australia and The Cardiac Society of Australia and New Zealand

Relational attributes

Related metadata references:

Supersedes Aspirin therapy status, version 1, DE, NHDD, NHIMG, Superseded 01/03/2005

Implementation in Data Set Specifications:

Acute coronary syndrome (clinical) DSS NHIG, Standard 07/12/2005

Implementation start date: 07/12/2005

Information specific to this data set:

For Acute coronary syndrome (ACS) reporting, can be collected at any time point during the management of the current event (i.e. at the time of triage, at times during the admission, or at the time of discharge).

Acute coronary syndrome (clinical) DSS NHIG, Superseded 07/12/2005

Information specific to this data set:

For Acute coronary syndrome (ACS) reporting, can be collected at any time point during the management of the current event (i.e. at the time of triage, at times during the admission, or at the time of discharge).

Assistance with activities

Identifying and definitional attributes

<i>Metadata item type:</i>	Data Element
<i>Technical name:</i>	Person – need for assistance with activities in a life area, code N
<i>METeOR identifier:</i>	320213
<i>Registration status:</i>	NHIG, Standard 29/11/2006 NCSIMG, Standard 16/10/2006
<i>Definition:</i>	The level of help and/or supervision a person requires (or would require if the person currently helping/supervising was not available) to perform tasks and actions in a specified life area, as represented by a code.
<i>Context:</i>	Human functioning and disability

Data element concept attributes

<i>Data element concept:</i>	Person – need for assistance with activities in a life area
<i>Definition:</i>	The personal assistance and/or supervision a person needs to perform tasks and actions in a life area.
<i>Object class:</i>	Person
<i>Property:</i>	Need for assistance with activities in a life area

Value domain attributes

Representational attributes

<i>Classification scheme:</i>	International Classification of Functioning, Disability and Health 2001	
<i>Representation class:</i>	Code	
<i>Data type:</i>	Number	
<i>Format:</i>	N	
<i>Maximum character length:</i>	1	
<i>Permissible values:</i>	Value	Meaning
	0	Does not need help/supervision
	1	Sometimes needs help/supervision
	2	Always needs help/supervision
	3	Unable to do this task or action, even with assistance
<i>Supplementary values:</i>	8	Not specified
	9	Not applicable

Collection and usage attributes

<i>Guide for use:</i>	This metadata item contributes to the definition of the concept ' Disability ' and gives an indication of the experience of disability for a person. In the context of health, an activity is the execution of a task or action by an individual. Activity limitations are difficulties an individual may have in executing an activity. Activity limitation varies with the environment and is assessed in relation to a particular environment; the absence or presence
-----------------------	--

of assistance, including aids and equipment, is an aspect of the environment.

This value domain records the level of a person's need for help or supervision, in a specified domain, in their overall life. This means that the need for assistance may not be directly relevant to the health or community care service being provided.

Where a life area includes a range of examples, (e.g. domestic life includes cooking, cleaning and shopping), if a person requires assistance in any of the areas then the highest level of assistance should be recorded.

Where need for assistance varies markedly over time (e.g. episodic psychiatric conditions) please record the average level of assistance needed.

The presence of an activity limitation with a given domain is indicated by a non-zero response in this value domain. Activity is limited when an individual, in the context of a health condition, either has need for assistance in performing an activity in an expected manner, or cannot perform the activity at all.

CODE 0 is used when the person has no need for supervision or help and can undertake the activity independently.

CODE 1 is used when the person sometimes needs assistance to perform an activity.

CODE 2 is used when the person always needs assistance to undertake the activity and cannot do the activity without assistance.

CODE 3 is used when the person cannot do the activity even with assistance

CODE 8 is used when a person's need for assistance to undertake the activity is unknown or there is insufficient information to use codes 0-3.

CODE 9 is used where the need for help or supervision is due to the person's age. For example, Education for persons less than 5 years and work for persons less than 15 years.

Source and reference attributes

Submitting organisation: Australian Institute of Health and Welfare (AIHW) which is the Australian Collaborating Centre for the World Health Organization Family of International Classifications.

Origin: WHO 2001. ICF: International Classification of Functioning, Disability and Health. Geneva: WHO
AIHW 2003. ICF Australian User Guide Version 1.0. Canberra: AIHW

Reference documents: Further information on the ICF, including more detailed codes, can be found in the ICF itself and the ICF Australian User Guide (AIHW 2003), at the following websites:

- WHO ICF website
<http://www.who.int/classifications/icf/en/>
- Australian Collaborating Centre ICF website
<http://www.aihw.gov.au/disability/icf/index.html>

Data element attributes

Collection and usage attributes

Guide for use: This data element, in conjunction with Person – activities and participation life area, code (ICF 2001) AN[NNN], indicates a person's need for assistance in a given domain of activity.

Source and reference attributes

Submitting organisation: Australian Institute of Health and Welfare (AIHW) which is the Australian Collaborating Centre for the World Health Organization Family of International Classifications.

Relational attributes

Implementation in Data Set Specifications: Activities and Participation cluster NHIG, Standard 29/11/2006
NCSIMG, Standard 16/10/2006

Australian State/Territory identifier (establishment)

Identifying and definitional attributes

<i>Metadata item type:</i>	Data Element
<i>Technical name:</i>	Establishment – Australian state/territory identifier, code N
<i>METeOR identifier:</i>	269941
<i>Registration status:</i>	NHIG, Standard 01/03/2005
<i>Definition:</i>	An identifier of the Australian state or territory in which an establishment is located, as represented by a code.

Data element concept attributes

<i>Data element concept:</i>	Establishment – Australian state/territory identifier
<i>Definition:</i>	An identifier of the Australian state or territory in which an establishment is located.
<i>Object class:</i>	Establishment
<i>Property:</i>	Australian state/territory identifier

Value domain attributes

Representational attributes

<i>Representation class:</i>	Code																				
<i>Data type:</i>	Number																				
<i>Format:</i>	N																				
<i>Maximum character length:</i>	1																				
<i>Permissible values:</i>	<table><thead><tr><th>Value</th><th>Meaning</th></tr></thead><tbody><tr><td>1</td><td>New South Wales</td></tr><tr><td>2</td><td>Victoria</td></tr><tr><td>3</td><td>Queensland</td></tr><tr><td>4</td><td>South Australia</td></tr><tr><td>5</td><td>Western Australia</td></tr><tr><td>6</td><td>Tasmania</td></tr><tr><td>7</td><td>Northern Territory</td></tr><tr><td>8</td><td>Australian Capital Territory</td></tr><tr><td>9</td><td>Other territories (Cocos (Keeling) Islands, Christmas Island and Jervis Bay Territory)</td></tr></tbody></table>	Value	Meaning	1	New South Wales	2	Victoria	3	Queensland	4	South Australia	5	Western Australia	6	Tasmania	7	Northern Territory	8	Australian Capital Territory	9	Other territories (Cocos (Keeling) Islands, Christmas Island and Jervis Bay Territory)
Value	Meaning																				
1	New South Wales																				
2	Victoria																				
3	Queensland																				
4	South Australia																				
5	Western Australia																				
6	Tasmania																				
7	Northern Territory																				
8	Australian Capital Territory																				
9	Other territories (Cocos (Keeling) Islands, Christmas Island and Jervis Bay Territory)																				

Collection and usage attributes

<i>Guide for use:</i>	The order presented here is the standard for the Australian Bureau of Statistics (ABS). Other organisations (including the Australian Institute of Health and Welfare) publish data in state order based on population (that is, Western Australia before South Australia and Australian Capital Territory before Northern Territory).
-----------------------	--

Source and reference attributes

<i>Reference documents:</i>	Australian Bureau of Statistics 2005. Australian Standard Geographical Classification (ASGC). Cat. no. 1216.0. Canberra:
-----------------------------	--

Data element attributes

Collection and usage attributes

Guide for use: This metadata item applies to the location of the establishment and not to the patient's area of usual residence.

Source and reference attributes

Submitting organisation: Australian Institute of Health and Welfare
Origin: National Health Data Committee
National Community Services Data Committee

Relational attributes

Related metadata references: Supersedes Australian State/Territory identifier, version 4, DE, Int. NCSDD & NHDD, NCSIMG & NHIMG, Superseded 01/03/2005
Is used in the formation of Service delivery outlet – geographic location, code (ASGC 2006) NNNNN NHIG, Standard 14/09/2006
Is used in the formation of Establishment – geographical location, code (ASGC 2006) NNNNN NHIG, Standard 14/09/2006
Is used in the formation of Establishment – geographical location, code (ASGC 2005) NNNNN NHIG, Superseded 14/09/2006
Is used in the formation of Service delivery outlet – geographic location, code (ASGC 2005) NNNNN NHIG, Superseded 14/09/2006
Is used in the formation of Establishment – organisation identifier (Australian), NNX[X]NNNNN NHIG, Standard 01/03/2005
Is used in the formation of Service delivery outlet – geographic location, code (ASGC 2004) NNNNN NHIG, Superseded 21/03/2006

Implementation in Data Set Specifications:

Admitted patient care NMDS NHIG, Superseded 07/12/2005

Implementation start date: 01/07/2005

Implementation end date: 30/06/2006

Information specific to this data set:

This data element applies to the location of the establishment and not to the patient's area of usual residence.

Admitted patient care NMDS 2007-2008 NHIG, Standard 29/11/2006

Implementation start date: 01/07/2007

Information specific to this data set:

This data element applies to the location of the establishment and not to the patient's area of usual residence.

Community mental health care NMDS 2005-2006 NHIG,
Superseded 07/12/2005

Implementation start date: 01/07/2005

Implementation end date: 30/06/2006

Community mental health care NMDS 2006-2007 NHIG,
Superseded 23/10/2006

Implementation start date: 01/07/2006

Implementation end date: 30/06/2007

Community mental health care NMDS 2007-2008 NHIG,
Standard 23/10/2006

Implementation start date: 01/07/2007

Mental health establishments NMDS 2005-2006 NHIG,
Superseded 07/12/2005

Implementation start date: 01/07/2005

Mental health establishments NMDS 2005-2006 NHIG,
Superseded 21/03/2006

Implementation start date: 01/07/2005

Implementation end date: 30/06/2006

Mental health establishments NMDS 2006-2007 NHIG,
Superseded 23/10/2006

Implementation start date: 01/07/2006

Implementation end date: 30/06/2007

Mental health establishments NMDS 2007-2008 NHIG,
Standard 23/10/2006

Implementation start date: 01/07/2007

Residential mental health care NMDS 2005-2006 NHIG,
Superseded 07/12/2005

Implementation start date: 01/07/2005

Implementation end date: 30/06/2006

Residential mental health care NMDS 2006-2007 NHIG,
Superseded 23/10/2006

Implementation start date: 01/07/2006

Implementation end date: 30/06/2007

Residential mental health care NMDS 2007-2008 NHIG,
Standard 23/10/2006

Implementation start date: 01/07/2007

Australian state/territory identifier

Identifying and definitional attributes

<i>Metadata item type:</i>	Data Element
<i>Technical name:</i>	Person – Australian state/territory identifier, code N
<i>METeOR identifier:</i>	286919
<i>Registration status:</i>	NHIG, Standard 04/05/2005 NCSIMG, Standard 25/08/2005 NHDAMG, Standard 10/02/2006
<i>Definition:</i>	The Australian state or territory where a person can be located, as represented by a code.

Data element concept attributes

<i>Data element concept:</i>	Person – Australian state/territory identifier
<i>Definition:</i>	The Australian state or territory where a person can be located.
<i>Object class:</i>	Person
<i>Property:</i>	Australian state/territory identifier

Value domain attributes

Representational attributes

<i>Representation class:</i>	Code																				
<i>Data type:</i>	Number																				
<i>Format:</i>	N																				
<i>Maximum character length:</i>	1																				
<i>Permissible values:</i>	<table><thead><tr><th>Value</th><th>Meaning</th></tr></thead><tbody><tr><td>1</td><td>New South Wales</td></tr><tr><td>2</td><td>Victoria</td></tr><tr><td>3</td><td>Queensland</td></tr><tr><td>4</td><td>South Australia</td></tr><tr><td>5</td><td>Western Australia</td></tr><tr><td>6</td><td>Tasmania</td></tr><tr><td>7</td><td>Northern Territory</td></tr><tr><td>8</td><td>Australian Capital Territory</td></tr><tr><td>9</td><td>Other territories (Cocos (Keeling) Islands, Christmas Island and Jervis Bay Territory)</td></tr></tbody></table>	Value	Meaning	1	New South Wales	2	Victoria	3	Queensland	4	South Australia	5	Western Australia	6	Tasmania	7	Northern Territory	8	Australian Capital Territory	9	Other territories (Cocos (Keeling) Islands, Christmas Island and Jervis Bay Territory)
Value	Meaning																				
1	New South Wales																				
2	Victoria																				
3	Queensland																				
4	South Australia																				
5	Western Australia																				
6	Tasmania																				
7	Northern Territory																				
8	Australian Capital Territory																				
9	Other territories (Cocos (Keeling) Islands, Christmas Island and Jervis Bay Territory)																				

Collection and usage attributes

<i>Guide for use:</i>	The order presented here is the standard for the Australian Bureau of Statistics (ABS). Other organisations (including the Australian Institute of Health and Welfare) publish data in state order based on population (that is, Western Australia before South Australia and Australian Capital Territory before Northern Territory).
-----------------------	--

Source and reference attributes

<i>Reference documents:</i>	Australian Bureau of Statistics 2005. Australian Standard
-----------------------------	---

Data element attributes

Collection and usage attributes

Collection methods: Irrespective of how the information is coded, conversion of the codes to the ABS standard must be possible.

Source and reference attributes

Origin: Australian Bureau of Statistics 2004. Australian Standard Geographical Classification (ASGC) (Cat. no. 1216.0). Viewed 13 October 2005.

Reference documents: AS4846 Health Care Provider Identification, 2004, Sydney: Standards Australia
AS5017 Health Care Client Identification, 2004, Sydney: Standards Australia
In AS4846 and AS5017 alternative codes are presented. Refer to the current standard for more details.

Relational attributes

Related metadata references: See also Person (address) – Australian postcode, code (Postcode datafile) {NNNN} NHIG, Standard 04/05/2005, NCSIMG, Standard 25/08/2005, NHDAMG, Standard 10/02/2006

Implementation in Data Set Specifications: Health care client identification DSS NHIG, Standard 04/05/2005
NCSIMG, Standard 03/10/2006
Health care provider identification DSS NHIG, Standard 04/05/2005

Australian state/territory identifier (service provider organisation)

Identifying and definitional attributes

<i>Metadata item type:</i>	Data Element
<i>Technical name:</i>	Service provider organisation – Australian state/territory identifier, code N
<i>METeOR identifier:</i>	289083
<i>Registration status:</i>	NHIG, Standard 04/05/2005 NCSIMG, Standard 07/12/2005
<i>Definition:</i>	An identifier of the Australian state or territory where an organisation or agency can be located, as represented by a code.

Data element concept attributes

<i>Data element concept:</i>	Service provider organisation – Australian state/territory identifier
<i>Definition:</i>	An identifier of the Australian state or territory where an organisation or agency can be located.
<i>Context:</i>	This is a geographic indicator which is used for analysis of the distribution of agencies or establishments and services.
<i>Object class:</i>	Service provider organisation
<i>Property:</i>	Australian state/territory identifier

Value domain attributes

Representational attributes

<i>Representation class:</i>	Code																				
<i>Data type:</i>	Number																				
<i>Format:</i>	N																				
<i>Maximum character length:</i>	1																				
<i>Permissible values:</i>	<table><thead><tr><th>Value</th><th>Meaning</th></tr></thead><tbody><tr><td>1</td><td>New South Wales</td></tr><tr><td>2</td><td>Victoria</td></tr><tr><td>3</td><td>Queensland</td></tr><tr><td>4</td><td>South Australia</td></tr><tr><td>5</td><td>Western Australia</td></tr><tr><td>6</td><td>Tasmania</td></tr><tr><td>7</td><td>Northern Territory</td></tr><tr><td>8</td><td>Australian Capital Territory</td></tr><tr><td>9</td><td>Other territories (Cocos (Keeling) Islands, Christmas Island and Jervis Bay Territory)</td></tr></tbody></table>	Value	Meaning	1	New South Wales	2	Victoria	3	Queensland	4	South Australia	5	Western Australia	6	Tasmania	7	Northern Territory	8	Australian Capital Territory	9	Other territories (Cocos (Keeling) Islands, Christmas Island and Jervis Bay Territory)
Value	Meaning																				
1	New South Wales																				
2	Victoria																				
3	Queensland																				
4	South Australia																				
5	Western Australia																				
6	Tasmania																				
7	Northern Territory																				
8	Australian Capital Territory																				
9	Other territories (Cocos (Keeling) Islands, Christmas Island and Jervis Bay Territory)																				

Collection and usage attributes

<i>Guide for use:</i>	The order presented here is the standard for the Australian Bureau of Statistics (ABS). Other organisations (including the Australian Institute of Health and Welfare) publish data in state order based on population (that is, Western Australia before
-----------------------	---

South Australia and Australian Capital Territory before Northern Territory).

Source and reference attributes

Reference documents: Australian Bureau of Statistics 2005. Australian Standard Geographical Classification (ASGC). Cat. no. 1216.0. Canberra: ABS. Viewed on 30/09/2005

Data element attributes

Collection and usage attributes

Collection methods: Irrespective of how the information is coded, conversion of the codes to the ABS standard must be possible.

Source and reference attributes

Submitting organisation: Australian Institute of Health and Welfare

Origin: Health Data Standard Committee
National Community Services Data Committee

Reference documents: AS4846 Health Care Provider Identification, 2004, Sydney: Standards Australia
AS5017 Health Care Client Identification, 2002, Sydney: Standards Australia
In AS4846 and AS5017 alternative codes are presented. Refer to the current standard for more details.

Relational attributes

Implementation in Data Set Specifications: Health care provider identification DSS NHIG, Standard 04/05/2005

Information specific to this data set:

When used specifically in the collection of address information for a client, the following local implementation rules may be applied:
NULL may be used to signify an unknown address State;
and Code 0 may be used to signify an overseas address.

Behaviour-related risk factor intervention - purpose

Identifying and definitional attributes

<i>Metadata item type:</i>	Data Element
<i>Technical name:</i>	Episode of care – behaviour-related risk factor intervention purpose, code N
<i>METeOR identifier:</i>	270338
<i>Registration status:</i>	NHIG, Standard 01/03/2005
<i>Definition:</i>	The behaviour-related risk factor(s) associated with an intervention(s), as represented by a code.

Data element concept attributes

<i>Data element concept:</i>	Episode of care – behaviour-related risk factor intervention purpose
<i>Definition:</i>	The behaviour-related risk factor(s) associated with an intervention(s).
<i>Context:</i>	Public health, health care and clinical settings: The presence of one or more behaviour-related risk factors can be used to help determine the risk of future adverse health events and the development of chronic diseases.
<i>Object class:</i>	Episode of care
<i>Property:</i>	Behaviour-related risk factor intervention purpose

Value domain attributes

Representational attributes

<i>Representation class:</i>	Code														
<i>Data type:</i>	Number														
<i>Format:</i>	N														
<i>Maximum character length:</i>	1														
<i>Permissible values:</i>	<table><thead><tr><th>Value</th><th>Meaning</th></tr></thead><tbody><tr><td>1</td><td>Smoking</td></tr><tr><td>2</td><td>Nutrition</td></tr><tr><td>3</td><td>Alcohol misuse</td></tr><tr><td>4</td><td>Physical inactivity</td></tr><tr><td>8</td><td>Other</td></tr><tr><td>9</td><td>Not stated/inadequately described</td></tr></tbody></table>	Value	Meaning	1	Smoking	2	Nutrition	3	Alcohol misuse	4	Physical inactivity	8	Other	9	Not stated/inadequately described
Value	Meaning														
1	Smoking														
2	Nutrition														
3	Alcohol misuse														
4	Physical inactivity														
8	Other														
9	Not stated/inadequately described														
<i>Supplementary values:</i>															

Source and reference attributes

<i>Steward:</i>	Australian Bureau of Statistics (ABS)
-----------------	---------------------------------------

Data element attributes

Collection and usage attributes

<i>Guide for use:</i>	More than one code can be recorded.
-----------------------	-------------------------------------

Source and reference attributes

Submitting organisation:

Cardiovascular Data Working Group

Origin:

Smoking, Nutrition, Alcohol, Physical Activity (SNAP) Framework - Commonwealth Department of Health and Ageing - June 2001.
Australian Institute of Health and Welfare 2002. Chronic Diseases and associated risk factors in Australians, 2001; Canberra.

Relational attributes

Related metadata references:

Supersedes Behaviour-related risk factor intervention - purpose, version 1, DE, NHDD, NHIMG, Superseded 01/03/2005

Implementation in Data Set Specifications:

Cardiovascular disease (clinical) DSS NHIG, Superseded 15/02/2006

Information specific to this data set:

Behaviour-related risk factors include tobacco smoking, nutrition patterns that are high in saturated fats and excessive energy (calories / kilojoules) (National Heart Foundation of Australia - A review of the relationship between dietary fat and cardiovascular disease, AJND, 1999. 56 (Supp) S5-S22), alcohol misuse and physical inactivity.

The importance of behaviour-related risk factors in health has become increasingly relevant in recent times because chronic diseases have emerged as the principal threat to the health of Australians. Most of the chronic diseases have their roots in these risk-taking behaviours (Chronic Diseases and associated risk factors in Australians, 2001; AIHW 2002 Canberra).

Smoking, Nutrition, Alcohol, Physical Activity (SNAP) initiative:

SNAP Framework for General Practice is an initiative of the Joint Advisory Group (JAG) on General Practice and Population Health.

The lifestyle-related behavioural risk factors of smoking, poor nutrition (and associated overweight and obesity) and harmful and hazardous alcohol use and declining levels of physical activity have been identified as significant contributors to the burden of disease in Australia, and particularly towards the National Health Priority Areas (NHPAs) of diabetes, cardiovascular disease, some cancers, injury, mental health and asthma. The NHPAs represent about 70% of the burden of illness and injury in Australia. Substantial health gains could occur by public health interventions that address these contributory factors.

Around 86% of the Australian population attends a general practice at least once a year. There is therefore substantial opportunity for general practitioners to observe and influence the lifestyle risk behaviours of their patients. Many general practitioners already undertake risk factor management with their patients. There are also a number of initiatives within general practices, Divisions of General Practice, state/territory and Commonwealth Governments and peak non-government organisations

aimed at reducing disease related to these four behavioural risk factors. Within the health system, there is potential for greater collaboration and integration of approaches for influencing risk factor behaviour based on system-wide roll-out of evidence-based best practice interventions.

The aim of the SNAP initiative is to reduce the health and socioeconomic impact of smoking, poor nutrition, harmful and hazardous alcohol use and physical inactivity on patients and the community through a systematic approach to behavioural interventions in primary care. This will provide an opportunity to make better use of evidence-based interventions and to ensure adoption of best practice initiatives widely through general practice.

Cardiovascular disease (clinical) DSS NHIG, Standard 15/02/2006

Information specific to this data set:

Behaviour-related risk factors include tobacco smoking, nutrition patterns that are high in saturated fats and excessive energy (calories /kilojoules) (National Heart Foundation of Australia - A review of the relationship between dietary fat and cardiovascular disease, AJND, 1999. 56 (Supp) S5-S22), alcohol misuse and physical inactivity.

The importance of behaviour-related risk factors in health has become increasingly relevant in recent times because chronic diseases have emerged as the principal threat to the health of Australians. Most of the chronic diseases have their roots in these risk-taking behaviours (Chronic Diseases and associated risk factors in Australians, 2001; AIHW 2002 Canberra).

Smoking, Nutrition, Alcohol, Physical Activity (SNAP) initiative:

SNAP Framework for General Practice is an initiative of the Joint Advisory Group (JAG) on General Practice and Population Health.

The lifestyle-related behavioural risk factors of smoking, poor nutrition (and associated overweight and obesity) and harmful and hazardous alcohol use and declining levels of physical activity have been identified as significant contributors to the burden of disease in Australia, and particularly towards the National Health Priority Areas (NHPAs) of diabetes, cardiovascular disease, some cancers, injury, mental health and asthma. The NHPAs represent about 70% of the burden of illness and injury in Australia. Substantial health gains could occur by public health interventions that address these contributory factors.

Around 86% of the Australian population attends a general practice at least once a year. There is therefore substantial opportunity for general practitioners to observe and influence the lifestyle risk behaviours of their patients. Many general practitioners already undertake risk factor management with their patients. There are also

a number of initiatives within general practices, Divisions of General Practice, state/territory and Commonwealth Governments and peak non-government organisations aimed at reducing disease related to these four behavioural risk factors. Within the health system, there is potential for greater collaboration and integration of approaches for influencing risk factor behaviour based on system-wide roll-out of evidence-based best practice interventions.

The aim of the SNAP initiative is to reduce the health and socioeconomic impact of smoking, poor nutrition, harmful and hazardous alcohol use and physical inactivity on patients and the community through a systematic approach to behavioural interventions in primary care. This will provide an opportunity to make better use of evidence-based interventions and to ensure adoption of best practice initiatives widely through general practice.

Behaviour-related risk factor intervention purpose

Identifying and definitional attributes

<i>Metadata item type:</i>	Data Element
<i>Technical name:</i>	Episode of care – behaviour-related risk factor intervention, code NN
<i>METeOR identifier:</i>	270165
<i>Registration status:</i>	NHIG, Standard 01/03/2005
<i>Definition:</i>	The intervention taken to modify or manage the patient's behaviour-related risk factor(s), as represented by a code.

Data element concept attributes

<i>Data element concept:</i>	Episode of care – behaviour-related risk factor intervention
<i>Definition:</i>	The intervention taken to modify or manage the patient's behaviour-related risk factor(s).
<i>Context:</i>	Public health, health care and clinical settings: To enable analysis of the interventions within an episode of care, in relation to the outcome of this care, especially when linked to information on risk factors. The recording of Clinician's management interventions is critical information for health service monitoring, planning and patient outcomes. It is a major descriptor of the care provided throughout an episode of care.
<i>Object class:</i>	Episode of care
<i>Property:</i>	Behaviour-related risk factor intervention

Value domain attributes

Representational attributes

<i>Representation class:</i>	Code																				
<i>Data type:</i>	String																				
<i>Format:</i>	NN																				
<i>Maximum character length:</i>	2																				
<i>Permissible values:</i>	<table><thead><tr><th>Value</th><th>Meaning</th></tr></thead><tbody><tr><td>01</td><td>No intervention</td></tr><tr><td>02</td><td>Information and education (not including written regimen)</td></tr><tr><td>03</td><td>Counselling</td></tr><tr><td>04</td><td>Pharmacotherapy</td></tr><tr><td>05</td><td>Referral provided to a health professional</td></tr><tr><td>06</td><td>Referral to a community program, support group or service</td></tr><tr><td>07</td><td>Written regimen provided</td></tr><tr><td>08</td><td>Surgery</td></tr><tr><td>98</td><td>Other</td></tr></tbody></table>	Value	Meaning	01	No intervention	02	Information and education (not including written regimen)	03	Counselling	04	Pharmacotherapy	05	Referral provided to a health professional	06	Referral to a community program, support group or service	07	Written regimen provided	08	Surgery	98	Other
Value	Meaning																				
01	No intervention																				
02	Information and education (not including written regimen)																				
03	Counselling																				
04	Pharmacotherapy																				
05	Referral provided to a health professional																				
06	Referral to a community program, support group or service																				
07	Written regimen provided																				
08	Surgery																				
98	Other																				
<i>Supplementary values:</i>	99 Not stated/inadequately defined																				

Collection and usage attributes

<i>Guide for use:</i>	CODE 01 No intervention Refers to no intervention taken with regard to the behaviour-related risk factor intervention-purpose.
	CODE 02 Information and education (not including written regimen) Refers to where there is no treatment provided to the patient for a behaviour-related risk factor intervention-purpose other than information and education.
	CODE 03 Counselling Refers to any method of individual or group counselling directed towards the behaviour-related risk factor intervention-purpose. This code excludes counselling activities that are part of referral options as defined in code 05 and 06.
	CODE 04 Pharmacotherapy Refers to pharmacotherapies that are prescribed or recommended for the management of the behaviour-related risk factor intervention-purpose.
	CODE 05 Referral provided to a health professional Refers to a referral to a health professional who has the expertise to assist the patient manage the behaviour-related risk factor intervention-purpose.
	CODE 06 Referral to a community program, support group or service Refers to a referral to community program, support group or service that has the expertise and resources to assist the patient manage the behaviour-related risk factor intervention-purpose.
	CODE 07 Written regimen provided Refers to the provision of a written regimen (nutrition plan, exercise prescription, smoking contract) given to the patient to assist them with the management of the behaviour-related risk factor intervention-purpose.
	CODE 08 Surgery Refers to a surgical procedure undertaken to assist the patient with the management of the behaviour-related risk factor intervention-purpose.

Data element attributes

Collection and usage attributes

Guide for use: More than one code can be recorded.

Source and reference attributes

Submitting organisation: Cardiovascular Data Working Group

Relational attributes

Related metadata references: Supersedes Behaviour-related risk factor intervention, version 1, DE, NHDD, NHIMG, Superseded 01/03/2005

Implementation in Data Set Specifications: Cardiovascular disease (clinical) DSS NHIG, Superseded 15/02/2006

Cardiovascular disease (clinical) DSS NHIG, Standard 15/02/2006

Beta-blocker therapy status

Identifying and definitional attributes

<i>Metadata item type:</i>	Data Element
<i>Technical name:</i>	Person – beta-blocker therapy status, code NN
<i>METeOR identifier:</i>	284802
<i>Registration status:</i>	NHIG, Standard 04/06/2004
<i>Definition:</i>	The person's beta-blocker therapy status, as represented by a code.

Data element concept attributes

<i>Data element concept:</i>	Person – beta-blocker therapy status
<i>Definition:</i>	Identifies the person's beta-blocker therapy status.
<i>Context:</i>	Health care and clinical settings.
<i>Object class:</i>	Person
<i>Property:</i>	Beta-blocker therapy status

Value domain attributes

Representational attributes

<i>Representation class:</i>	Code																						
<i>Data type:</i>	Number																						
<i>Format:</i>	NN																						
<i>Maximum character length:</i>	2																						
<i>Permissible values:</i>	<table><thead><tr><th>Value</th><th>Meaning</th></tr></thead><tbody><tr><td>10</td><td>Given</td></tr><tr><td>21</td><td>Not given - patient refusal</td></tr><tr><td>22</td><td>Not given - allergy or history of intolerance</td></tr><tr><td>23</td><td>Not given - bradycardia (heart rate less than 50 beats per minute)</td></tr><tr><td>24</td><td>Not given - symptomatic acute heart failure</td></tr><tr><td>25</td><td>Not given - systolic blood pressure of less than 90 mmHg</td></tr><tr><td>26</td><td>Not given - PR interval greater than 0.24 seconds</td></tr><tr><td>27</td><td>Not given - second and third degree heart block or bifascicular heart block</td></tr><tr><td>28</td><td>Not given - asthma/airways hyper-reactivity</td></tr><tr><td>29</td><td>Not given - other</td></tr></tbody></table>	Value	Meaning	10	Given	21	Not given - patient refusal	22	Not given - allergy or history of intolerance	23	Not given - bradycardia (heart rate less than 50 beats per minute)	24	Not given - symptomatic acute heart failure	25	Not given - systolic blood pressure of less than 90 mmHg	26	Not given - PR interval greater than 0.24 seconds	27	Not given - second and third degree heart block or bifascicular heart block	28	Not given - asthma/airways hyper-reactivity	29	Not given - other
Value	Meaning																						
10	Given																						
21	Not given - patient refusal																						
22	Not given - allergy or history of intolerance																						
23	Not given - bradycardia (heart rate less than 50 beats per minute)																						
24	Not given - symptomatic acute heart failure																						
25	Not given - systolic blood pressure of less than 90 mmHg																						
26	Not given - PR interval greater than 0.24 seconds																						
27	Not given - second and third degree heart block or bifascicular heart block																						
28	Not given - asthma/airways hyper-reactivity																						
29	Not given - other																						
<i>Supplementary values:</i>	90 Not stated/inadequately described																						

Collection and usage attributes

<i>Guide for use:</i>	CODES 15 - 29 Not given If recording 'Not given', record the principal reason if more than one code applies.
-----------------------	---

Source and reference attributes

Submitting organisation: Australian Institute of Health and Welfare

Data element attributes

Source and reference attributes

Submitting organisation: Acute coronary syndrome data working group

Steward: The National Heart Foundation of Australia and The Cardiac Society of Australia and New Zealand

Relational attributes

Related metadata references: Supersedes Beta-blocker therapy status, version 1, DE, NHDD, NHIMG, Superseded 01/03/2005

Implementation in Data Set Specifications: Acute coronary syndrome (clinical) DSS NHIG, Standard 07/12/2005

Implementation start date: 07/12/2005

Information specific to this data set:

For Acute coronary syndrome (ACS) reporting, can be collected at any time point during the management of the current event (i.e. at the time of triage, at times during the admission, or at the time of discharge).

Acute coronary syndrome (clinical) DSS NHIG, Superseded 07/12/2005

Information specific to this data set:

For Acute coronary syndrome (ACS) reporting, can be collected at any time point during the management of the current event (i.e. at the time of triage, at times during the admission, or at the time of discharge).

Birth order

Identifying and definitional attributes

<i>Metadata item type:</i>	Data Element
<i>Technical name:</i>	Birth – birth order, code N
<i>METeOR identifier:</i>	269992
<i>Registration status:</i>	NHIG, Standard 01/03/2005
<i>Definition:</i>	The sequential order of each baby of a multiple birth, as represented by a code.

Data element concept attributes

<i>Data element concept:</i>	Birth – birth order
<i>Definition:</i>	The sequential order of each baby of a multiple birth.
<i>Object class:</i>	Birth
<i>Property:</i>	Birth order

Value domain attributes

Representational attributes

<i>Representation class:</i>	Code																		
<i>Data type:</i>	Number																		
<i>Format:</i>	N																		
<i>Maximum character length:</i>	1																		
<i>Permissible values:</i>	<table><thead><tr><th>Value</th><th>Meaning</th></tr></thead><tbody><tr><td>1</td><td>Singleton or first of a multiple birth</td></tr><tr><td>2</td><td>Second of a multiple birth</td></tr><tr><td>3</td><td>Third of a multiple birth</td></tr><tr><td>4</td><td>Fourth of a multiple birth</td></tr><tr><td>5</td><td>Fifth of a multiple birth</td></tr><tr><td>6</td><td>Sixth of a multiple birth</td></tr><tr><td>8</td><td>Other</td></tr><tr><td>9</td><td>Not stated</td></tr></tbody></table>	Value	Meaning	1	Singleton or first of a multiple birth	2	Second of a multiple birth	3	Third of a multiple birth	4	Fourth of a multiple birth	5	Fifth of a multiple birth	6	Sixth of a multiple birth	8	Other	9	Not stated
Value	Meaning																		
1	Singleton or first of a multiple birth																		
2	Second of a multiple birth																		
3	Third of a multiple birth																		
4	Fourth of a multiple birth																		
5	Fifth of a multiple birth																		
6	Sixth of a multiple birth																		
8	Other																		
9	Not stated																		
<i>Supplementary values:</i>																			

Data element attributes

Collection and usage attributes

<i>Guide for use:</i>	CODE 2 Second of a multiple birth Stillborns are counted such that, if twins were born, the first stillborn and the second live-born, the second twin would be recorded as code 2 Second of a multiple birth (and not code 1 Singleton or first of a multiple birth).
<i>Collection methods:</i>	This data should be collected routinely for persons aged 28 days or less.

Source and reference attributes

<i>Submitting organisation:</i>	National Perinatal Data Development Committee
---------------------------------	---

Relational attributes

Related metadata references:

Supersedes Birth order, version 2, DE, NHDD, NHIMG,
Superseded 01/03/2005

Implementation in Data Set Specifications:

Health care client identification NHIG, Superseded 04/05/2005

Implementation start date: 01/01/2003

Health care client identification DSS NHIG, Standard
04/05/2005

NCSIMG, Standard 03/10/2006

Perinatal NMDS NHIG, Superseded 07/12/2005

Implementation start date: 01/07/2005

Implementation end date: 30/06/2006

Perinatal NMDS NHIG, Superseded 06/09/2006

Implementation start date: 01/07/2006

Implementation end date: 30/06/2007

Perinatal NMDS 2007-2008 NHIG, Standard 06/09/2006

Implementation start date: 01/07/2007

Birth plurality

Identifying and definitional attributes

<i>Metadata item type:</i>	Data Element
<i>Technical name:</i>	Birth event – birth plurality, code N
<i>Synonymous names:</i>	Multiple birth
<i>METeOR identifier:</i>	269994
<i>Registration status:</i>	NHIG, Standard 01/03/2005
<i>Definition:</i>	The number of babies resulting from a single pregnancy, as represented by a code.

Data element concept attributes

<i>Data element concept:</i>	Birth event – birth plurality
<i>Definition:</i>	The number of babies resulting from a single pregnancy.
<i>Object class:</i>	Birth event
<i>Property:</i>	Birth plurality

Value domain attributes

Representational attributes

<i>Representation class:</i>	Code																
<i>Data type:</i>	Number																
<i>Format:</i>	N																
<i>Maximum character length:</i>	1																
<i>Permissible values:</i>	<table><thead><tr><th>Value</th><th>Meaning</th></tr></thead><tbody><tr><td>1</td><td>Singleton</td></tr><tr><td>2</td><td>Twins</td></tr><tr><td>3</td><td>Triplets</td></tr><tr><td>4</td><td>Quadruplets</td></tr><tr><td>5</td><td>Quintuplets</td></tr><tr><td>6</td><td>Sextuplets</td></tr><tr><td>8</td><td>Other</td></tr></tbody></table>	Value	Meaning	1	Singleton	2	Twins	3	Triplets	4	Quadruplets	5	Quintuplets	6	Sextuplets	8	Other
Value	Meaning																
1	Singleton																
2	Twins																
3	Triplets																
4	Quadruplets																
5	Quintuplets																
6	Sextuplets																
8	Other																
<i>Supplementary values:</i>	<table><tbody><tr><td>9</td><td>Not stated</td></tr></tbody></table>	9	Not stated														
9	Not stated																

Data element attributes

Collection and usage attributes

<i>Guide for use:</i>	Plurality of a pregnancy is determined by the number of live births or by the number of fetuses that remain in utero at 20 weeks gestation and that are subsequently born separately. In multiple pregnancies, or if gestational age is unknown, only live births of any birthweight or gestational age, or fetuses weighing 400 g or more, are taken into account in determining plurality. Fetuses aborted before 20 completed weeks or fetuses compressed in the placenta at 20 or more weeks are excluded.
<i>Collection methods:</i>	This data should be collected routinely for persons aged 28 days or less.

Source and reference attributes

Submitting organisation: National Perinatal Data Development Committee

Relational attributes

Related metadata references: Supersedes Birth plurality, version 1, DE, NHDD, NHIMG, Superseded 01/03/2005

Implementation in Data Set Specifications: Health care client identification NHIG, Superseded 04/05/2005
Implementation start date: 01/01/2003

Health care client identification DSS NHIG, Standard 04/05/2005

NCSIMG, Standard 03/10/2006

Perinatal NMDS NHIG, Superseded 07/12/2005

Implementation start date: 01/07/2005

Implementation end date: 30/06/2006

Perinatal NMDS NHIG, Superseded 06/09/2006

Implementation start date: 01/07/2006

Implementation end date: 30/06/2007

Perinatal NMDS 2007-2008 NHIG, Standard 06/09/2006

Implementation start date: 01/07/2007

Bleeding episode using TIMI criteria (status)

Identifying and definitional attributes

<i>Metadata item type:</i>	Data Element
<i>Technical name:</i>	Person – bleeding episode status, code N
<i>METeOR identifier:</i>	284812
<i>Registration status:</i>	NHIG, Standard 04/06/2004
<i>Definition:</i>	A person's episode of bleeding as described by the Thrombolysis In Myocardial Infarction (TIMI) criteria, as represented by a code.

Data element concept attributes

<i>Data element concept:</i>	Person – bleeding episode status
<i>Definition:</i>	A person's episode of bleeding as described by the Thrombolysis In Myocardial Infarction (TIMI) criteria.
<i>Context:</i>	Health care and clinical settings.
<i>Object class:</i>	Person
<i>Property:</i>	Bleeding episode status

Value domain attributes

Representational attributes

<i>Representation class:</i>	Code										
<i>Data type:</i>	Number										
<i>Format:</i>	N										
<i>Maximum character length:</i>	1										
<i>Permissible values:</i>	<table><thead><tr><th>Value</th><th>Meaning</th></tr></thead><tbody><tr><td>1</td><td>Major</td></tr><tr><td>2</td><td>Minor</td></tr><tr><td>3</td><td>Non TIMI bleeding</td></tr><tr><td>4</td><td>None</td></tr></tbody></table>	Value	Meaning	1	Major	2	Minor	3	Non TIMI bleeding	4	None
Value	Meaning										
1	Major										
2	Minor										
3	Non TIMI bleeding										
4	None										
<i>Supplementary values:</i>	9 Not stated/inadequately described										

Collection and usage attributes

<i>Guide for use:</i>	<p>Note in calculating the fall in haemoglobin or haematocrit, transfusion of whole blood or packed red blood cells is counted as 1g/dl (0.1g/l) haemoglobin or 3% absolute haematocrit.</p> <p>CODE 1 Major Overt clinical bleeding (or documented intracranial or retroperitoneal haemorrhage) associated with a drop in haemoglobin of greater than 5g/dl (0.5g/l) or a haematocrit of greater than 15% (absolute).</p> <p>CODE 2 Minor Overt clinical bleeding associated with a fall in haemoglobin of 3 or less than or equal to 5g/dl (0.5g/l) or a haematocrit of 9% to less than or equal to 15% (absolute).</p> <p>CODE 3 Non TIMI Bleeding Bleeding event that does not meet the major or minor</p>
-----------------------	--

definition.
CODE 4 None
No bleeding event

Source and reference attributes

Submitting organisation: Australian Institute of Health and Welfare

Data element attributes

Source and reference attributes

Submitting organisation: Acute coronary syndrome data working group
Steward: The National Heart Foundation of Australia and The Cardiac Society of Australia and New Zealand
Origin: Rao AK, Pratt C, Berke A, et al. Thrombolysis in Myocardial Infarction (TIMI) Trial, phase I: hemorrhagic manifestations and changes in plasma fibrinogen and the fibrinolytic system in patients with recombinant tissue plasminogen activator and streptokinase. J Am Coll Cardiol 1988; 11:1-11.

Relational attributes

Related metadata references: Supersedes Bleeding episode using TIMI criteria - status, version 1, DE, NHDD, NHIMG, Superseded 01/03/2005
Implementation in Data Set Specifications: Acute coronary syndrome (clinical) DSS NHIG, Standard 07/12/2005
Implementation start date: 07/12/2005
Information specific to this data set:

Can be collected at any time point during the management of the current event (i.e. at the time of triage, at times during the admission, or at the time of discharge).

Acute coronary syndrome (clinical) DSS NHIG, Superseded 07/12/2005
Information specific to this data set:

Can be collected at any time point during the management of the current event (i.e. at the time of triage, at times during the admission, or at the time of discharge).

Blindness (diabetes complication)

Identifying and definitional attributes

<i>Metadata item type:</i>	Data Element
<i>Technical name:</i>	Person – blindness, code N
<i>METeOR identifier:</i>	270065
<i>Registration status:</i>	NHIG, Standard 01/03/2005
<i>Definition:</i>	Whether the individual has become legally blind in either or both eyes, as represented by a code.

Data element concept attributes

<i>Data element concept:</i>	Person – blindness
<i>Definition:</i>	Whether the individual has become legally blind in either or both eyes.
<i>Context:</i>	Diabetes mellitus specific metadata item.
<i>Object class:</i>	Person
<i>Property:</i>	Blindness

Value domain attributes

Representational attributes

<i>Representation class:</i>	Code										
<i>Data type:</i>	Number										
<i>Format:</i>	N										
<i>Maximum character length:</i>	1										
<i>Permissible values:</i>	<table><thead><tr><th>Value</th><th>Meaning</th></tr></thead><tbody><tr><td>1</td><td>Blindness - (</td></tr><tr><td>2</td><td>Blindness - (</td></tr><tr><td>3</td><td>Blindness - (</td></tr><tr><td>4</td><td>No blindness</td></tr></tbody></table>	Value	Meaning	1	Blindness - (2	Blindness - (3	Blindness - (4	No blindness
Value	Meaning										
1	Blindness - (
2	Blindness - (
3	Blindness - (
4	No blindness										
<i>Supplementary values:</i>	9 Not stated/inadequately described										

Collection and usage attributes

<i>Guide for use:</i>	CODE 3 Blindness - (< 6/60) occurred in one eye within 12 months and in the other eye prior to the last 12 months Blindness can be diagnosed in one eye within 12 months even though it has been previously diagnosed on the other eye.
<i>Collection methods:</i>	Ask the individual if he/she has been diagnosed as legally blind (< 6/60) in both or either eye. If so record whether it has occurred within or prior to the last 12 months. Alternatively determine blindness from appropriate documentation obtained from an ophthalmologist or optometrist.

Data element attributes

Source and reference attributes

Submitting organisation:

National Diabetes Data Working Group

Origin:

National Diabetes Outcomes Quality Review Initiative (NDOQRIN) data dictionary.

Relational attributes

Related metadata references:

Supersedes Blindness - diabetes complication, version 1, DE, NHDD, NHIMG, Superseded 01/03/2005

Implementation in Data Set Specifications:

Diabetes (clinical) DSS NHIG, Superseded 21/09/2005

Information specific to this data set:

Patients with diabetes have an increased risk of developing several eye complications including retinopathy, cataract and glaucoma that lead to loss of vision.

Diabetic retinopathy is a leading cause of blindness. Retinopathy is characterised by proliferation of the retina's blood vessels, which may project into the vitreous, causing vitreous haemorrhage, proliferation of fibrous tissue and retinal detachment. It is often accompanied by microaneurysms and macular oedema, which can express as blurred vision. The prevalence of retinopathy increases with increasing duration of diabetes. In the early stage, retinopathy is asymptomatic. Up to 20% of people with diabetes Type 2 have retinopathy at the time of diagnosis of diabetes. The cumulative prevalence of proliferation diabetic retinopathy and macular oedema after 20 years of type 1 diabetes is about 40%. The Diabetic Retinopathy Study Group showed that panretinal photocoagulation reduces the risk of severe loss of vision by 50%.

Although diabetes retinopathy cannot totally be prevented, better control of blood sugar level slows the onset and progression of retinopathy (The Diabetes Control and Complications Trial - DCCT). Cataract and glaucoma are also associated diabetic eye problems that could lead to blindness.

Regular eye checkups are important for patients suffering from diabetes mellitus. This helps to early detect abnormalities and to avoid or postpone vision-threatening complications.

According to the NSW Principles of Care and Guidelines for the Clinical Management of Diabetes Mellitus, a comprehensive ophthalmological examination should be carried out:

- At diagnosis and then every 1-2 years for patients whose diabetes onset was at age 30 years or more.
- Within five years of diagnosis and then every 1-2 years for patients whose diabetes onset was at age less than 30 years.

If retinopathy is detected, review diabetes control and improve if necessary.

References:

Vision Australia, No 2, 1997/8; University of Melbourne.

The Diabetic Retinopathy Study Research Group.

Photocoagulation treatment of proliferative diabetic retinopathy.

Clinical application of Diabetic Retinopathy Study (DRS)

finding, DRS Report Number8. Ophthalmology. 1981; 88:583/600).

Diabetes Control and Complications Trial: DCCT New England Journal of Medicine, 329(14), September 30, 1993.

Diabetes (clinical) DSS NHIG, Standard 21/09/2005

Information specific to this data set:

Patients with diabetes have an increased risk of developing several eye complications including retinopathy, cataract and glaucoma that lead to loss of vision.

Diabetic retinopathy is a leading cause of blindness. Retinopathy is characterised by proliferation of the retina's blood vessels, which may project into the vitreous, causing vitreous haemorrhage, proliferation of fibrous tissue and retinal detachment. It is often accompanied by microaneurysms and macular oedema, which can express as blurred vision. The prevalence of retinopathy increases with increasing duration of diabetes. In the early stage, retinopathy is asymptomatic. Up to 20% of people with diabetes Type 2 have retinopathy at the time of diagnosis of diabetes. The cumulative prevalence of proliferative diabetic retinopathy and macular oedema after 20 years of type 1 diabetes is about 40%. The Diabetic Retinopathy Study Group showed that panretinal photocoagulation reduces the risk of severe loss of vision by 50%.

Although diabetes retinopathy cannot totally be prevented, better control of blood sugar level slows the onset and progression of retinopathy (The Diabetes Control and Complications Trial - DCCT). Cataract and glaucoma are also associated diabetic eye problems that could lead to blindness.

Regular eye checkups are important for patients suffering from diabetes mellitus. This helps to early detect abnormalities and to avoid or postpone vision-threatening complications.

According to the NSW Principles of Care and Guidelines for the Clinical Management of Diabetes Mellitus, a comprehensive ophthalmological examination should be carried out:

- At diagnosis and then every 1-2 years for patients whose diabetes onset was at age 30 years or more.
- Within five years of diagnosis and then every 1-2 years for patients whose diabetes onset was at age less than 30 years.

If retinopathy is detected, review diabetes control and improve if necessary.

References:

Vision Australia, No 2, 1997/8; University of Melbourne.

The Diabetic Retinopathy Study Research Group.

Photocoagulation treatment of proliferative diabetic retinopathy.

Clinical application of Diabetic Retinopathy Study (DRS)

finding, DRS Report Number8. Ophthalmology. 1981;

88:583/600).

Diabetes Control and Complications Trial: DCCT New England

Blood pressure—diastolic (measured)

Identifying and definitional attributes

<i>Metadata item type:</i>	Data Element
<i>Technical name:</i>	Person—blood pressure (diastolic) (measured), millimetres of mercury NN[N]
<i>METeOR identifier:</i>	270072
<i>Registration status:</i>	NHIG, Standard 01/03/2005
<i>Definition:</i>	The person's diastolic blood pressure , measured in millimetres of mercury (mmHg).

Data element concept attributes

<i>Data element concept:</i>	Person—blood pressure (diastolic)
<i>Definition:</i>	The person's diastolic blood pressure .
<i>Context:</i>	Public health, health care and clinical settings
<i>Object class:</i>	Person
<i>Property:</i>	Blood pressure

Value domain attributes

Representational attributes

<i>Representation class:</i>	Total				
<i>Data type:</i>	Number				
<i>Format:</i>	NN[N]				
<i>Maximum character length:</i>	3				
<i>Supplementary values:</i>	<table><thead><tr><th>Value</th><th>Meaning</th></tr></thead><tbody><tr><td>999</td><td>Not stated/inadequately described</td></tr></tbody></table>	Value	Meaning	999	Not stated/inadequately described
Value	Meaning				
999	Not stated/inadequately described				
<i>Unit of measure:</i>	Millimetre of mercury (mmHg)				

Data element attributes

Collection and usage attributes

<i>Guide for use:</i>	<p>The diastolic pressure is recorded as phase V Korotkoff (disappearance of sound) however phase IV Korotkoff (muffling of sound) is used if the sound continues towards zero but does not cease.</p> <p>If Blood pressure - diastolic is not collected or not able to be collected, code 999.</p>
<i>Collection methods:</i>	<p>Measurement protocol for resting blood pressure:</p> <p>The diastolic blood pressure is one component of a routine blood pressure measurement (i.e. systolic/diastolic) and reflects the minimum pressure to which the arteries are exposed.</p> <ul style="list-style-type: none">• The patient should be relaxed and seated, preferably for several minutes, (at least 5 minutes). Ideally, patients should not take caffeine-containing beverages or smoke for two hours before blood pressure is measured.• Ideally, patients should not exercise within half an hour of the measurement being taken (National Nutrition Survey

User's Guide).

- Use a mercury sphygmomanometer. All other sphygmomanometers should be calibrated regularly against mercury sphygmomanometers to ensure accuracy.
- Bladder length should be at least 80%, and width at least 40% of the circumference of the mid-upper arm. If the velcro on the cuff is not totally attached, the cuff is probably too small.
- Wrap cuff snugly around upper arm, with the centre of the bladder of the cuff positioned over the brachial artery and the lower border of the cuff about 2 cm above the bend of the elbow.
- Ensure cuff is at heart level, whatever the position of the patient.
- Palpate the radial pulse of the arm in which the blood pressure is being measured.
- Inflate cuff to the pressure at which the radial pulse disappears and note this value. Deflate cuff, wait 30 seconds, and then inflate cuff to 30 mm Hg above the pressure at which the radial pulse disappeared.
- Deflate the cuff at a rate of 2-3 mm Hg/beat (2-3 mm Hg/sec) or less.
- Recording the diastolic pressure use phase V Korotkoff (disappearance of sound). Use phase IV Korotkoff (muffling of sound) only if sound continues towards zero but does not cease. Wait 30 seconds before repeating the procedure in the same arm. Average the readings.
- If the first two readings differ by more than 4 mmHg diastolic or if initial readings are high, take several readings after five minutes of quiet rest.

Comments:

The pressure head is the height difference a pressure can raise a fluid's equilibrium level above the surface subjected to pressure. (Blood pressure is usually measured as a head of Mercury, and this is the unit of measure nominated for this metadata item.)

The current (2002) definition of hypertension is based on the level of blood pressure above which treatment is recommended, and this depends on the presence of other risk factors, e.g. age, diabetes etc. (NHF 1999 Guide to Management of Hypertension).

Source and reference attributes

Submitting organisation:

Cardiovascular Data Working Group
National Diabetes Data Working Group

Origin:

The National Heart Foundation Blood Pressure Advisory Committee's 'Guidelines for the Management of Hypertension - 1999' which are largely based on World Health Organization Recommendations. (Guidelines Subcommittee of the WHO-ISH: 1999 WHO-ISH guidelines for management of hypertension. J Hypertension 1999; 17:151-83).
Australian Bureau of Statistics 1998. National Nutrition Survey User's Guide 1995. Cat. No. 4801.0. Canberra: ABS. (p. 20).
National Diabetes Outcomes Quality Review Initiative (NDOQRIN) data dictionary.

Reference documents:

'Guidelines for the Management of Hypertension - 1999' largely based on World Health Organization Recommendations. (Guidelines Subcommittee of the WHO) J Hypertension 1999; 17: 151-83.).

Diabetes Control and Complications Trial: DCCT New England Journal of Medicine, 329(14), September 30, 1993.

UKPDS 38 Tight blood pressure control and risk of macrovascular and microvascular complications in type 2 diabetes: UK Prospective Diabetes Study Group. British Medical Journal (1998); 317: 703-713.

Relational attributes

Related metadata references:

Supersedes Blood pressure - diastolic measured, version 1, DE, NHDD, NHIMG, Superseded 01/03/2005

Implementation in Data Set Specifications:

Acute coronary syndrome (clinical) DSS NHIG, Standard 07/12/2005

Implementation start date: 07/12/2005

Acute coronary syndrome (clinical) DSS NHIG, Superseded 07/12/2005

Cardiovascular disease (clinical) DSS NHIG, Superseded 15/02/2006

Information specific to this data set:

In the primary care setting, blood pressure on both arms should be measured at the first visit, particularly if there is evidence of peripheral vascular disease.

Variation of up to 5 mm Hg in blood pressure between arms can be acceptable. In certain conditions (e.g. chronic aortic dissection, subclavian artery stenosis) all blood pressure recordings should be taken from the arm with the highest reading.

Measure sitting and standing blood pressures in elderly and diabetic patients or in other situations in which orthostatic hypotension might be suspected.

Measure and record heart rate and rhythm. Note: Atrial fibrillation in a patient with hypertension indicates increased risk of stroke.

In all patients, consideration should be given to obtaining blood pressure measurements outside the clinic setting either by self-measurement of blood pressure at home or by non-invasive ambulatory blood pressure monitoring. Target-organ damage and cardiovascular outcome relate more closely to blood pressures measured outside the clinic, particularly with ambulatory monitoring. An accurate, reliable machine and technique are essential if home blood pressure monitoring is to be used. In up to 30% of patients who are hypertensive in the clinic, blood pressure outside the clinic is within acceptable limits ('white coat' hypertension).

High blood pressure is a major risk factor for coronary heart disease, heart failure, stroke, and renal failure with the risk increasing along with the level of blood pressure (Ashwell 1997; DSHS 1994b; Whelton 1994; Kannel 1991). The higher the blood pressure, the higher the risk of both stroke and coronary heart disease. The dividing line

between normotension and hypertension is arbitrary. Both systolic and diastolic blood pressures are predictors of heart, stroke and vascular disease at all ages (Kannel 1991), although diastolic blood pressure is a weaker predictor of death due to coronary heart disease (Neaton & Wentworth 1992).

The risk of disease increases as the level of blood pressure increases. When blood pressure is lowered by 4-6 mm Hg over two to three years, it is estimated that the risk reduces by 14 per cent in patients with coronary heart disease and by 42 per cent in stroke patients (Collins et al 1990; Rose 1992.) When high blood pressure is controlled by medication, the risk of cardiovascular disease is reduced, but not to the levels of unaffected people.

In settings such as general practice where the monitoring of a person's health is ongoing and where a measure can change over time, the service contact date should be recorded.

Cardiovascular disease (clinical) DSS NHIG, Standard
15/02/2006

Information specific to this data set:

In the primary care setting, blood pressure on both arms should be measured at the first visit, particularly if there is evidence of peripheral vascular disease.

Variation of up to 5 mm Hg in blood pressure between arms can be acceptable. In certain conditions (e.g. chronic aortic dissection, subclavian artery stenosis) all blood pressure recordings should be taken from the arm with the highest reading.

Measure sitting and standing blood pressures in elderly and diabetic patients or in other situations in which orthostatic hypotension might be suspected.

Measure and record heart rate and rhythm. Note: Atrial fibrillation in a patient with hypertension indicates increased risk of stroke.

In all patients, consideration should be given to obtaining blood pressure measurements outside the clinic setting either by self-measurement of blood pressure at home or by non-invasive ambulatory blood pressure monitoring. Target-organ damage and cardiovascular outcome relate more closely to blood pressures measured outside the clinic, particularly with ambulatory monitoring. An accurate, reliable machine and technique are essential if home blood pressure monitoring is to be used. In up to 30% of patients who are hypertensive in the clinic, blood pressure outside the clinic is within acceptable limits ('white coat' hypertension).

High blood pressure is a major risk factor for coronary heart disease, heart failure, stroke, and renal failure with the risk increasing along with the level of blood pressure (Ashwell 1997; DSHS 1994b; Whelton 1994; Kannel 1991). The higher the blood pressure, the higher the risk of both stroke and coronary heart disease. The dividing line between normotension and hypertension is arbitrary. Both systolic and diastolic blood pressures are predictors

of heart, stroke and vascular disease at all ages (Kannel 1991), although diastolic blood pressure is a weaker predictor of death due to coronary heart disease (Neaton & Wentworth 1992).

The risk of disease increases as the level of blood pressure increases. When blood pressure is lowered by 4-6 mm Hg over two to three years, it is estimated that the risk reduces by 14 per cent in patients with coronary heart disease and by 42 per cent in stroke patients (Collins et al 1990; Rose 1992.) When high blood pressure is controlled by medication, the risk of cardiovascular disease is reduced, but not to the levels of unaffected people.

In settings such as general practice where the monitoring of a person's health is ongoing and where a measure can change over time, the service contact date should be recorded.

Diabetes (clinical) DSS NHIG, Superseded 21/09/2005

Information specific to this data set:

The United Kingdom Prospective Diabetes Study (1987 to 1998) showed major benefit from lowering blood pressure in preventing diabetes complications.

A target for blood pressure for people who suffer from diabetes is 130/85 mm Hg or less; recommended by the Australian Diabetes Society (if proteinuria is detected it is less than 125/75 mm Hg) Australian Medicines Handbook: last modified February, 2001).

Following the NSW Principles of Care and Guidelines for the Clinical Management of Diabetes Mellitus for patients who suffer from hypertension, if pharmacological intervention is required, ACE inhibitors are the preferred agents for treating hypertension in people with diabetes (unless contraindicated).

Diabetes (clinical) DSS NHIG, Standard 21/09/2005

Information specific to this data set:

The United Kingdom Prospective Diabetes Study (1987 to 1998) showed major benefit from lowering blood pressure in preventing diabetes complications.

A target for blood pressure for people who suffer from diabetes is 130/85 mm Hg or less; recommended by the Australian Diabetes Society (if proteinuria is detected it is less than 125/75 mm Hg) Australian Medicines Handbook: last modified February, 2001).

Following the NSW Principles of Care and Guidelines for the Clinical Management of Diabetes Mellitus for patients who suffer from hypertension, if pharmacological intervention is required, ACE inhibitors are the preferred agents for treating hypertension in people with diabetes (unless contraindicated).

High blood pressure is a major risk factor for coronary heart disease, heart failure, stroke, and renal failure with the risk increasing along with the level of blood pressure (Ashwell 1997; DSHS 1994b; Whelton 1994; Kannel 1991).

Blood pressure—systolic (measured)

Identifying and definitional attributes

<i>Metadata item type:</i>	Data Element
<i>Technical name:</i>	Person—blood pressure (systolic) (measured), millimetres of mercury NN[N]
<i>METeOR identifier:</i>	270073
<i>Registration status:</i>	NHIG, Standard 01/03/2005
<i>Definition:</i>	The person's systolic blood pressure , measured in millimetres of mercury (mmHg).

Data element concept attributes

<i>Data element concept:</i>	Person—blood pressure (systolic)
<i>Definition:</i>	The person's systolic blood pressure .
<i>Context:</i>	Public health, health care and clinical settings
<i>Object class:</i>	Person
<i>Property:</i>	Blood pressure

Value domain attributes

Representational attributes

<i>Representation class:</i>	Total				
<i>Data type:</i>	Number				
<i>Format:</i>	NN[N]				
<i>Maximum character length:</i>	3				
<i>Supplementary values:</i>	<table><thead><tr><th>Value</th><th>Meaning</th></tr></thead><tbody><tr><td>999</td><td>Not stated/inadequately described</td></tr></tbody></table>	Value	Meaning	999	Not stated/inadequately described
Value	Meaning				
999	Not stated/inadequately described				
<i>Unit of measure:</i>	Millimetre of mercury (mmHg)				

Data element attributes

Collection and usage attributes

<i>Guide for use:</i>	For recording the systolic reading, use phase I Korotkoff (the first appearance of sound). If Blood pressure - systolic is not collected or not able to be collected, code 999.
<i>Collection methods:</i>	<p>Measurement protocol for resting blood pressure:</p> <p>The systolic blood pressure is one component of a routine blood pressure measurement (i.e. systolic/diastolic) and reflects the maximum pressure to which the arteries are exposed.</p> <ul style="list-style-type: none">• The patient should be relaxed and seated, preferably for several minutes, (at least 5 minutes). Ideally, patients should not take caffeine-containing beverages or smoke for two hours before blood pressure is measured.• Ideally, patients should not exercise within half an hour of the measurement being taken (National Nutrition Survey User's Guide).• Use a mercury sphygmomanometer. All other sphygmomanometers should be calibrated regularly

- against mercury sphygmomanometers to ensure accuracy.
- Bladder length should be at least 80%, and width at least 40% of the circumference of the mid-upper arm. If the Velcro on the cuff is not totally attached, the cuff is probably too small.
 - Wrap cuff snugly around upper arm, with the centre of the bladder of the cuff positioned over the brachial artery and the lower border of the cuff about 2 cm above the bend of the elbow.
 - Ensure cuff is at heart level, whatever the position of the patient.
 - Palpate the radial pulse of the arm in which the blood pressure is being measured.
 - Inflate cuff to the pressure at which the radial pulse disappears and note this value. Deflate cuff, wait 30 seconds, and then inflate cuff to 30 mm Hg above the pressure at which the radial pulse disappeared.
 - Deflate the cuff at a rate of 2-3 mm Hg/beat (2-3 mm Hg/sec) or less.
 - For recording the systolic reading, use phase I Korotkoff (the first appearance of sound). Wait 30 seconds before repeating the procedure in the same arm. Average the readings. If the first two readings differ by more than 6 mm Hg systolic or if initial readings are high, take several readings after five minutes of quiet rest.

Comments:

The pressure head is the height difference a pressure can raise a fluid's equilibrium level above the surface subjected to pressure. (Blood pressure is usually measured as a head of Mercury, and this is the unit of measure nominated for this metadata item.)

The current (2002) definition of hypertension is based on the level of blood pressure above which treatment is recommended, and this depends on the presence of other risk factors, e.g. age, diabetes etc. (NHF 1999 Guide to Management of Hypertension).

Source and reference attributes

Submitting organisation:

Cardiovascular Data Working Group
National Diabetes Data Working Group

Origin:

The National Heart Foundation Blood Pressure Advisory Committee's 'Guidelines for the Management of Hypertension - 1999' which are largely based on World Health Organization Recommendations. (Guidelines Subcommittee of the WHO-SH: 1999 WHO-ISH guidelines for management of hypertension. J Hypertension 1999; 17:151-83).

Australian Bureau of Statistics 1998. National Nutrition Survey User's Guide 1995. Cat. No. 4801.0. Canberra: ABS. (p. 20).

National Diabetes Outcomes Quality Review Initiative (NDOQRIN) data dictionary.

Reference documents:

'Guidelines for the Management of Hypertension - 1999' largely based on World Health Organization Recommendations. (Guidelines Subcommittee of the WHO) J Hypertension 1999; 17: 151-83.).

Diabetes Control and Complications Trial: DCCT New England

Journal of Medicine, 329(14), September 30, 1993.
UKPDS 38 Tight blood pressure control and risk of macrovascular and microvascular complications in type 2 diabetes: UK Prospective Diabetes Study Group. British Medical Journal (1998); 317: 703-713.

Relational attributes

Related metadata references:

Supersedes Blood pressure - systolic measured, version 1, DE, NHDD, NHIMG, Superseded 01/03/2005

Implementation in Data Set Specifications:

Acute coronary syndrome (clinical) DSS NHIG, Standard 07/12/2005

Implementation start date: 07/12/2005

Acute coronary syndrome (clinical) DSS NHIG, Superseded 07/12/2005

Cardiovascular disease (clinical) DSS NHIG, Superseded 15/02/2006

Information specific to this data set:

In the primary care setting, blood pressure on both arms should be measured at the first visit, particularly if there is evidence of peripheral vascular disease.

Variation of up to 5 mm Hg in blood pressure between arms can be acceptable. In certain conditions (e.g. chronic aortic dissection, subclavian artery stenosis) all blood pressure recordings should be taken from the arm with the highest reading.

Measure sitting and standing blood pressures in elderly and diabetic patients or in other situations in which orthostatic hypotension might be suspected.

Measure and record heart rate and rhythm. Note: Atrial fibrillation in a patient with hypertension indicates increased risk of stroke.

In all patients, consideration should be given to obtaining blood pressure measurements outside the clinic setting either by self-measurement of blood pressure at home or by non-invasive ambulatory blood pressure monitoring.

Target-organ damage and cardiovascular outcome relate more closely to blood pressures measured outside the clinic, particularly with ambulatory monitoring. An accurate, reliable machine and technique are essential if home blood pressure monitoring is to be used. In up to 30% of patients who are hypertensive in the clinic, blood pressure outside the clinic is within acceptable limits ('white coat' hypertension).

High blood pressure is a major risk factor for coronary heart disease, heart failure, stroke, and renal failure with the risk increasing along with the level of blood pressure (Ashwell 1997; DSHS 1994b; Whelton 1994; Kannel 1991). The higher the blood pressure, the higher the risk of both stroke and coronary heart disease. The dividing line between normotension and hypertension is arbitrary.

Both systolic and diastolic blood pressures are predictors of heart, stroke and vascular disease at all ages (Kannel 1991), although diastolic blood pressure is a weaker predictor of death due to coronary heart disease (Neaton &

Wentworth 1992).

The risk of disease increases as the level of blood pressure increases. When blood pressure is lowered by 4-6 mm Hg over two to three years, it is estimated that the risk reduces by 14 per cent in patients with coronary heart disease and by 42 per cent in stroke patients (Collins et al 1990; Rose 1992.) When high blood pressure is controlled by medication, the risk of cardiovascular disease is reduced, but not to the levels of unaffected people.

In settings such as general practice where the monitoring of a person's health is ongoing and where a measure can change over time, the service contact date should be recorded.

Cardiovascular disease (clinical) DSS NHIG, Standard
15/02/2006

Information specific to this data set:

In the primary care setting, blood pressure on both arms should be measured at the first visit, particularly if there is evidence of peripheral vascular disease.

Variation of up to 5 mm Hg in blood pressure between arms can be acceptable. In certain conditions (e.g. chronic aortic dissection, subclavian artery stenosis) all blood pressure recordings should be taken from the arm with the highest reading.

Measure sitting and standing blood pressures in elderly and diabetic patients or in other situations in which orthostatic hypotension might be suspected.

Measure and record heart rate and rhythm. Note: Atrial fibrillation in a patient with hypertension indicates increased risk of stroke.

In all patients, consideration should be given to obtaining blood pressure measurements outside the clinic setting either by self-measurement of blood pressure at home or by non-invasive ambulatory blood pressure monitoring.

Target-organ damage and cardiovascular outcome relate more closely to blood pressures measured outside the clinic, particularly with ambulatory monitoring. An accurate, reliable machine and technique are essential if home blood pressure monitoring is to be used. In up to 30% of patients who are hypertensive in the clinic, blood pressure outside the clinic is within acceptable limits ('white coat' hypertension).

High blood pressure is a major risk factor for coronary heart disease, heart failure, stroke, and renal failure with the risk increasing along with the level of blood pressure (Ashwell 1997; DSHS 1994b; Whelton 1994; Kannel 1991). The higher the blood pressure, the higher the risk of both stroke and coronary heart disease. The dividing line between normotension and hypertension is arbitrary.

Both systolic and diastolic blood pressures are predictors of heart, stroke and vascular disease at all ages (Kannel 1991), although diastolic blood pressure is a weaker predictor of death due to coronary heart disease (Neaton & Wentworth 1992).

The risk of disease increases as the level of blood pressure

increases. When blood pressure is lowered by 4-6 mm Hg over two to three years, it is estimated that the risk reduces by 14 per cent in patients with coronary heart disease and by 42 per cent in stroke patients (Collins et al 1990; Rose 1992.) When high blood pressure is controlled by medication, the risk of cardiovascular disease is reduced, but not to the levels of unaffected people.

In settings such as general practice where the monitoring of a person's health is ongoing and where a measure can change over time, the service contact date should be recorded.

Diabetes (clinical) DSS NHIG, Superseded 21/09/2005

Information specific to this data set:

The United Kingdom Prospective Diabetes Study (1987 to 1998) showed major benefit from lowering blood pressure in preventing diabetes complications.

A target for blood pressure for people who suffer from diabetes is 130/85 mm Hg or less; recommended by the Australian Diabetes Society (if proteinuria is detected it is less than 125/75 mm Hg) Australian Medicines Handbook: last modified February, 2001).

Following the NSW Principles of Care and Guidelines for the Clinical Management of Diabetes Mellitus for patients who suffer from hypertension, if pharmacological intervention is required, ACE inhibitors are the preferred agents for treating hypertension in people with diabetes (unless contraindicated).

Diabetes (clinical) DSS NHIG, Standard 21/09/2005

Information specific to this data set:

The United Kingdom Prospective Diabetes Study (1987 to 1998) showed major benefit from lowering blood pressure in preventing diabetes complications.

A target for blood pressure for people who suffer from diabetes is 130/85 mm Hg or less; recommended by the Australian Diabetes Society (if proteinuria is detected it is less than 125/75 mm Hg) Australian Medicines Handbook: last modified February, 2001).

Following the NSW Principles of Care and Guidelines for the Clinical Management of Diabetes Mellitus for patients who suffer from hypertension, if pharmacological intervention is required, ACE inhibitors are the preferred agents for treating hypertension in people with diabetes (unless contraindicated).

High blood pressure is a major risk factor for coronary heart disease, heart failure, stroke, and renal failure with the risk increasing along with the level of blood pressure (Ashwell 1997; DSHS 1994b; Whelton 1994; Kannel 1991).

Bodily location of main injury

Identifying and definitional attributes

<i>Metadata item type:</i>	Data Element
<i>Technical name:</i>	Person – bodily location of main injury, code NN
<i>METeOR identifier:</i>	268943
<i>Registration status:</i>	NHIG, Standard 01/03/2005
<i>Definition:</i>	The bodily location of the injury chiefly responsible for the attendance of the person at the health care facility, as represented by a code.

Data element concept attributes

<i>Data element concept:</i>	Person – bodily location of main injury
<i>Definition:</i>	The bodily location of the injury chiefly responsible for the attendance of the person at the health care facility.
<i>Context:</i>	Injury surveillance
<i>Object class:</i>	Person
<i>Property:</i>	Bodily location of main injury

Value domain attributes

Representational attributes

<i>Representation class:</i>	Code																																		
<i>Data type:</i>	String																																		
<i>Format:</i>	NN																																		
<i>Maximum character length:</i>	2																																		
<i>Permissible values:</i>	<table><thead><tr><th>Value</th><th>Meaning</th></tr></thead><tbody><tr><td>01</td><td>Head (excludes face)</td></tr><tr><td>02</td><td>Face (excludes eye)</td></tr><tr><td>03</td><td>Neck</td></tr><tr><td>04</td><td>Thorax</td></tr><tr><td>05</td><td>Abdomen</td></tr><tr><td>06</td><td>Lower back (includes loin)</td></tr><tr><td>07</td><td>Pelvis (includes perineum, anogenital area and buttocks)</td></tr><tr><td>08</td><td>Shoulder</td></tr><tr><td>09</td><td>Upper arm</td></tr><tr><td>10</td><td>Elbow</td></tr><tr><td>11</td><td>Forearm</td></tr><tr><td>12</td><td>Wrist</td></tr><tr><td>13</td><td>Hand (include fingers)</td></tr><tr><td>14</td><td>Hip</td></tr><tr><td>15</td><td>Thigh</td></tr><tr><td>16</td><td>Knee</td></tr></tbody></table>	Value	Meaning	01	Head (excludes face)	02	Face (excludes eye)	03	Neck	04	Thorax	05	Abdomen	06	Lower back (includes loin)	07	Pelvis (includes perineum, anogenital area and buttocks)	08	Shoulder	09	Upper arm	10	Elbow	11	Forearm	12	Wrist	13	Hand (include fingers)	14	Hip	15	Thigh	16	Knee
Value	Meaning																																		
01	Head (excludes face)																																		
02	Face (excludes eye)																																		
03	Neck																																		
04	Thorax																																		
05	Abdomen																																		
06	Lower back (includes loin)																																		
07	Pelvis (includes perineum, anogenital area and buttocks)																																		
08	Shoulder																																		
09	Upper arm																																		
10	Elbow																																		
11	Forearm																																		
12	Wrist																																		
13	Hand (include fingers)																																		
14	Hip																																		
15	Thigh																																		
16	Knee																																		

17	Lower leg
18	Ankle
19	Foot (include toes)
20	Unspecified bodily location
21	Multiple injuries (involving more than one bodily location)
22	Bodily location not required

Data element attributes

Collection and usage attributes

Guide for use:

If the full International Classification of Diseases - Tenth Revision - Australian Modification 3rd Edition 2002 (ICD-10-AM 3rd edition) code is used to code the injury, this metadata item is not required (see metadata items Episode of care – principal diagnosis, code (ICD-10-AM 3rd edn) ANN{.N[N]} and Episode of care – additional diagnosis, code (ICD-10-AM 3rd edn) ANN{.N[N]}).

If any code from 01 to 12 or 26 to 29 in the metadata item Non-admitted patient service event – nature of main injury, code NN{.N} has been selected, the body region affected by that injury must be specified.

Select the category that best describes the location of the injury. If two or more categories are judged to be equally appropriate, select the one that comes first on the code list. A major injury, if present, should always be coded rather than a minor injury. If a major injury has been sustained (e.g. a fractured femur), along with one or more minor injuries (e.g. some small abrasions), the major injury should be coded in preference to coding 'multiple injuries'. As a general guide, an injury which, on its own, would be unlikely to have led to the attendance may be regarded as 'minor'. Bodily location of main injury is not required with other nature of main injury codes (code 22 may be used as a filler to indicate that a specific body region code is not required).

Comments:

The injury diagnosis is necessary for purposes including epidemiological research, casemix studies and planning. The nature of main injury together with the bodily location of the main injury indicates the diagnosis.

This metadata item is related to the ICD-10-AM (3rd edition) injury and poisoning classification. However, coding to the full ICD-10-AM (3rd edition) injury and poisoning classification (see metadata item Episode of care – principal diagnosis, code (ICD-10-AM 3rd edn) ANN{.N[N]}) is not available in most settings where basic injury surveillance is undertaken. This metadata item, in combination with the metadata item Non-admitted patient service event – nature of main injury, code NN{.N}, is a practicable alternative. Data coded to the full ICD-10-AM (3rd edition) codes can be aggregated to match this item, facilitating data comparison. Further information on the national injury surveillance program can be obtained from the National Injury Surveillance Unit, Flinders University, Adelaide.

Source and reference attributes

Submitting organisation: National Injury Surveillance Unit, Flinders University, Adelaide
National Data Standards for Injury Surveillance Advisory Group

Relational attributes

Related metadata references: Supersedes Bodily location of main injury, version 1, DE, NHDD, NHIMG, Superseded 01/03/2005
See also Injury event – nature of main injury, non-admitted patient code NN{.N} NHIG, Standard 01/03/2005

Implementation in Data Set Specifications: Injury surveillance DSS NHIG, Standard 03/05/2006
Injury surveillance NMDS NHIG, Superseded 03/05/2006

Implementation start date: 01/07/2005

Implementation end date: 30/06/2006

Injury surveillance NMDS NHIG, Superseded 07/12/2005

Body function

Identifying and definitional attributes

<i>Metadata item type:</i>	Data Element
<i>Technical name:</i>	Person – body function, code (ICF 2001) AN[NNNN]
<i>Synonymous names:</i>	Body function code
<i>METeOR identifier:</i>	320141
<i>Registration status:</i>	NHIG, Standard 29/11/2006 NCSIMG, Standard 16/10/2006
<i>Definition:</i>	The physiological or psychological function of a person's body system, as represented by a code.

Data element concept attributes

<i>Data element concept:</i>	Person – body function
<i>Definition:</i>	The physiological or psychological function of a person's body system.
<i>Object class:</i>	Person
<i>Property:</i>	Body function

Value domain attributes

Representational attributes

<i>Classification scheme:</i>	International Classification of Functioning, Disability and Health 2001
<i>Representation class:</i>	Code
<i>Data type:</i>	String
<i>Format:</i>	AN[NNNN]
<i>Maximum character length:</i>	6

Collection and usage attributes

<i>Guide for use:</i>	<p>This metadata item contributes to the definition of the concept 'Disability' and gives an indication of the experience of disability for a person.</p> <p>Data can be collected at the three digit level in one chapter and at the chapter level in another. However it is only possible to collect data at a single level of the hierarchy in a single chapter to maintain mutual exclusivity. For example, it is not permitted to collect both Exercise tolerance functions (3 digit level) and 'fatiguability' (4-digit level) as the former includes the latter.</p> <p>The value domain below refers to the highest hierarchical level (ICF chapter level). Data collected at this level, in association with <i>Impairment extent code N</i> will use the codes as indicated.</p> <p>CODE b1 Mental functions CODE b2 Sensory functions and pain CODE b3 Voice and speech functions CODE b4 Functions of the cardiovascular, haematological, immunological and respiratory systems CODE b5 Functions of the digestive, metabolic and the endocrine system CODE b6 Genitourinary and reproductive functions</p>
-----------------------	--

CODE b7 Neuromusculoskeletal and movement-related functions
CODE b8 Functions of the skin and related structures

Data collected at this level will provide a general description of the structures and can only be compared with data collected at the same level.

Each chapter contains categories at different levels ordered from general to detailed. For more detailed information the user should follow the structure of the ICF; the codes should be drawn from the same hierarchical level within any particular chapter. The full range of permissible values together, with definitions is listed in the *Body Functions* component of the ICF.

An example of a value domain at the 3 digit level from the Sensory functions and pain chapter may include:

CODE b210 Seeing functions
CODE b230 Hearing functions
CODE b235 Vestibular functions
CODE b250 Taste functions
CODE b255 Smell functions
CODE b260 Proprioceptive functions
CODE b265 Touch functions
CODE b270 Sensory functions related to temperature and other stimuli
CODE b279 Additional sensory functions, other specified and unspecified

An example of a value domain at the 4 digit level from the body function component may include:

CODE b1300 Energy level
CODE b1400 Sustaining attention
CODE b1442 Retrieval of memory
CODE b1521 Regulation of emotion
CODE b1641 Organization and planning

The prefix *b* denotes the domains within the component of *Body Functions*.

Source and reference attributes

<i>Submitting organisation:</i>	Australian Institute of Health and Welfare which is the Australian Collaborating Centre for the World Health Organization Family of International Classifications.
<i>Origin:</i>	WHO 2001. ICF: International Classification of Functioning, Disability and Health. Geneva: WHO AIHW 2003. ICF Australian User Guide Version 1.0. Canberra: AIHW
<i>Reference documents:</i>	Further information on the ICF, including more detailed codes, can be found in the ICF itself and the ICF Australian User Guide (AIHW 2003), at the following websites: <ul style="list-style-type: none">• WHO ICF website http://www.who.int/classifications/icf/en/• Australian Collaborating Centre ICF website http://www.aihw.gov.au/disability/icf/index.html

Data element attributes

Collection and usage attributes

Guide for use:

This data element can be used to record positive or neutral body function, as well as impairment of body function when used in conjunction with the metadata item Person—extent of impairment of body function, code (ICF 2001)N.

Where multiple body functions or impairments of body functions are recorded, the following prioritising system should be useful.

- The first recorded body function or impairment of body function is the one having the greatest impact on the individual.
- Second and subsequent body function or impairment of body function is also of relevance to the individual.

Source and reference attributes

Submitting organisation:

Australian Institute of Health and Welfare (AIHW) which is the Australian Collaborating Centre for the World Health Organization Family of International Classifications.

Relational attributes

Implementation in Data Set Specifications:

Body functions cluster NHIG, Standard 29/11/2006
NCSIMG, Standard 16/10/2006

Body mass index—adult (measured)

Identifying and definitional attributes

<i>Metadata item type:</i>	Data Element
<i>Technical name:</i>	Adult—body mass index (measured), ratio NN[N].N[N]
<i>METeOR identifier:</i>	270084
<i>Registration status:</i>	NHIG, Standard 01/03/2005
<i>Definition:</i>	A measure of an adult's weight (body mass) relative to height used to assess the extent of weight deficit or excess where height and weight have been measured.

Data element concept attributes

<i>Data element concept:</i>	Adult—body mass index
<i>Definition:</i>	Weight (body mass) relative to height used to assess the extent of weight deficit or excess in adults.
<i>Context:</i>	<p>Public health and health care:</p> <p>Body Mass Index (BMI) is used as an indicator of underweight, normal or healthy weight and overweight and obesity in adults and overweight and obesity in children and adolescents. On a population basis there is a strong association between BMI and health risk such as coronary heart disease, non-insulin dependant diabetes mellitus and high blood pressure in adults. In population based surveys, BMI may be used:</p> <ul style="list-style-type: none">• to indicate the prevalence of thinness and overweight and their sociodemographic distribution (problem identification)• to evaluate health promotion and disease prevention programs (assessment of interventions)• to monitor progress towards National public health policy• to ascertain determinants and consequences of thinness and overweight• in nutrition and physical activity surveillance and long-term planning.
<i>Object class:</i>	Adult
<i>Property:</i>	Body mass index

Value domain attributes

Representational attributes

<i>Representation class:</i>	Ratio						
<i>Data type:</i>	Number						
<i>Format:</i>	NN[N].N[N]						
<i>Maximum character length:</i>	5						
<i>Supplementary values:</i>	<table><thead><tr><th>Value</th><th>Meaning</th></tr></thead><tbody><tr><td>888.8</td><td>Unknown</td></tr><tr><td>999.9</td><td>Not reported</td></tr></tbody></table>	Value	Meaning	888.8	Unknown	999.9	Not reported
Value	Meaning						
888.8	Unknown						
999.9	Not reported						

Data element attributes

Collection and usage attributes

Guide for use:

Formula: BMI = weight (kg) divided by height (m) squared.

Body mass index is a continuous variable.

Code body mass index to one or two decimal places (i.e. 99.99 or 99.9). If any component necessary for its calculation (i.e. weight or height for adults) is unknown or has not been collected, code to 888.8, 999.9.

Collection methods:

NN.NN for BMI calculated from measured height and weight.

BMI should be derived after data entry of weight and height. It should be stored on the raw data set as a continuous variable and should not be aggregated or rounded.

Comments:

This metadata item applies to persons aged 2 years or older. It is recommended for use in population surveys and health care settings for adults and population surveys only for children and adolescents. It is recommended that calculated BMI for children and adolescents be compared with a suitable growth reference such as the United States Centers for Disease Control 2000 BMI-for-age chart be used for in health care settings such as hospitals, clinics and in general practice. A BMI greater than the 85th percentile would be classified as overweight, while a BMI greater than the 95th percentile would be classified as obese. These percentiles are arbitrary and do not relate to morbidity as the BMI cut-points do in adults.

BMI is relatively easy to determine, and has been validated against more direct measures of adiposity such as Magnetic Resonance Imaging and Dual X-ray Absorptiometry.

BMI is a low cost technique, with low respondent and investigator burden. In addition, it offers low inter-observer and intra-observer error, therefore offering good reliability.

Overweight and obesity, as defined by the World Health Organisation (WHO) for the interpretation of BMI (WHO 2000), are exceedingly common in Australia and their prevalence is increasing.

It is recommended that in population surveys, sociodemographic data including ethnicity should be collected, as well as other risk factors including physiological status (e.g. pregnancy), physical activity, smoking and alcohol consumption. Summary statistics may need to be adjusted for these variables.

National health metadata items currently exist for sex, date of birth, country of birth, Indigenous status and smoking.

Metadata items are being developed for physical activity.

Presentation of data:

Means, 95% confidence intervals, medians and centiles should be reported to one decimal place. Where the sample permits, population estimates should be presented by sex and 5-year age groups. Estimates based on sample surveys may need to take into account sampling weights.

For consistency with conventional practice, and for current comparability with international data sets, recommended centiles are 5, 10, 15, 25, 50, 75, 85, 90 and 95. To estimate the 5th and 95th centiles a sample size of at least 200 is recommended for each group for which the centiles are being specified.

Body mass index can be calculated from measured height and

weight, or self-reported height and weight, however for children and adolescents, self-reported or parentally reported data should be used cautiously if at all.

For adults, body mass index tends to be underestimated when based on self-reported, rather than measured, height and weight. This is due to the fact that, on average, height tends to be overestimated and weight tends to be underestimated when self-reported by respondents.

There are many individuals for whom BMI is an inappropriate measure of body fatness. These are individuals whose high body mass is due to excess muscle rather than fat (e.g. body builders or others in whom the level of physical activity promotes an increase in muscle mass); or in those with osteoporosis who will have a lower than usual BMI; or those who have a different body build (e.g. individuals with unusually long or short legs or a different body fat distribution) (WHO Expert Committee 1995).

This is particularly important when assessing individuals but should also be taken into account in interpreting data from populations in which there are sub-groups with genetic or environmental differences in body build, composition, skeletal proportions or body fat distribution. As such, both BMI and a measure of fat distribution (waist circumference or waist: hip ratio) are important in calculating the risk of obesity comorbidities.

Epidemiological research shows that there is a strong association between BMI and health risk. Excess adipose tissue in adults is associated with excess morbidity and mortality from conditions such as hypertension, unfavourable blood lipid concentrations, diabetes mellitus, coronary heart disease, some cancers, gall bladder disease, and osteoarthritis. It may also lead to social and economic disadvantage as well as psychosocial problems. It is a major public health issue in most industrialised societies.

Thinness (low BMI) is also an indicator of health risk, often being associated with general illness, anorexia, cigarette smoking, drug addiction and alcoholism. Low BMI is consistently associated with increased risk of osteoporosis and fractures in the elderly.

Source and reference attributes

Submitting organisation:

The Commonwealth Department of Health and Ageing based on the work of the consortium to develop an Australian standard definition of child/adolescent overweight and obesity; based at the Children Hospital at Westmead.

Origin:

Obesity: Preventing and Managing the Global Epidemic. Report of a WHO Consultation. 2000. World Health Organization.

Relational attributes

Related metadata references:

Is formed using Person – weight (measured), total kilograms
N[NN].N NHIG, Standard 01/03/2005

Is formed using Person – height (measured), total centimetres
NN[N].N NHIG, Standard 01/03/2005

Supersedes Body mass index, version 2, Derived DE, NHDD,
NHIMG, Superseded 01/03/2005

See also Person – body mass index (classification), code N[.N]

Body mass index—adult (self-reported)

Identifying and definitional attributes

<i>Metadata item type:</i>	Data Element
<i>Technical name:</i>	Adult—body mass index (self-reported), ratio NN[N].N[N]
<i>METeOR identifier:</i>	270086
<i>Registration status:</i>	NHIG, Standard 01/03/2005
<i>Definition:</i>	A measure of an adult's weight (body mass) relative to height used to assess the extent of weight deficit or excess where at least one of the measures is self reported.

Data element concept attributes

<i>Data element concept:</i>	Adult—body mass index
<i>Definition:</i>	Weight (body mass) relative to height used to assess the extent of weight deficit or excess in adults.
<i>Context:</i>	<p>Public health and health care:</p> <p>Body Mass Index (BMI) is used as an indicator of underweight, normal or healthy weight and overweight and obesity in adults and overweight and obesity in children and adolescents. On a population basis there is a strong association between BMI and health risk such as coronary heart disease, non-insulin dependant diabetes mellitus and high blood pressure in adults. In population based surveys, BMI may be used:</p> <ul style="list-style-type: none">• to indicate the prevalence of thinness and overweight and their sociodemographic distribution (problem identification)• to evaluate health promotion and disease prevention programs (assessment of interventions)• to monitor progress towards National public health policy• to ascertain determinants and consequences of thinness and overweight• in nutrition and physical activity surveillance and long-term planning.
<i>Object class:</i>	Adult
<i>Property:</i>	Body mass index

Value domain attributes

Representational attributes

<i>Representation class:</i>	Ratio						
<i>Data type:</i>	Number						
<i>Format:</i>	NN[N].N[N]						
<i>Maximum character length:</i>	5						
<i>Supplementary values:</i>	<table><thead><tr><th>Value</th><th>Meaning</th></tr></thead><tbody><tr><td>888.8</td><td>Unknown</td></tr><tr><td>999.9</td><td>Not reported</td></tr></tbody></table>	Value	Meaning	888.8	Unknown	999.9	Not reported
Value	Meaning						
888.8	Unknown						
999.9	Not reported						

Data element attributes

Collection and usage attributes

Collection methods:

NN.N for BMI calculated from either self-reported height and/or self-reported weight.

BMI calculated from measured height and weight should be distinguished from BMI calculated from self-reported height and/or weight. When either self-reported height or self-reported weight is used in the calculation, BMI should be recorded as self-reported BMI. Self-reported or parentally reported height and weight for children and adolescents should be used cautiously if at all.

BMI should be derived after the data entry of weight and height. It should be stored on the raw data set as a continuous variable and should not be aggregated or rounded.

Comments:

This metadata item applies to persons aged 2 years or older. It is recommended for use in population surveys and health care settings for adults and population surveys only for children and adolescents. It is recommended that calculated BMI for children and adolescents be compared with a suitable growth reference such as the United States Centers for Disease Control 2000 BMI-for-age chart be used for in health care settings such as hospitals, clinics and in general practice. A BMI greater than the 85th percentile would be classified as overweight, while a BMI greater than the 95th percentile would be classified as obese. These percentiles are arbitrary and do not relate to morbidity as the BMI cut-points do in adults.

BMI is relatively easy to determine, and has been validated against more direct measures of adiposity such as Magnetic Resonance Imaging and Dual X-ray Absorptiometry.

BMI is a low cost technique, with low respondent and investigator burden. In addition, it offers low inter-observer and intra-observer error, therefore offering good reliability.

Overweight and obesity, as defined by the World Health Organisation (WHO) for the interpretation of BMI (WHO 2000), are exceedingly common in Australia and their prevalence is increasing.

It is recommended that in population surveys, sociodemographic data including ethnicity should be collected, as well as other risk factors including physiological status (e.g. pregnancy), physical activity, smoking and alcohol consumption. Summary statistics may need to be adjusted for these variables.

National health metadata items currently exist for sex, date of birth, country of birth, Indigenous status and smoking.

Metadata items are being developed for physical activity.

Presentation of data:

Means, 95% confidence intervals, medians and centiles should be reported to one decimal place. Where the sample permits, population estimates should be presented by sex and 5-year age groups. Estimates based on sample surveys may need to take into account sampling weights.

For consistency with conventional practice, and for current comparability with international data sets, recommended centiles are 5, 10, 15, 25, 50, 75, 85, 90 and 95. To estimate the 5th and 95th centiles a sample size of at least 200 is recommended for each group for which the centiles are being specified.

Body mass index can be calculated from measured height and weight, or self-reported height and weight, however for children and adolescents, self-reported or parentally reported data should be used cautiously if at all.

For adults, body mass index tends to be underestimated when based on self-reported, rather than measured, height and weight. This is due to the fact that, on average, height tends to be overestimated and weight tends to be underestimated when self-reported by respondents.

There are many individuals for whom BMI is an inappropriate measure of body fatness. These are individuals whose high body mass is due to excess muscle rather than fat (e.g. body builders or others in whom the level of physical activity promotes an increase in muscle mass); or in those with osteoporosis who will have a lower than usual BMI; or those who have a different body build (e.g. individuals with unusually long or short legs or a different body fat distribution) (WHO Expert Committee 1995).

This is particularly important when assessing individuals but should also be taken into account in interpreting data from populations in which there are sub-groups with genetic or environmental differences in body build, composition, skeletal proportions or body fat distribution. As such, both BMI and a measure of fat distribution (waist circumference or waist: hip ratio) are important in calculating the risk of obesity comorbidities.

Epidemiological research shows that there is a strong association between BMI and health risk. Excess adipose tissue in adults is associated with excess morbidity and mortality from conditions such as hypertension, unfavourable blood lipid concentrations, diabetes mellitus, coronary heart disease, some cancers, gall bladder disease, and osteoarthritis. It may also lead to social and economic disadvantage as well as psychosocial problems. It is a major public health issue in most industrialised societies.

Thinness (low BMI) is also an indicator of health risk, often being associated with general illness, anorexia, cigarette smoking, drug addiction and alcoholism. Low BMI is consistently associated with increased risk of osteoporosis and fractures in the elderly.

Source and reference attributes

Submitting organisation:

The Commonwealth Department of Health and Ageing based on the work of the consortium to develop an Australian standard definition of child/adolescent overweight and obesity; based at the Children Hospital at Westmead.

Origin:

Obesity: Preventing and Managing the Global Epidemic. Report of a WHO Consultation. 2000. World Health Organization.

Relational attributes

Related metadata references:

See also Person – body mass index (classification), code N[.N] NHIG, Standard 01/03/2005

Is formed using Person – weight (measured), total kilograms N[NN].N NHIG, Standard 01/03/2005

Is formed using Person – height (self-reported), total centimetres NN[N] NHIG, Standard 01/03/2005

Is formed using Person – height (measured), total centimetres
NN[N].N NHIG, Standard 01/03/2005

Supersedes Body mass index, version 2, Derived DE, NHDD,
NHIMG, Superseded 01/03/2005

Is formed using Person – weight (self-reported), total kilograms
NN[N] NHIG, Standard 14/07/2005

Body mass index—child (measured)

Identifying and definitional attributes

<i>Metadata item type:</i>	Data Element
<i>Technical name:</i>	Child – body mass index (measured), ratio NN[N].N[N]
<i>METeOR identifier:</i>	270085
<i>Registration status:</i>	NHIG, Standard 01/03/2005
<i>Definition:</i>	A measure of a child's weight (body mass) relative to height used to assess the extent of weight excess where height and weight have been measured.

Data element concept attributes

<i>Data element concept:</i>	Child – body mass index
<i>Definition:</i>	Weight (body mass) relative to height used to assess the extent of weight excess in children and adolescents.
<i>Context:</i>	Public health and health care: Body Mass Index (BMI) is used as an indicator of underweight, normal or healthy weight and overweight and obesity in adults and overweight and obesity in children and adolescents. In population based surveys, BMI may be used: <ul style="list-style-type: none">• to indicate the prevalence of thinness and overweight and their sociodemographic distribution (problem identification)• to evaluate health promotion and disease prevention programs (assessment of interventions)• to monitor progress towards National public health policy• to ascertain determinants and consequences of thinness and overweight• in nutrition and physical activity surveillance and long-term planning.
<i>Object class:</i>	Child
<i>Property:</i>	Body mass index

Value domain attributes

Representational attributes

<i>Representation class:</i>	Ratio						
<i>Data type:</i>	Number						
<i>Format:</i>	NN[N].N[N]						
<i>Maximum character length:</i>	5						
<i>Supplementary values:</i>	<table><thead><tr><th>Value</th><th>Meaning</th></tr></thead><tbody><tr><td>888.8</td><td>Unknown</td></tr><tr><td>999.9</td><td>Not reported</td></tr></tbody></table>	Value	Meaning	888.8	Unknown	999.9	Not reported
Value	Meaning						
888.8	Unknown						
999.9	Not reported						

Data element attributes

Collection and usage attributes

<i>Collection methods:</i>	NN.NN for BMI calculated from measured height and weight.
----------------------------	---

BMI should be derived after the data entry of weight and height. It should be stored on the raw data set as a continuous variable and should not be aggregated or rounded.

Comments:

This metadata item applies to persons aged 2 years or older. It is recommended for use in population surveys and health care settings for adults and population surveys only for children and adolescents. It is recommended that calculated BMI for children and adolescents be compared with a suitable growth reference such as the United States Centers for Disease Control 2000 BMI-for-age chart be used for in health care settings such as hospitals, clinics and in general practice. A BMI greater than the 85th percentile would be classified as overweight, while a BMI greater than the 95th percentile would be classified as obese. These percentiles are arbitrary and do not relate to morbidity as the BMI cut-points do in adults.

BMI is relatively easy to determine, and has been validated against more direct measures of adiposity such as Magnetic Resonance Imaging and Dual X-ray Absorptiometry.

BMI is a low cost technique, with low respondent and investigator burden. In addition, it offers low inter-observer and intra-observer error, therefore offering good reliability.

Overweight and obesity, as defined by the World Health Organisation (WHO) for the interpretation of BMI (WHO 2000), are exceedingly common in Australia and their prevalence is increasing.

It is recommended that in population surveys, sociodemographic data including ethnicity should be collected, as well as other risk factors including physiological status (e.g. pregnancy), physical activity, smoking and alcohol consumption. Summary statistics may need to be adjusted for these variables.

National health metadata items currently exist for sex, date of birth, country of birth, Indigenous status and smoking. Metadata items are being developed for physical activity.

Presentation of data:

Means, 95% confidence intervals, medians and centiles should be reported to one decimal place. Where the sample permits, population estimates should be presented by sex and 5-year age groups. Estimates based on sample surveys may need to take into account sampling weights.

For consistency with conventional practice, and for current comparability with international data sets, recommended centiles are 5, 10, 15, 25, 50, 75, 85, 90 and 95. To estimate the 5th and 95th centiles a sample size of at least 200 is recommended for each group for which the centiles are being specified.

Body mass index can be calculated from measured height and weight, or self-reported height and weight, however for children and adolescents, self-reported or parentally reported data should be used cautiously if at all.

For adults, body mass index tends to be underestimated when based on self-reported, rather than measured, height and weight. This is due to the fact that, on average, height tends to be overestimated and weight tends to be underestimated when self-reported by respondents.

There are many individuals for whom BMI is an inappropriate measure of body fatness. These are individuals whose high

body mass is due to excess muscle rather than fat (e.g. body builders or others in whom the level of physical activity promotes an increase in muscle mass); or in those with osteoporosis who will have a lower than usual BMI; or those who have a different body build (e.g. individuals with unusually long or short legs or a different body fat distribution) (WHO Expert Committee 1995).

This is particularly important when assessing individuals but should also be taken into account in interpreting data from populations in which there are sub-groups with genetic or environmental differences in body build, composition, skeletal proportions or body fat distribution. As such, both BMI and a measure of fat distribution (waist circumference or waist: hip ratio) are important in calculating the risk of obesity comorbidities.

Epidemiological research shows that there is a strong association between BMI and health risk. Excess adipose tissue in adults is associated with excess morbidity and mortality from conditions such as hypertension, unfavourable blood lipid concentrations, diabetes mellitus, coronary heart disease, some cancers, gall bladder disease, and osteoarthritis. It may also lead to social and economic disadvantage as well as psychosocial problems. It is a major public health issue in most industrialised societies.

Thinness (low BMI) is also an indicator of health risk, often being associated with general illness, anorexia, cigarette smoking, drug addiction and alcoholism. Low BMI is consistently associated with increased risk of osteoporosis and fractures in the elderly.

Source and reference attributes

Submitting organisation:

The Commonwealth Department of Health and Ageing based on the work of the consortium to develop an Australian standard definition of child/adolescent overweight and obesity; based at the Children Hospital at Westmead.

Origin:

Obesity: Preventing and Managing the Global Epidemic. Report of a WHO Consultation. 2000. World Health Organization.
Cole TJ, Bellizzi MC, Flegal KM, Bietz WH. Establishing a standard definition for child overweight and obesity worldwide: international survey. *British Medical Journal* 2000; 320: 1240-1243

Relational attributes

Related metadata references:

See also Person – body mass index (classification), code N[.N] NHIG, Standard 01/03/2005

Supersedes Body mass index, version 2, Derived DE, NHDD, NHIMG, Superseded 01/03/2005

Is formed using Person – height (measured), total centimetres NN[N].N NHIG, Standard 01/03/2005

Is formed using Person – weight (measured), total kilograms N[NN].N NHIG, Standard 01/03/2005

Body mass index—child (self-reported)

Identifying and definitional attributes

<i>Metadata item type:</i>	Data Element
<i>Technical name:</i>	Child – body mass index (self-reported), ratio NN[N].N[N]
<i>METeOR identifier:</i>	270087
<i>Registration status:</i>	NHIG, Standard 01/03/2005
<i>Definition:</i>	A measure of a child's weight (body mass) relative to height used to assess the extent of weight excess where at least one of the measures is self reported.

Data element concept attributes

<i>Data element concept:</i>	Child – body mass index
<i>Definition:</i>	Weight (body mass) relative to height used to assess the extent of weight excess in children and adolescents.
<i>Context:</i>	Public health and health care: Body Mass Index (BMI) is used as an indicator of underweight, normal or healthy weight and overweight and obesity in adults and overweight and obesity in children and adolescents. In population based surveys, BMI may be used: <ul style="list-style-type: none">• to indicate the prevalence of thinness and overweight and their sociodemographic distribution (problem identification)• to evaluate health promotion and disease prevention programs (assessment of interventions)• to monitor progress towards National public health policy• to ascertain determinants and consequences of thinness and overweight• in nutrition and physical activity surveillance and long-term planning.
<i>Object class:</i>	Child
<i>Property:</i>	Body mass index

Value domain attributes

Representational attributes

<i>Representation class:</i>	Ratio						
<i>Data type:</i>	Number						
<i>Format:</i>	NN[N].N[N]						
<i>Maximum character length:</i>	5						
<i>Supplementary values:</i>	<table><thead><tr><th>Value</th><th>Meaning</th></tr></thead><tbody><tr><td>888.8</td><td>Unknown</td></tr><tr><td>999.9</td><td>Not reported</td></tr></tbody></table>	Value	Meaning	888.8	Unknown	999.9	Not reported
Value	Meaning						
888.8	Unknown						
999.9	Not reported						

Data element attributes

Collection and usage attributes

<i>Collection methods:</i>	NN.N for BMI calculated from either self-reported height
----------------------------	--

and/or self-reported weight.

BMI calculated from measured height and weight should be distinguished from BMI calculated from self-reported height and/or weight. When either self-reported height or self-reported weight is used in the calculation, BMI should be recorded as self-reported BMI. Self-reported or parentally reported height and weight for children and adolescents should be used cautiously if at all.

BMI should be derived after the data entry of weight and height. It should be stored on the raw data set as a continuous variable and should not be aggregated or rounded.

Comments:

This metadata item applies to persons aged 2 years or older. It is recommended for use in population surveys and health care settings for adults and population surveys only for children and adolescents. It is recommended that calculated BMI for children and adolescents be compared with a suitable growth reference such as the United States Centers for Disease Control 2000 BMI-for-age chart be used for in health care settings such as hospitals, clinics and in general practice. A BMI greater than the 85th percentile would be classified as overweight, while a BMI greater than the 95th percentile would be classified as obese. These percentiles are arbitrary and do not relate to morbidity as the BMI cut-points do in adults.

BMI is relatively easy to determine, and has been validated against more direct measures of adiposity such as Magnetic Resonance Imaging and Dual X-ray Absorptiometry.

BMI is a low cost technique, with low respondent and investigator burden. In addition, it offers low inter-observer and intra-observer error, therefore offering good reliability.

Overweight and obesity, as defined by the World Health Organisation (WHO) for the interpretation of BMI (WHO 2000), are exceedingly common in Australia and their prevalence is increasing.

It is recommended that in population surveys, sociodemographic data including ethnicity should be collected, as well as other risk factors including physiological status (e.g. pregnancy), physical activity, smoking and alcohol consumption. Summary statistics may need to be adjusted for these variables.

National health metadata items currently exist for sex, date of birth, country of birth, Indigenous status and smoking.

Metadata items are being developed for physical activity.

Presentation of data:

Means, 95% confidence intervals, medians and centiles should be reported to one decimal place. Where the sample permits, population estimates should be presented by sex and 5-year age groups. Estimates based on sample surveys may need to take into account sampling weights.

For consistency with conventional practice, and for current comparability with international data sets, recommended centiles are 5, 10, 15, 25, 50, 75, 85, 90 and 95. To estimate the 5th and 95th centiles a sample size of at least 200 is recommended for each group for which the centiles are being specified.

Body mass index can be calculated from measured height and weight, or self-reported height and weight, however for children and adolescents, self-reported or parentally reported

data should be used cautiously if at all.

For adults, body mass index tends to be underestimated when based on self-reported, rather than measured, height and weight. This is due to the fact that, on average, height tends to be overestimated and weight tends to be underestimated when self-reported by respondents.

There are many individuals for whom BMI is an inappropriate measure of body fatness. These are individuals whose high body mass is due to excess muscle rather than fat (e.g. body builders or others in whom the level of physical activity promotes an increase in muscle mass); or in those with osteoporosis who will have a lower than usual BMI; or those who have a different body build (e.g. individuals with unusually long or short legs or a different body fat distribution) (WHO Expert Committee 1995).

This is particularly important when assessing individuals but should also be taken into account in interpreting data from populations in which there are sub-groups with genetic or environmental differences in body build, composition, skeletal proportions or body fat distribution. As such, both BMI and a measure of fat distribution (waist circumference or waist: hip ratio) are important in calculating the risk of obesity comorbidities.

Epidemiological research shows that there is a strong association between BMI and health risk. Excess adipose tissue in adults is associated with excess morbidity and mortality from conditions such as hypertension, unfavourable blood lipid concentrations, diabetes mellitus, coronary heart disease, some cancers, gall bladder disease, and osteoarthritis. It may also lead to social and economic disadvantage as well as psychosocial problems. It is a major public health issue in most industrialised societies.

Thinness (low BMI) is also an indicator of health risk, often being associated with general illness, anorexia, cigarette smoking, drug addiction and alcoholism. Low BMI is consistently associated with increased risk of osteoporosis and fractures in the elderly.

Source and reference attributes

Submitting organisation:

The Commonwealth Department of Health and Ageing based on the work of the consortium to develop an Australian standard definition of child/adolescent overweight and obesity; based at the Children Hospital at Westmead.

Origin:

Obesity: Preventing and Managing the Global Epidemic. Report of a WHO Consultation. 2000. World Health Organization.

Cole TJ, Bellizzi MC, Flegal KM, Bietz WH. Establishing a standard definition for child overweight and obesity worldwide: international survey. *British Medical Journal* 2000; 320: 1240-1243

Relational attributes

Related metadata references:

Supersedes Body mass index, version 2, Derived DE, NHDD, NHIMG, Superseded 01/03/2005

Is formed using Person – height (measured), total centimetres NN[N].N NHIG, Standard 01/03/2005

Is formed using Person – height (self-reported), total

centimetres NN[N] NHIG, Standard 01/03/2005

Is formed using Person – weight (measured), total kilograms
N[NN].N NHIG, Standard 01/03/2005

See also Person – body mass index (classification), code N[.N]
NHIG, Standard 01/03/2005

Is formed using Person – weight (self-reported), total kilograms
NN[N] NHIG, Standard 14/07/2005

Body mass index—classification

Identifying and definitional attributes

<i>Metadata item type:</i>	Data Element
<i>Technical name:</i>	Person—body mass index (classification), code N[.N]
<i>METeOR identifier:</i>	270474
<i>Registration status:</i>	NHIG, Standard 01/03/2005

Data element concept attributes

<i>Data element concept:</i>	Person—body mass index (classification)
<i>Definition:</i>	The category of weight deficit or excess in adults and weight excess only in children and adolescents.
<i>Context:</i>	Public health and health care: BMI is used as an indicator of underweight, normal or healthy weight and overweight and obesity in adults and of overweight and obesity in children and adolescents. On a population basis there is a strong association between BMI and health risk.
<i>Object class:</i>	Person
<i>Property:</i>	Body mass index

Source and reference attributes

<i>Submitting organisation:</i>	World Health Organization (see also Comments) and the consortium to develop an Australian standard definition of child/adolescent overweight and obesity; at the Children's Hospital at Westmead on behalf of the Commonwealth Department of Health & Ageing
---------------------------------	--

Value domain attributes

Representational attributes

<i>Representation class:</i>	Code																						
<i>Data type:</i>	Number																						
<i>Format:</i>	N[.N]																						
<i>Maximum character length:</i>	2																						
<i>Permissible values:</i>	<table><thead><tr><th>Value</th><th>Meaning</th></tr></thead><tbody><tr><td>1</td><td>Not overweight or obese</td></tr><tr><td>1.1</td><td>Underweight</td></tr><tr><td>1.2</td><td>Normal range 18.50 - 24.99 Average</td></tr><tr><td>2</td><td>Overweight >= 25.00 Average</td></tr><tr><td>2.1</td><td>Overweight >= 25.0 Average</td></tr><tr><td>2.2</td><td>Pre Obese 25.00 - 29.99 Increased</td></tr><tr><td>3</td><td>Obese >= 30 Increased</td></tr><tr><td>3.1</td><td>Obese class 1 30.00 - 34.99 Moderate</td></tr><tr><td>3.2</td><td>Obese class 2 35.00 - 39.99 Severe</td></tr><tr><td>3.3</td><td>Obese class 3 >= 40.00 Very severe</td></tr></tbody></table>	Value	Meaning	1	Not overweight or obese	1.1	Underweight	1.2	Normal range 18.50 - 24.99 Average	2	Overweight >= 25.00 Average	2.1	Overweight >= 25.0 Average	2.2	Pre Obese 25.00 - 29.99 Increased	3	Obese >= 30 Increased	3.1	Obese class 1 30.00 - 34.99 Moderate	3.2	Obese class 2 35.00 - 39.99 Severe	3.3	Obese class 3 >= 40.00 Very severe
Value	Meaning																						
1	Not overweight or obese																						
1.1	Underweight																						
1.2	Normal range 18.50 - 24.99 Average																						
2	Overweight >= 25.00 Average																						
2.1	Overweight >= 25.0 Average																						
2.2	Pre Obese 25.00 - 29.99 Increased																						
3	Obese >= 30 Increased																						
3.1	Obese class 1 30.00 - 34.99 Moderate																						
3.2	Obese class 2 35.00 - 39.99 Severe																						
3.3	Obese class 3 >= 40.00 Very severe																						
<i>Supplementary values:</i>	9 Not stated/inadequately described																						

Collection and usage attributes

Guide for use:

Adults:

Body mass index for adults cannot be calculated if components necessary for its calculation (weight or height) is unknown or has not been collected (i.e is coded to 888.8 or 999.9).

BMI for adults is categorised according to the range it falls within as indicated by codes 1.1, 1.2, 2.1, 2.2, 3.1, 3.2, 3.3 or 9.9. For consistency, when the sample includes children and adolescents, adults can be analysed under the broader categories of 1,2,3 or 9 as used for categorising children and adolescents.

Children/adolescents:

Body mass index for children and adolescents aged 2 to 17 years cannot be calculated if components necessary for its calculation (date of birth, sex, weight or height) is unknown or has not been collected (i.e is coded to 888.8, 999.9 or 9).

Self-reported or parentally reported height and weight for children and adolescents should be used cautiously if at all.

To determine overweight and obesity in children and adolescents, compare the derived BMI against those recorded for the relevant age and sex of the subject to be classified, against Table 1: Classification of BMI for children and adolescents, based on BMI cut-points developed by Cole et al (see below). For example, an 11 year old boy with a BMI of 21 would be considered overweight (i.e coded as 2), or a 7 year old girl with a BMI of 17.5 would be considered not overweight or obese (i.e coded as 1).

Using this method, children and adolescents can only be coded as 1, 2, 3 or 9.

Collection methods:

Use N for BMI category determined (1,2,3 or 9) for persons (children and adolescents) aged 2 to 17 years.

Use N.N for BMI category determined (1.1,1.2,2.1,2.2,3.1,3.2,3.3 or 9.9) for persons aged 18 years or older.

Standard definitions of overweight and obesity in terms of BMI are used to derive age-specific and age-adjusted indicators of overweight and obesity for reporting progress towards National public health policy .

Data element attributes

Collection and usage attributes

Guide for use:

Age(years)	BMI equivalent to 25 kg/m ²		BMI equivalent to 30 kg/m ²	
	Males	Females	Males	Females
2	18.41	18.02	20.09	19.81
2.5	18.13	17.76	19.80	19.55
3	17.89	17.56	19.57	19.36
3.5	17.69	17.40	19.39	19.23
4	17.55	17.28	19.29	19.15

4.5	17.47	17.19	19.26	19.12
5	17.42	17.15	19.30	19.17
5.5	17.45	17.20	19.47	19.34
6	17.55	17.34	19.78	19.65
6.5	17.71	17.53	20.23	20.08
7	17.92	17.75	20.63	20.51
7.5	18.16	18.03	21.09	21.01
8	18.44	18.35	21.60	21.57
8.5	18.76	18.69	22.17	22.18
9	19.10	19.07	22.77	22.81
9.5	19.46	19.45	23.39	23.46
10	19.84	19.86	24.00	24.11
10.5	20.20	20.29	24.57	24.77
11	20.55	20.74	25.10	25.42
11.5	20.89	21.20	25.58	26.05
12	21.22	21.68	26.02	26.67
12.5	21.56	22.14	26.43	27.24
13	21.91	22.58	26.84	27.76
13.5	22.27	22.98	27.25	28.20
14	22.62	23.34	27.63	28.57
14.5	22.96	23.66	27.98	28.87
15	23.29	23.94	28.30	29.11
15.5	23.60	24.17	28.60	29.29
16	23.90	24.37	28.88	29.43
16.5	24.19	24.54	29.14	29.56
17	24.46	24.70	29.41	26.69
17.5	24.73	24.85	29.70	29.84
18	25.00	25.00	30.00	30.00

Comments:

This metadata item applies to persons aged 2 years or older. It is recommended for use in population surveys and health care settings for adults and population surveys only for children and adolescents. It is recommended that calculated BMI for children and adolescents be compared with a suitable growth reference such as the US Centers for Disease Control 2000 BMI- for-age chart in health care settings such as hospitals, clinics and in general practice. A BMI greater than the 85th percentile would be classified as overweight, while a BMI greater than the 95th percentile would be classified as obese. These percentiles are arbitrary and do not relate to morbidity as the BMI cut-points do in adults.

BMI can be considered to provide the most useful, albeit crude, population-level measure of obesity. The robust nature of the measurements and the widespread routine inclusion of weights and heights in clinical and population health surveys mean that a more selective measure of adiposity, such as skinfold thickness measurements, provides additional rather than primary information. BMI can be used to estimate the prevalence of obesity within a population and the risks

associated with it, but does not, however, account for the wide variation in the nature of obesity between different individuals and populations (WHO 2000).

BMI values for adults are age-independent and the same for both sexes.

However, BMI values for children and adolescents aged 2 to 17 years are age and sex specific and are classified by comparing against the above table, Table 1: Classification of BMI for children and adolescents.

For adults and children and adolescents BMI may not correspond to the same degree of fatness in different populations due, in part, to differences in body proportions. The classification table shows a simplistic relationship between BMI and the risk of comorbidity, which can be affected by a range of factors, including the nature of the diet, ethnic group and activity level. The risks associated with increasing BMI are continuous and graded and begin at a BMI of 25 (or equivalent to 25 for children and adolescents). The interpretation of BMI grades in relation to risk may differ for different populations. Both BMI and a measure of fat distribution (waist circumference or waist: hip ratio in adults) are important in calculating the risk of obesity comorbidities.

It is recommended that in population surveys, sociodemographic data including ethnicity should be collected, as well as other risk factors including physiological status (e.g. pregnancy), physical activity, smoking and alcohol consumption. Summary statistics may need to be adjusted for these variables.

National health metadata items currently exist for sex, date of birth, country of birth, Indigenous Status and smoking. Metadata items are being developed for physical activity.

Presentation of data:

Methods used to establish cut-off points for overweight have been arbitrary and, as a result, cut-off points vary between countries. The data are derived mainly from studies of mortality and morbidity risk performed in people living in western Europe or the United States of America, and cut-off points for BMI as an indicator of adiposity and risk in populations who differ in body build and genetic disposition are likely to vary.

Caution is required in relation to BMI cut-off points when used for different ethnic groups because of limited outcome data for some ethnic groups, e.g. Aboriginal and Torres Strait Islander peoples. As with overweight the cut-off points for a given level of risk are likely to vary with body build, genetic background and physical activity.

The classification above is different to ones that have been used in the past and it is important that in any trend analysis consistent definitions are used.

BMI should not be rounded before categorisation to the classification above.

Source and reference attributes

Submitting organisation:

World Health Organization (see also Comments) and the consortium to develop an Australian standard definition of child/adolescent overweight and obesity; at the Children's

Origin:

Hospital at Westmead on behalf of the Commonwealth
Department of Health & Ageing

Obesity: Preventing and Managing the Global Epidemic
(Report of a WHO Consultation: World Health Organization
2000);

Cole TJ, Bellizzi MC, Flegal KM, Dietz WH. Establishing a
standard definition for child overweight and obesity
worldwide: international survey. British Medical Journal 2000;
320: 1240-1243

Relational attributes

Related metadata references:

Supersedes Body mass index - classification, version 2, Derived
DE, NHDD, NHIMG, Superseded 01/03/2005

Body structure

Identifying and definitional attributes

<i>Metadata item type:</i>	Data Element
<i>Technical name:</i>	Person – body structure, code (ICF 2001) AN[NNNN]
<i>Synonymous names:</i>	Body structure code
<i>METeOR identifier:</i>	320147
<i>Registration status:</i>	NHIG, Standard 29/11/2006 NCSIMG, Standard 16/10/2006
<i>Definition:</i>	An anatomical part of a person's body such as organs, limbs or their components, as represented by a code.

Data element concept attributes

<i>Data element concept:</i>	Person – body structure
<i>Definition:</i>	An anatomical part of a person's body such as organs, limbs or their components.
<i>Object class:</i>	Person
<i>Property:</i>	Body structure

Value domain attributes

Representational attributes

<i>Classification scheme:</i>	International Classification of Functioning, Disability and Health 2001
<i>Representation class:</i>	Code
<i>Data type:</i>	String
<i>Format:</i>	AN[NNNN]
<i>Maximum character length:</i>	6

Collection and usage attributes

<i>Guide for use:</i>	<p>This metadata item contributes to the definition of the concept <i>disability</i> and gives an indication of the experience of disability for a person.</p> <p>Data can be collected at the three digit level in one chapter and at the chapter level in another. However it is only possible to collect data at a single level of the hierarchy in a single chapter to maintain mutual exclusivity. For example, it is not permitted to collect both 'Skin and related structures' (chapter level) and 'Structure of nails' (3 digit level) as the former includes the latter.</p> <p>The value domain below refers to the highest hierarchical level (ICF chapter level). Data collected at this level, in association with respective qualifiers (<i>Impairment extent code N, Impairment nature code N, Impairment location code N</i>) will use the codes as indicated.</p> <p>CODE s1 Structures of the nervous system CODE s2 The eye, ear and related structures CODE s3 Structures involved in voice and speech CODE s4 Structures of the cardiovascular, immunological and respiratory systems</p>
-----------------------	---

CODE s5 Structures related to the digestive, metabolic and endocrine systems
CODE s6 Structures related to the genitourinary and reproductive systems
CODE s7 Structures related to movement
CODE s8 Skin and related structures

Data collected at this level will provide a general description of the structures and can only be compared with data collected at the same level.

Each chapter contains categories at different levels ordered from general to detailed. For more detailed information the user should follow the structure of the ICF; the codes should be drawn from the same hierarchical level within any particular chapter. The full range of permissible values together with definitions is listed in the Body Structures component of the ICF.

An example of a value domain at the 3 digit level from the Structures of the nervous system chapter may include:

CODE s110 Structure of the brain
CODE s120 Spinal cord and related structures
CODE s130 Structure of the meninges
CODE s140 Structure of sympathetic nervous system
CODE s150 Structure of parasympathetic nervous system
CODE s198 Structure of the nervous system, other specified
CODE s199 Structure of the nervous system, unspecified

An example of a value domain at the 4 digit level from the Structures related to movement chapter may include:

CODE s7300 Structure of upper arm
CODE s7301 Structure of forearm
CODE s7302 Structure of hand
CODE s7500 Structure of thigh
CODE s7501 Structure of lower leg
CODE s7502 Structure of ankle and foot
CODE s7600 Structure of vertebral column

The prefix *s* denotes the domains within the component of *Body Structures*.

Source and reference attributes

<i>Submitting organisation:</i>	Australian Institute of Health and Welfare which is the Australian Collaborating Centre for the World Health Organization Family of International Classifications.
<i>Origin:</i>	WHO 2001. ICF: International Classification of Functioning, Disability and Health. Geneva: WHO AIHW 2003. ICF Australian User Guide Version 1.0. Canberra: AIHW
<i>Reference documents:</i>	Further information on the ICF, including more detailed codes, can be found in the ICF itself and the ICF Australian User Guide (AIHW 2003), at the following websites: <ul style="list-style-type: none">• WHO ICF website http://www.who.int/classifications/icf/en/• Australian Collaborating Centre ICF website http://www.aihw.gov.au/disability/icf/index.html

Data element attributes

Collection and usage attributes

Guide for use:

This data element consists of a single, neutral list of body structures that can be used to record positive or neutral body function. In conjunction with *Impairment extent code N*, it enables the provision of information about the presence and extent of impairment for any given body structures; with *Impairment nature code N*, the provision of information about the nature of the impairment for given body functions; and *Impairment location code N*, the location of the impairment for given body functions.

Where multiple body structures or **impairments of body structures** are recorded, the following prioritising system should be useful:

- The first recorded body structure or impairment of body function is the one having the greatest impact on the individual.
- Second and subsequent body structure or impairment of body function is also of relevance to the individual.

Source and reference attributes

Submitting organisation:

Australian Institute of Health and Welfare (AIHW) which is the Australian Collaborating Centre for the World Health Organization Family of International Classifications.

Relational attributes

Implementation in Data Set Specifications:

Body structures cluster NHIG, Standard 29/11/2006
NCSIMG, Standard 16/10/2006

Building/complex sub-unit number (person)

Identifying and definitional attributes

<i>Metadata item type:</i>	Data Element
<i>Technical name:</i>	Person (address)—building/complex sub-unit identifier, [X(7)]
<i>METeOR identifier:</i>	270018
<i>Registration status:</i>	NHIG, Standard 01/03/2005 NCSIMG, Standard 30/09/2005
<i>Definition:</i>	The unique number or identifier for a building/complex, marina, etc. where a person resides.

Data element concept attributes

<i>Data element concept:</i>	Person (address)—building/complex sub-unit identifier
<i>Definition:</i>	The number or identifier of a building/complex, marina, etc. where a person resides.
<i>Object class:</i>	Person
<i>Property:</i>	Building/complex sub-unit identifier

Value domain attributes

Representational attributes

<i>Representation class:</i>	Identifier
<i>Data type:</i>	String
<i>Format:</i>	[X(7)]
<i>Maximum character length:</i>	7

Data element attributes

Collection and usage attributes

<i>Guide for use:</i>	The building/complex sub-unit number must be recorded with its corresponding building/complex unit type - abbreviation. Where applicable, the number may be followed by an alphanumeric suffix.
<i>Collection methods:</i>	To be collected in conjunction with building/complex sub-unit type - abbreviation.

Source and reference attributes

<i>Submitting organisation:</i>	Australian Institute of Health and Welfare
<i>Origin:</i>	Australia Post Address Presentation Standard

Relational attributes

<i>Related metadata references:</i>	Supersedes Building/complex sub-unit number, version 1, DE, NHDD, NHIMG, Superseded 01/03/2005 Is used in the formation of Person (address)—address line, text [X(180)] NHIG, Standard 04/05/2005, NCSIMG, Standard 30/09/2005 Is used in the formation of Person (address)—health address line, text [X(180)] NHIG, Superseded 04/05/2005
-------------------------------------	--

*Implementation in Data Set
Specifications:*

Health care client identification DSS NHIG, Standard
04/05/2005

NCSIMG, Standard 03/10/2006

Health care provider identification DSS NHIG, Standard
04/05/2005

Building/complex sub-unit number (service provider organisation)

Identifying and definitional attributes

<i>Metadata item type:</i>	Data Element
<i>Technical name:</i>	Service provider organisation (address) – building/complex sub-unit identifier, [X(7)]
<i>METeOR identifier:</i>	290291
<i>Registration status:</i>	NHIG, Standard 04/05/2005 NCSIMG, Standard 30/09/2005
<i>Definition:</i>	The unique number or identifier of a building/complex, marina, etc. where an organisation is located.

Data element concept attributes

<i>Data element concept:</i>	Service provider organisation (address) – building/complex sub-unit identifier
<i>Definition:</i>	The number or identifier of a building/complex, marina, etc. where an organisation is located.
<i>Object class:</i>	Service provider organisation
<i>Property:</i>	Building/complex sub-unit identifier

Value domain attributes

Representational attributes

<i>Representation class:</i>	Identifier
<i>Data type:</i>	String
<i>Format:</i>	[X(7)]
<i>Maximum character length:</i>	7

Data element attributes

Source and reference attributes

<i>Submitting organisation:</i>	Australian Institute of Health and Welfare
<i>Origin:</i>	Australia Post Address Presentation Standard

Relational attributes

<i>Related metadata references:</i>	Is used in the formation of Service provider organisation (address) – address line, text [X(180)] NHIG, Standard 04/05/2005, NCSIMG, Standard 30/09/2005
<i>Implementation in Data Set Specifications:</i>	Health care provider identification DSS NHIG, Standard 04/05/2005

Building/complex sub-unit type—abbreviation (person)

Identifying and definitional attributes

<i>Metadata item type:</i>	Data Element
<i>Technical name:</i>	Person (address)—building/complex sub-unit type, code A[AAA]
<i>METeOR identifier:</i>	270023
<i>Registration status:</i>	NHIG, Standard 01/03/2005 NCSIMG, Standard 30/09/2005
<i>Definition:</i>	The type of building/complex where a person can be located, as represented by a code.

Data element concept attributes

<i>Data element concept:</i>	Person (address)—building/complex sub-unit type
<i>Definition:</i>	The type of building/complex where a person can be located.
<i>Object class:</i>	Person
<i>Property:</i>	Building/complex sub-unit type

Value domain attributes

Representational attributes

<i>Representation class:</i>	Code																																								
<i>Data type:</i>	String																																								
<i>Format:</i>	A[AAA]																																								
<i>Maximum character length:</i>	4																																								
<i>Permissible values:</i>	<table><thead><tr><th>Value</th><th>Meaning</th></tr></thead><tbody><tr><td>APT</td><td>Apartment</td></tr><tr><td>CTGE</td><td>Cottage</td></tr><tr><td>DUP</td><td>Duplex</td></tr><tr><td>FY</td><td>Factory</td></tr><tr><td>F</td><td>Flat</td></tr><tr><td>HSE</td><td>House</td></tr><tr><td>KSK</td><td>Kiosk</td></tr><tr><td>MSNT</td><td>Maisonette</td></tr><tr><td>MB</td><td>Marine Berth</td></tr><tr><td>OFF</td><td>Office</td></tr><tr><td>PTHS</td><td>Penthouse</td></tr><tr><td>RM</td><td>Room</td></tr><tr><td>SHED</td><td>Shed</td></tr><tr><td>SHOP</td><td>Shop</td></tr><tr><td>SITE</td><td>Site</td></tr><tr><td>SL</td><td>Stall</td></tr><tr><td>STU</td><td>Studio</td></tr><tr><td>SE</td><td>Suite</td></tr><tr><td>TNHS</td><td>Townhouse</td></tr></tbody></table>	Value	Meaning	APT	Apartment	CTGE	Cottage	DUP	Duplex	FY	Factory	F	Flat	HSE	House	KSK	Kiosk	MSNT	Maisonette	MB	Marine Berth	OFF	Office	PTHS	Penthouse	RM	Room	SHED	Shed	SHOP	Shop	SITE	Site	SL	Stall	STU	Studio	SE	Suite	TNHS	Townhouse
Value	Meaning																																								
APT	Apartment																																								
CTGE	Cottage																																								
DUP	Duplex																																								
FY	Factory																																								
F	Flat																																								
HSE	House																																								
KSK	Kiosk																																								
MSNT	Maisonette																																								
MB	Marine Berth																																								
OFF	Office																																								
PTHS	Penthouse																																								
RM	Room																																								
SHED	Shed																																								
SHOP	Shop																																								
SITE	Site																																								
SL	Stall																																								
STU	Studio																																								
SE	Suite																																								
TNHS	Townhouse																																								

U	Unit
VLLA	Villa
WARD	Ward
WE	Warehouse

Collection and usage attributes

Guide for use:

Addresses may contain multiple instances of building/complex type. Record each instance of building/complex type with its corresponding building/complex number when appropriate.

Examples:

APT 6

SHOP 3A

U 6

PTHS

Data element attributes

Collection and usage attributes

Collection methods:

To be collected in conjunction with building/complex sub unit number.

Source and reference attributes

Submitting organisation:

Australian Institute of Health and Welfare

Origin:

Health Data Standards Committee

Relational attributes

Related metadata references:

Supersedes Building/complex sub-unit type - abbreviation, version 1, DE, NHDD, NHIMG, Superseded 01/03/2005

Is used in the formation of Person (address) – address line, text [X(180)] NHIG, Standard 04/05/2005, NCSIMG, Standard 30/09/2005

Is used in the formation of Person (address) – health address line, text [X(180)] NHIG, Superseded 04/05/2005

Implementation in Data Set Specifications:

Health care client identification DSS NHIG, Standard 04/05/2005

NCSIMG, Standard 03/10/2006

Health care provider identification DSS NHIG, Standard 04/05/2005

Building/complex sub-unit type—abbreviation (service provider organisation)

Identifying and definitional attributes

<i>Metadata item type:</i>	Data Element
<i>Technical name:</i>	Service provider organisation (address) – building/complex sub-unit type, code A[AAA]
<i>METeOR identifier:</i>	290278
<i>Registration status:</i>	NHIG, Standard 04/05/2005 NCSIMG, Standard 30/09/2005
<i>Definition:</i>	The type of building/complex where an organisation can be located, as represented by a code.

Data element concept attributes

<i>Data element concept:</i>	Service provider organisation (address) – building/complex sub-unit type
<i>Definition:</i>	The type of building/complex where an organisation can be located.
<i>Object class:</i>	Service provider organisation
<i>Property:</i>	Building/complex sub-unit type

Value domain attributes

Representational attributes

<i>Representation class:</i>	Code																																		
<i>Data type:</i>	String																																		
<i>Format:</i>	A[AAA]																																		
<i>Maximum character length:</i>	4																																		
<i>Permissible values:</i>	<table><thead><tr><th>Value</th><th>Meaning</th></tr></thead><tbody><tr><td>APT</td><td>Apartment</td></tr><tr><td>CTGE</td><td>Cottage</td></tr><tr><td>DUP</td><td>Duplex</td></tr><tr><td>FY</td><td>Factory</td></tr><tr><td>F</td><td>Flat</td></tr><tr><td>HSE</td><td>House</td></tr><tr><td>KSK</td><td>Kiosk</td></tr><tr><td>MSNT</td><td>Maisonette</td></tr><tr><td>MB</td><td>Marine Berth</td></tr><tr><td>OFF</td><td>Office</td></tr><tr><td>PTHS</td><td>Penthouse</td></tr><tr><td>RM</td><td>Room</td></tr><tr><td>SHED</td><td>Shed</td></tr><tr><td>SHOP</td><td>Shop</td></tr><tr><td>SITE</td><td>Site</td></tr><tr><td>SL</td><td>Stall</td></tr></tbody></table>	Value	Meaning	APT	Apartment	CTGE	Cottage	DUP	Duplex	FY	Factory	F	Flat	HSE	House	KSK	Kiosk	MSNT	Maisonette	MB	Marine Berth	OFF	Office	PTHS	Penthouse	RM	Room	SHED	Shed	SHOP	Shop	SITE	Site	SL	Stall
Value	Meaning																																		
APT	Apartment																																		
CTGE	Cottage																																		
DUP	Duplex																																		
FY	Factory																																		
F	Flat																																		
HSE	House																																		
KSK	Kiosk																																		
MSNT	Maisonette																																		
MB	Marine Berth																																		
OFF	Office																																		
PTHS	Penthouse																																		
RM	Room																																		
SHED	Shed																																		
SHOP	Shop																																		
SITE	Site																																		
SL	Stall																																		

STU	Studio
SE	Suite
TNHS	Townhouse
U	Unit
VLLA	Villa
WARD	Ward
WE	Warehouse

Collection and usage attributes

Guide for use:

Addresses may contain multiple instances of building/complex type. Record each instance of building/complex type with its corresponding building/complex number when appropriate.

Examples:

APT 6

SHOP 3A

U 6

PTHS

Data element attributes

Source and reference attributes

Submitting organisation:

Australian Institute of Health and Welfare

Origin:

Health Data Standards Committee

Relational attributes

Related metadata references:

Is used in the formation of Service provider organisation (address) – address line, text [X(180)] NHIG, Standard 04/05/2005, NCSIMG, Standard 30/09/2005

Implementation in Data Set Specifications:

Health care provider identification DSS NHIG, Standard 04/05/2005

Building/property name (person)

Identifying and definitional attributes

<i>Metadata item type:</i>	Data Element
<i>Technical name:</i>	Person (address)—building/property name, text [X(30)]
<i>METeOR identifier:</i>	270028
<i>Registration status:</i>	NHIG, Standard 01/03/2005 NCSIMG, Standard 30/09/2005
<i>Definition:</i>	The name of a building or property where a person resides, as represented by text.

Data element concept attributes

<i>Data element concept:</i>	Person (address)—building/property name
<i>Definition:</i>	The name of a building or property where a person resides.
<i>Object class:</i>	Person
<i>Property:</i>	Building/property name

Value domain attributes

Representational attributes

<i>Representation class:</i>	Text
<i>Data type:</i>	String
<i>Format:</i>	[X(30)]
<i>Maximum character length:</i>	30

Data element attributes

Collection and usage attributes

<i>Guide for use:</i>	Usually this information is not abbreviated. Should include any reference to a wing or other components of a building complex, if applicable. A comma is to be used to separate the wing reference from the rest of the building name. Record each Building/property name relevant to the address: <ul style="list-style-type: none">• Building/property name 1 (30 alphanumeric characters)• Building/property name 2 (30 alphanumeric characters) For example: Building - TREASURY BUILDING Property - BRINDABELLA STATION
-----------------------	--

Source and reference attributes

<i>Origin:</i>	Health Data Standards Committee Australia Post Address Presentation Standard
----------------	---

Relational attributes

<i>Related metadata references:</i>	Supersedes Building/property name, version 1, DE, NHDD, NHIMG, Superseded 01/03/2005 Is used in the formation of Person (address)—address line, text
-------------------------------------	---

Implementation in Data Set Specifications:

[X(180)] NHIG, Standard 04/05/2005, NCSIMG, Standard 30/09/2005

Is used in the formation of Person (address) – health address line, text [X(180)] NHIG, Superseded 04/05/2005

Health care client identification DSS NHIG, Standard 04/05/2005

NCSIMG, Standard 03/10/2006

Health care provider identification DSS NHIG, Standard 04/05/2005

Building/property name (service provider organisation)

Identifying and definitional attributes

<i>Metadata item type:</i>	Data Element
<i>Technical name:</i>	Service provider organisation (address) – building/property name, text [X(30)]
<i>METeOR identifier:</i>	290295
<i>Registration status:</i>	NHIG, Standard 04/05/2005 NCSIMG, Standard 30/09/2005
<i>Definition:</i>	The name of a building or property where an organisation is located, as represented by text.

Data element concept attributes

<i>Data element concept:</i>	Service provider organisation (address) – building/property name
<i>Definition:</i>	The name of a building or property where an organisation is located.
<i>Object class:</i>	Service provider organisation
<i>Property:</i>	Building/property name

Value domain attributes

Representational attributes

<i>Representation class:</i>	Text
<i>Data type:</i>	String
<i>Format:</i>	[X(30)]
<i>Maximum character length:</i>	30

Data element attributes

Collection and usage attributes

<i>Guide for use:</i>	Usually this information is not abbreviated. Should include any reference to a wing or other components of a building complex, if applicable. A comma is to be used to separate the wing reference from the rest of the building name. Record each Building/property name relevant to the address: <ul style="list-style-type: none">• Building/property name 1 (30 alphanumeric characters)• Building/property name 2 (30 alphanumeric characters) For example: Building - TREASURY BUILDING Property - BRINDABELLA STATION
-----------------------	--

Source and reference attributes

<i>Origin:</i>	Health Data Standards Committee Australia Post Address Presentation Standard
----------------	---

Relational attributes

Related metadata references: Is used in the formation of Service provider organisation (address) – address line, text [X(180)] NHIG, Standard 04/05/2005, NCSIMG, Standard 30/09/2005

Implementation in Data Set Specifications: Health care provider identification DSS NHIG, Standard 04/05/2005

CVD drug therapy—condition

Identifying and definitional attributes

<i>Metadata item type:</i>	Data Element
<i>Technical name:</i>	Person – cardiovascular disease condition targeted by drug therapy, code NN
<i>METeOR identifier:</i>	270193
<i>Registration status:</i>	NHIG, Standard 01/03/2005
<i>Definition:</i>	The condition(s) for which drug therapy is being used for the prevention or long-term treatment of cardiovascular disease, as represented by a code.

Data element concept attributes

<i>Data element concept:</i>	Person – cardiovascular disease condition targeted by drug therapy
<i>Definition:</i>	Describes the condition(s) for which drug therapy is being used for the prevention or long-term treatment of cardiovascular disease.
<i>Context:</i>	Public health, health care and clinical settings: Its main use is to enable categorisation of drug management regimens used in the community for the long-term care of patients with or at increased risk of vascular disease.
<i>Object class:</i>	Person
<i>Property:</i>	Cardiovascular disease condition targeted by drug therapy

Value domain attributes

Representational attributes

<i>Representation class:</i>	Code																												
<i>Data type:</i>	String																												
<i>Format:</i>	NN																												
<i>Maximum character length:</i>	2																												
<i>Permissible values:</i>	<table><thead><tr><th>Value</th><th>Meaning</th></tr></thead><tbody><tr><td>01</td><td>Heart failure</td></tr><tr><td>02</td><td>Ischaemic heart disease</td></tr><tr><td>03</td><td>Hypertension</td></tr><tr><td>04</td><td>Atrial fibrillation (AF)</td></tr><tr><td>05</td><td>Other dysrhythmia or conductive disorder</td></tr><tr><td>06</td><td>Dyslipidaemia</td></tr><tr><td>07</td><td>Peripheral vascular disease (PVD)</td></tr><tr><td>08</td><td>Renal vascular disease</td></tr><tr><td>09</td><td>Stroke</td></tr><tr><td>10</td><td>Transient ischaemic attack (TIA)</td></tr><tr><td>97</td><td>Other</td></tr><tr><td>98</td><td>No CVD drugs prescribed</td></tr><tr><td>99</td><td>Not recorded</td></tr></tbody></table>	Value	Meaning	01	Heart failure	02	Ischaemic heart disease	03	Hypertension	04	Atrial fibrillation (AF)	05	Other dysrhythmia or conductive disorder	06	Dyslipidaemia	07	Peripheral vascular disease (PVD)	08	Renal vascular disease	09	Stroke	10	Transient ischaemic attack (TIA)	97	Other	98	No CVD drugs prescribed	99	Not recorded
Value	Meaning																												
01	Heart failure																												
02	Ischaemic heart disease																												
03	Hypertension																												
04	Atrial fibrillation (AF)																												
05	Other dysrhythmia or conductive disorder																												
06	Dyslipidaemia																												
07	Peripheral vascular disease (PVD)																												
08	Renal vascular disease																												
09	Stroke																												
10	Transient ischaemic attack (TIA)																												
97	Other																												
98	No CVD drugs prescribed																												
99	Not recorded																												
<i>Supplementary values:</i>																													

Collection and usage attributes

Guide for use: The categorisations may be made using the most recent version of the Australian Modification of the appropriate International Classification of Diseases codes.

Data element attributes

Collection and usage attributes

Guide for use: More than one code can be recorded.

Comments: References such as the Australian Medicines Handbook can be used to identify specific drugs that are appropriate for use in the management of the conditions identified in the value domain.

Source and reference attributes

Submitting organisation: Cardiovascular Data Working Group

Relational attributes

Related metadata references: Supersedes CVD drug therapy - condition, version 1, DE, NHDD, NHIMG, Superseded 01/03/2005

Implementation in Data Set Specifications: Cardiovascular disease (clinical) DSS NHIG, Superseded 15/02/2006

Cardiovascular disease (clinical) DSS NHIG, Standard 15/02/2006

Caesarean section indicator, last previous birth

Identifying and definitional attributes

<i>Metadata item type:</i>	Data Element
<i>Technical name:</i>	Female – caesarean section indicator (last previous birth) code N
<i>METeOR identifier:</i>	301993
<i>Registration status:</i>	NHIG, Standard 29/11/2006
<i>Definition:</i>	Whether a caesarean section was performed for the woman's last previous birth, as represented by a code.

Data element concept attributes

<i>Data element concept:</i>	Female – caesarean section indicator
<i>Definition:</i>	Whether a caesarean section was performed on the woman.
<i>Context:</i>	Perinatal statistics:
<i>Object class:</i>	Female
<i>Property:</i>	Caesarean section indicator

Source and reference attributes

<i>Submitting organisation:</i>	National Perinatal Data Development Committee
---------------------------------	---

Value domain attributes

Representational attributes

<i>Representation class:</i>	Code	
<i>Data type:</i>	Number	
<i>Format:</i>	N	
<i>Maximum character length:</i>	1	
<i>Permissible values:</i>	Value	Meaning
	1	Yes
	2	No

Data element attributes

Collection and usage attributes

<i>Guide for use:</i>	This item should be completed if there has been a previous birth. In the case of no previous births, the item should be left blank.
<i>Comments:</i>	Previous caesarean sections are associated with a higher risk of obstetric complications, and when used with other indicators provides important information on the quality of obstetric care. This item can be used to determine vaginal births occurring after a caesarean section delivery (VBAC).

Source and reference attributes

<i>Submitting organisation:</i>	National Perinatal Data Development Committee
---------------------------------	---

Cancer initial treatment completion date

Identifying and definitional attributes

<i>Metadata item type:</i>	Data Element
<i>Technical name:</i>	Cancer treatment— non-surgical cancer treatment completion date, DDMMYYYY
<i>METeOR identifier:</i>	288136
<i>Registration status:</i>	NHIG, Standard 04/06/2004
<i>Definition:</i>	The date on which the initial non-surgical treatment for cancer was completed.

Data element concept attributes

<i>Data element concept:</i>	Cancer treatment— non-surgical cancer treatment completion date
<i>Definition:</i>	The date on which the initial non-surgical treatment for cancer was completed.
<i>Object class:</i>	Cancer treatment
<i>Property:</i>	Non-surgical cancer treatment completion date

Value domain attributes

Representational attributes

<i>Representation class:</i>	Date
<i>Data type:</i>	Date/Time
<i>Format:</i>	DDMMYYYY
<i>Maximum character length:</i>	8

Data element attributes

Collection and usage attributes

<i>Guide for use:</i>	Collected for radiation therapy and systemic therapy.
-----------------------	---

Source and reference attributes

<i>Submitting organisation:</i>	National Cancer Control Initiative
<i>Origin:</i>	Commission on Cancer, American College of Surgeons
<i>Reference documents:</i>	Commission on Cancer, Standards of the Commission on Cancer Registry Operations and Data Standards (ROADS) Volume II (1998)

Relational attributes

<i>Related metadata references:</i>	Supersedes Cancer initial treatment - completion date, version 1, DE, NHDD, NHIMG, Superseded 01/03/2005
<i>Implementation in Data Set Specifications:</i>	Cancer (clinical) DSS NHIG, Standard 07/12/2005 <i>Information specific to this data set:</i>

This field must:

- be greater than or equal to the date of initial cancer

diagnosis, and

- be greater than or equal to the date of the initial course of treatment for cancer.

This item is collected for the analysis of outcome by treatment type.

Collecting dates for radiotherapy treatment and systemic therapy agent treatment will allow evaluation of treatments delivered and of time intervals from diagnosis to treatment, from treatment to recurrence and from treatment to death.

Cancer (clinical) DSS NHIG, Superseded 07/12/2005

Information specific to this data set:

This field must:

- be greater than or equal to the date of initial cancer diagnosis, and
- be greater than or equal to the date of the initial course of treatment for cancer.

This item is collected for the analysis of outcome by treatment type.

Collecting dates for radiotherapy treatment and systemic therapy agent treatment will allow evaluation of treatments delivered and of time intervals from diagnosis to treatment, from treatment to recurrence and from treatment to death.

Cancer initial treatment starting date

Identifying and definitional attributes

<i>Metadata item type:</i>	Data Element
<i>Technical name:</i>	Cancer treatment— non-surgical cancer treatment start date, DDMMYYYY
<i>METeOR identifier:</i>	288103
<i>Registration status:</i>	NHIG, Standard 04/06/2004
<i>Definition:</i>	The start date of the initial course of non-surgical treatment for cancer.

Data element concept attributes

<i>Data element concept:</i>	Cancer treatment— non-surgical cancer treatment start date
<i>Definition:</i>	The start date of the initial course of non-surgical treatment for cancer.
<i>Object class:</i>	Cancer treatment
<i>Property:</i>	Non-surgical cancer treatment start date

Value domain attributes

Representational attributes

<i>Representation class:</i>	Date
<i>Data type:</i>	Date/Time
<i>Format:</i>	DDMMYYYY
<i>Maximum character length:</i>	8

Data element attributes

Collection and usage attributes

<i>Guide for use:</i>	The start date of the treatment is recorded regardless of whether treatment is completed as intended or not. Treatment subsequent to a recurrence will not be recorded. Collected for radiation therapy and systemic therapy. Date of surgical treatment is collected as a separate item.
-----------------------	---

Source and reference attributes

<i>Submitting organisation:</i>	National Cancer Control Institute
<i>Origin:</i>	Commission on Cancer, Standards of the Commission on Cancer Registry Operations and Data Standards (ROADS) Volume II (1998).

Relational attributes

<i>Related metadata references:</i>	Supersedes Cancer initial treatment - starting date, version 1, DE, NHDD, NHIMG, Superseded 01/03/2005
<i>Implementation in Data Set Specifications:</i>	Cancer (clinical) DSS NHIG, Standard 07/12/2005 <i>Information specific to this data set:</i>

This field must:

- be greater than or equal to the date of initial cancer diagnosis, and
- be less than or equal to the date on which initial treatment for cancer was completed.

This metadata item is collected for the analysis of outcome by treatment type.

Collecting dates for radiotherapy treatment and systemic therapy agent treatment will allow evaluation of treatments delivered and of time intervals from diagnosis to treatment, from treatment to recurrence and from treatment to death.

Cancer (clinical) DSS NHIG, Superseded 07/12/2005

Information specific to this data set:

This field must:

- be greater than or equal to the date of initial cancer diagnosis, and
- be less than or equal to the date on which initial treatment for cancer was completed.

This metadata item is collected for the analysis of outcome by treatment type.

Collecting dates for radiotherapy treatment and systemic therapy agent treatment will allow evaluation of treatments delivered and of time intervals from diagnosis to treatment, from treatment to recurrence and from treatment to death.

Cancer staging—M stage code

Identifying and definitional attributes

<i>Metadata item type:</i>	Data Element
<i>Technical name:</i>	Person with cancer – distant metastasis status, M stage (UICC TNM Classification of Malignant Tumours 5th ed) code XX
<i>METeOR identifier:</i>	293231
<i>Registration status:</i>	NHIG, Standard 13/06/2004
<i>Definition:</i>	Absence or presence of distant metastasis at the time of diagnosis of the primary cancer, as represented by a code.

Data element concept attributes

<i>Data element concept:</i>	Person with cancer – distant metastasis status
<i>Definition:</i>	M stage is the coding system used to record the absence or presence of distant metastases at the time of diagnosis of the primary cancer. It is part of the TNM cancer staging system.
<i>Object class:</i>	Person with cancer
<i>Property:</i>	Distant metastasis status

Value domain attributes

Representational attributes

<i>Classification scheme:</i>	International Union against Cancer TNM Classification of Malignant Tumours 5th edition	
<i>Representation class:</i>	Code	
<i>Data type:</i>	String	
<i>Format:</i>	XX	
<i>Maximum character length:</i>	2	
<i>Supplementary values:</i>	Value	Meaning
	88	Not applicable

Collection and usage attributes

<i>Guide for use:</i>	Valid M codes from the current edition of the UICC TNM Classification of Malignant Tumours. Refer to the UICC reference manual, TNM Classification of Malignant Tumours for coding rules.
-----------------------	--

Data element attributes

Collection and usage attributes

<i>Guide for use:</i>	Choose the lower (less advanced) M category when there is any uncertainty.
<i>Collection methods:</i>	From information provided by the treating doctor and recorded on the patient's medical record.
<i>Comments:</i>	Cancer prognosis and survival can be related to the extent of the disease at diagnosis. Survival rates are generally higher if the disease is localised to the organ of origin compared with cases in which the tumour has spread beyond the primary site.

Staging systems seek to classify patients having a similar prognosis into groups or stages. TNM staging is an internationally agreed staging classification system based on the anatomical site of the primary tumour and its extent of spread. The T component refers to the size of the tumour and whether or not it has spread to surrounding tissues. The N component describes the presence or absence of tumour in regional lymph nodes. The M component refers to the presence or absence of tumour at sites distant from the primary site. TNM staging applies to solid tumours excluding brain tumours.

Source and reference attributes

Origin: International Union Against Cancer (UICC)
Commission on Cancer, American College of Surgeons

Reference documents: UICC TNM Classification of Malignant Tumours (5th Edition) (1997)
Commission on Cancer, Standards of the Commission on Cancer Registry Operations and Data Standards (ROADS) Volume II (1998).

Relational attributes

Related metadata references: Supersedes Cancer staging - M stage code, version 1, DE, NHDD, NHIMG, Superseded 01/03/2005
Is used in the formation of Person with cancer – extent of primary cancer, TNM stage (UICC TNM Classification of Malignant Tumours 5th ed) code XXXX{[X]XX} NHIG, Standard 04/06/2004

Implementation in Data Set Specifications: Cancer (clinical) DSS NHIG, Standard 07/12/2005
Information specific to this data set:

For survival analysis adjusted by stage at diagnosis and distribution of cancer cases by type and stage.

Cancer (clinical) DSS NHIG, Superseded 07/12/2005
Information specific to this data set:

For survival analysis adjusted by stage at diagnosis and distribution of cancer cases by type and stage.

Cancer staging—N stage code

Identifying and definitional attributes

<i>Metadata item type:</i>	Data Element
<i>Technical name:</i>	Person with cancer – regional lymph node metastasis status, N stage (UICC TNM Classification of Malignant Tumours 5th ed) code XX
<i>METeOR identifier:</i>	293254
<i>Registration status:</i>	NHIG, Standard 13/06/2004
<i>Definition:</i>	Extent of regional lymph node metastasis at the time of diagnosis of the primary cancer, as represented by a code.

Data element concept attributes

<i>Data element concept:</i>	Person with cancer – regional lymph node metastasis status
<i>Definition:</i>	N stage is the coding system used to denote the absence or presence of regional lymph node metastases. It classifies the extent of regional lymph node metastases at the time of diagnosis of the primary cancer. It is a part of the TNM cancer staging system.
<i>Object class:</i>	Person with cancer
<i>Property:</i>	Regional lymph node metastasis status

Value domain attributes

Representational attributes

<i>Classification scheme:</i>	International Union against Cancer TNM Classification of Malignant Tumours 5th edition	
<i>Representation class:</i>	Code	
<i>Data type:</i>	String	
<i>Format:</i>	XX	
<i>Maximum character length:</i>	2	
<i>Supplementary values:</i>	Value	Meaning
	88	Not applicable

Collection and usage attributes

<i>Guide for use:</i>	Valid N codes from the current edition of the UICC TNM Classification of Malignant Tumours. Refer to the UICC reference manual, TNM Classification of Malignant Tumours for coding rules.
-----------------------	--

Data element attributes

Collection and usage attributes

<i>Guide for use:</i>	Choose the lower (less advanced) N category when there is any uncertainty.
<i>Collection methods:</i>	From information provided by the treating doctor and recorded on the patient's medical record.
<i>Comments:</i>	Cancer prognosis and survival can be related to the extent of

the disease at diagnosis. Survival rates are generally higher if the disease is localised to the organ of origin compared with cases in which the tumour has spread beyond the primary site. Staging systems seek to classify patients having a similar prognosis into groups or stages. TNM staging is an internationally agreed staging classification system based on the anatomical site of the primary tumour and its extent of spread. The T component refers to the size of the tumour and whether or not it has spread to surrounding tissues. The N component describes the presence or absence of tumour in regional lymph nodes. The M component refers to the presence or absence of tumour at sites distant from the primary site. TNM staging applies to solid tumours excluding brain tumours.

Source and reference attributes

Reference documents:

Commission on Cancer, Standards of the Commission on Cancer Registry Operations and Data Standards (ROADS) Volume II (1998).

Relational attributes

Related metadata references:

Supersedes Cancer staging - N stage code, version 1, DE, NHDD, NHIMG, Superseded 01/03/2005

Is used in the formation of Person with cancer – extent of primary cancer, TNM stage (UICC TNM Classification of Malignant Tumours 5th ed) code XXXX{[X]XX} NHIG, Standard 04/06/2004

Implementation in Data Set Specifications:

Cancer (clinical) DSS NHIG, Standard 07/12/2005

Information specific to this data set:

For survival analysis adjusted by stage at diagnosis and distribution of cancer cases by type and stage.

Cancer (clinical) DSS NHIG, Superseded 07/12/2005

Information specific to this data set:

For survival analysis adjusted by stage at diagnosis and distribution of cancer cases by type and stage.

Cancer staging—T stage code

Identifying and definitional attributes

<i>Metadata item type:</i>	Data Element
<i>Technical name:</i>	Person with cancer – primary tumour status, T stage (UICC TNM Classification of Malignant Tumours 5th ed) code XX[X]
<i>METeOR identifier:</i>	293270
<i>Registration status:</i>	NHIG, Standard 13/06/2004
<i>Definition:</i>	Extent of primary cancer including tumour size, at the time of diagnosis, as represented by a code.

Data element concept attributes

<i>Data element concept:</i>	Person with cancer – primary tumour status
<i>Definition:</i>	T stage is the coding system used to identify the presence of primary tumour. It reflects the tumour size and extent of the primary cancer at the time of diagnosis. It is a part of the TNM cancer staging system.
<i>Object class:</i>	Person with cancer
<i>Property:</i>	Primary tumour status

Value domain attributes

Representational attributes

<i>Classification scheme:</i>	International Union against Cancer TNM Classification of Malignant Tumours 5th edition	
<i>Representation class:</i>	Code	
<i>Data type:</i>	String	
<i>Format:</i>	XX[X]	
<i>Maximum character length:</i>	3	
<i>Supplementary values:</i>	Value	Meaning
	88	Not applicable

Collection and usage attributes

<i>Guide for use:</i>	Valid T codes from the current edition of the UICC TNM Classification of Malignant Tumours. Refer to the UICC reference manual, TNM Classification of Malignant Tumours for coding rules.
-----------------------	--

Data element attributes

Collection and usage attributes

<i>Guide for use:</i>	Choose the lower (less advanced) T category when there is any uncertainty.
<i>Collection methods:</i>	From information provided by the treating doctor and recorded on the patient's medical record.
<i>Comments:</i>	Cancer prognosis and survival can be related to the extent of the disease at diagnosis. Survival rates are generally higher if the disease is localised to the organ of origin compared with

cases in which the tumour has spread beyond the primary site. Staging systems seek to classify patients having a similar prognosis into groups or stages. TNM staging is an internationally agreed staging classification system based on the anatomical site of the primary tumour and its extent of spread. The T component refers to the size of the tumour and whether or not it has spread to surrounding tissues. The N component describes the presence or absence of tumour in regional lymph nodes. The M component refers to the presence or absence of tumour at sites distant from the primary site. TNM staging applies to solid tumours excluding brain tumours.

Source and reference attributes

Reference documents:

Standards of the Commission on Cancer Registry Operations and Data Standards (ROADS) Volume II (1998).

Relational attributes

Related metadata references:

Supersedes Cancer staging - T stage code, version 1, DE, NHDD, NHIMG, Superseded 01/03/2005

Is used in the formation of Person with cancer – extent of primary cancer, TNM stage (UICC TNM Classification of Malignant Tumours 5th ed) code XXXX{[X]XX} NHIG, Standard 04/06/2004

Implementation in Data Set Specifications:

Cancer (clinical) DSS NHIG, Standard 07/12/2005

Information specific to this data set:

For survival analysis adjusted by stage at diagnosis and distribution of cancer cases by type and stage.

Cancer (clinical) DSS NHIG, Superseded 07/12/2005

Information specific to this data set:

For survival analysis adjusted by stage at diagnosis and distribution of cancer cases by type and stage.

Cancer staging—TNM stage grouping code

Identifying and definitional attributes

<i>Metadata item type:</i>	Data Element
<i>Technical name:</i>	Person with cancer – extent of primary cancer, TNM stage (UICC TNM Classification of Malignant Tumours 5th ed) code XXXX{[X]XX}
<i>METeOR identifier:</i>	296925
<i>Registration status:</i>	NHIG, Standard 04/06/2004
<i>Definition:</i>	The anatomical extent of disease at diagnosis based on the previously coded T,N and M stage categories, as represented by a code.

Data element concept attributes

<i>Data element concept:</i>	Person with cancer – extent of primary cancer
<i>Definition:</i>	The stage grouping defines the anatomical extent of disease at diagnosis based on the previously coded T, N and M stage categories.
<i>Object class:</i>	Person with cancer
<i>Property:</i>	Extent of primary cancer

Value domain attributes

Representational attributes

<i>Classification scheme:</i>	International Union against Cancer TNM Classification of Malignant Tumours 5th edition	
<i>Representation class:</i>	Code	
<i>Data type:</i>	String	
<i>Format:</i>	XXXX{[X]XX}	
<i>Maximum character length:</i>	6	
<i>Supplementary values:</i>	Value	Meaning
	8888	Not applicable
	9999	Unknown, Stage X

Collection and usage attributes

<i>Guide for use:</i>	Valid stage grouping codes from the current edition of the UICC TNM Classification of Malignant Tumours.
-----------------------	--

Data element attributes

Collection and usage attributes

<i>Guide for use:</i>	Refer to the UICC reference manual, TNM Classification of Malignant Tumours for coding rules. Choose the lower (less advanced) T category when there is any uncertainty.
<i>Collection methods:</i>	From information provided by the treating doctor and recorded on the patient's medical record.

Relational attributes

Related metadata references:

Supersedes Cancer staging - TNM stage grouping code, version 1, DE, NHDD, NHIMG, Superseded 01/03/2005

Is formed using Person with cancer – distant metastasis status, M stage (UICC TNM Classification of Malignant Tumours 5th ed) code XX NHIG, Standard 13/06/2004

Is formed using Person with cancer – regional lymph node metastasis status, N stage (UICC TNM Classification of Malignant Tumours 5th ed) code XX NHIG, Standard 13/06/2004

Is formed using Person with cancer – primary tumour status, T stage (UICC TNM Classification of Malignant Tumours 5th ed) code XX[X] NHIG, Standard 13/06/2004

Implementation in Data Set Specifications:

Cancer (clinical) DSS NHIG, Standard 07/12/2005

Information specific to this data set:

For survival analysis adjusted by stage at diagnosis and distribution of cancer cases by type and stage.

Cancer (clinical) DSS NHIG, Superseded 07/12/2005

Information specific to this data set:

For survival analysis adjusted by stage at diagnosis and distribution of cancer cases by type and stage.

Cancer treatment type

Identifying and definitional attributes

<i>Metadata item type:</i>	Data Element
<i>Technical name:</i>	Cancer treatment—cancer treatment type, code N
<i>METeOR identifier:</i>	288185
<i>Registration status:</i>	NHIG, Standard 04/06/2004
<i>Definition:</i>	The type of treatment for cancer given as initial treatment for the particular patient, as represented by a code.

Data element concept attributes

<i>Data element concept:</i>	Cancer treatment—cancer treatment type
<i>Definition:</i>	The type of treatment for cancer given as initial treatment for the particular patient.
<i>Context:</i>	This item is collected for surgical treatment, radiation therapy and systemic therapy. It is used for correlating outcome with original intent of the treatment.
<i>Object class:</i>	Cancer treatment
<i>Property:</i>	Cancer treatment type

Value domain attributes

Representational attributes

<i>Representation class:</i>	Code																		
<i>Data type:</i>	Number																		
<i>Format:</i>	N																		
<i>Maximum character length:</i>	1																		
<i>Permissible values:</i>	<table><thead><tr><th>Value</th><th>Meaning</th></tr></thead><tbody><tr><td>0</td><td>No treatment</td></tr><tr><td>1</td><td>Surgical treatment</td></tr><tr><td>2</td><td>Radiation therapy</td></tr><tr><td>3</td><td>Systemic agent therapy</td></tr><tr><td>4</td><td>Surgical and radiation treatment</td></tr><tr><td>5</td><td>Surgical treatment and systemic agent treatment</td></tr><tr><td>6</td><td>Radiation and systemic agent treatment</td></tr><tr><td>7</td><td>All three treatment types</td></tr></tbody></table>	Value	Meaning	0	No treatment	1	Surgical treatment	2	Radiation therapy	3	Systemic agent therapy	4	Surgical and radiation treatment	5	Surgical treatment and systemic agent treatment	6	Radiation and systemic agent treatment	7	All three treatment types
Value	Meaning																		
0	No treatment																		
1	Surgical treatment																		
2	Radiation therapy																		
3	Systemic agent therapy																		
4	Surgical and radiation treatment																		
5	Surgical treatment and systemic agent treatment																		
6	Radiation and systemic agent treatment																		
7	All three treatment types																		

Source and reference attributes

<i>Submitting organisation:</i>	Australian Institute of Health and Welfare
---------------------------------	--

Data element attributes

Source and reference attributes

<i>Origin:</i>	Commission on Cancer, American College of Surgeons. New South Wales Health Department.
----------------	---

Reference documents:

Commission on Cancer, Standards of the Commission on
Cancer Registry Operations and Data Standards (ROADS)
Volume II (1998)

Public Health Division NSW Clinical Cancer Data Collection for
Outcomes and Quality. Data Dictionary Version 1 Sydney NSW
Health Dept (2001)

Relational attributes

Related metadata references:

Supersedes Cancer treatment type, version 1, DE, NHDD,
NHIMG, Superseded 01/03/2005

*Implementation in Data Set
Specifications:*

Cancer (clinical) DSS NHIG, Standard 07/12/2005

Cancer (clinical) DSS NHIG, Superseded 07/12/2005

Cancer treatment—target site (ICD-10-AM)

Identifying and definitional attributes

<i>Metadata item type:</i>	Data Element
<i>Technical name:</i>	Cancer treatment—target site for cancer treatment, code (ICD-10-AM 5th edn) ANN{.N[N]}
<i>METeOR identifier:</i>	333822
<i>Registration status:</i>	NHIG, Standard 07/12/2005
<i>Definition:</i>	The site or region which is the target of particular surgical or radiotherapy treatment, as represented by an ICD-10-AM code.

Data element concept attributes

<i>Data element concept:</i>	Cancer treatment—target site for cancer treatment
<i>Definition:</i>	The site or region of cancer which is the target of a particular surgical or radiotherapy treatment.
<i>Object class:</i>	Cancer treatment
<i>Property:</i>	Target site for cancer treatment

Value domain attributes

Representational attributes

<i>Classification scheme:</i>	International Statistical Classification of Diseases and Related Health Problems, Tenth Revision, Australian Modification 5th edition
<i>Representation class:</i>	Code
<i>Data type:</i>	String
<i>Format:</i>	ANN{.N[N]}
<i>Maximum character length:</i>	6

Data element attributes

Collection and usage attributes

<i>Guide for use:</i>	This information is collected for surgical and radiotherapy treatments. Current edition of International Classification of Diseases (ICD-10-AM), Australian Modification, National Centre for Classification in Health, Sydney is used.
-----------------------	--

Relational attributes

<i>Related metadata references:</i>	Supersedes Cancer treatment—target site for cancer treatment, code (ICD-10-AM 4th edn) ANN{.N[N]} NHIG, Superseded 07/12/2005
<i>Implementation in Data Set Specifications:</i>	Cancer (clinical) DSS NHIG, Standard 07/12/2005

Cancer treatment—target site (ICDO-3)

Identifying and definitional attributes

<i>Metadata item type:</i>	Data Element
<i>Technical name:</i>	Cancer treatment—target site for cancer treatment, code (ICDO-3) ANN
<i>METeOR identifier:</i>	293161
<i>Registration status:</i>	NHIG, Standard 13/06/2004
<i>Definition:</i>	The site or region of cancer which is the target of a particular surgical or radiotherapy treatment, as represented by an ICDO-3 code.

Data element concept attributes

<i>Data element concept:</i>	Cancer treatment—target site for cancer treatment
<i>Definition:</i>	The site or region of cancer which is the target of a particular surgical or radiotherapy treatment.
<i>Object class:</i>	Cancer treatment
<i>Property:</i>	Target site for cancer treatment

Value domain attributes

Representational attributes

<i>Classification scheme:</i>	International Classification of Diseases for Oncology 3rd edition
<i>Representation class:</i>	Code
<i>Data type:</i>	String
<i>Format:</i>	ANN
<i>Maximum character length:</i>	3

Data element attributes

Collection and usage attributes

<i>Guide for use:</i>	This information is collected for surgical and radiotherapy treatments. Current edition of International Classification of Diseases for Oncology (ICD-O), World Health Organisation is used. Major organ only - first 3 characters.
-----------------------	---

Relational attributes

<i>Related metadata references:</i>	Supersedes Cancer treatment - target site, version 1, DE, NHDD, NHIMG, Superseded 01/03/2005
<i>Implementation in Data Set Specifications:</i>	Cancer (clinical) DSS NHIG, Standard 07/12/2005 Cancer (clinical) DSS NHIG, Superseded 07/12/2005

Cardiovascular medication (current)

Identifying and definitional attributes

<i>Metadata item type:</i>	Data Element
<i>Technical name:</i>	Person – cardiovascular medication taken (current), code N
<i>METeOR identifier:</i>	270237
<i>Registration status:</i>	NHIG, Standard 01/03/2005
<i>Definition:</i>	Whether the individual is currently taking cardiovascular medication, as represented by a code.

Data element concept attributes

<i>Data element concept:</i>	Person – cardiovascular medication taken
<i>Definition:</i>	Whether the individual is taking some of the following cardiovascular medications: <ul style="list-style-type: none">• Angiotensin converting enzyme (ACE) inhibitors• Angiotensin II [A2] antagonists• Beta blockers• Calcium antagonists
<i>Context:</i>	Public health, health care and clinical settings.
<i>Object class:</i>	Person
<i>Property:</i>	Cardiovascular medication taken

Value domain attributes

Representational attributes

<i>Representation class:</i>	Code														
<i>Data type:</i>	Number														
<i>Format:</i>	N														
<i>Maximum character length:</i>	1														
<i>Permissible values:</i>	<table><thead><tr><th>Value</th><th>Meaning</th></tr></thead><tbody><tr><td>1</td><td>Angiotensin converting enzyme (ACE) inhibitors</td></tr><tr><td>2</td><td>Angiotensin II (A2) receptor blockers</td></tr><tr><td>3</td><td>Beta blockers</td></tr><tr><td>4</td><td>Calcium antagonists</td></tr><tr><td>8</td><td>None of the above</td></tr><tr><td>9</td><td>Not stated/inadequately described</td></tr></tbody></table>	Value	Meaning	1	Angiotensin converting enzyme (ACE) inhibitors	2	Angiotensin II (A2) receptor blockers	3	Beta blockers	4	Calcium antagonists	8	None of the above	9	Not stated/inadequately described
Value	Meaning														
1	Angiotensin converting enzyme (ACE) inhibitors														
2	Angiotensin II (A2) receptor blockers														
3	Beta blockers														
4	Calcium antagonists														
8	None of the above														
9	Not stated/inadequately described														
<i>Supplementary values:</i>															

Collection and usage attributes

<i>Guide for use:</i>	CODE 1 Angiotensin converting enzyme (ACE) inhibitors Use this code for ACE inhibitors (captopril, enalapril, fosinopril, lisinopril, perindopril, quinapril, ramipril and trandolapril). CODE 2 Angiotensin II (A2) receptor blockers Use this code for Angiotensin II receptor blockers (candesartan, eprosartan, irbesartan and telmisartan). CODE 3 Beta blockers Use this code for Beta blockers (atenolol, carvedilol, labetalol,
-----------------------	--

metoprolol, oxprenolol, pindolol, propranolol and sotalol).

CODE 4 Calcium antagonists

Use this code for Calcium antagonists (amlodipine, diltiazem, felodipine, lercanidipine, nifedipine and verapamil).

CODE 8 None of the above

This code is used when none of the listed medications is being taken by the person.

CODE 9 Not stated/inadequately described

This code should only be used in situations where it is not practicable to ask the questions.

Collection methods:

The person should be asked a series of questions about any current medication for a cardiovascular condition as follows:

Are you currently taking any medication for a cardiovascular condition?

___Yes ___No

If the person answers 'NO', then code 8 should be applied.

If the person answers 'YES', then ask which one(s) (from the list of drugs in the Guide for use).

Ace Inhibitors ___Yes ___No

Angiotensin II receptor blockers ___Yes ___No

Beta blockers ___Yes ___No

Calcium antagonists ___Yes ___No

The appropriate code should be recorded for each type of medication currently in use.

Data element attributes

Collection and usage attributes

Collection methods:

A person may be taking one or more of the following medications for a cardiovascular condition. Therefore more than one code may be reported.

Source and reference attributes

Origin:

National Diabetes Outcomes Quality Review Initiative (NDOQRIN) data dictionary. Australian Medicines Handbook: last modified by February 2001 Contents of Cardiovascular, Version 3, 1999 Therapeutic Guidelines Limited (05.04.2002)].

Relational attributes

Related metadata references:

Supersedes Cardiovascular medication - Superseded 01/03/2005, version 1, DE, NHDD, NHIMG, Superseded 01/03/2005

Implementation in Data Set Specifications:

Diabetes (clinical) DSS NHIG, Superseded 21/09/2005

Information specific to this data set:

A person may be taking one or more of the following medications for a cardiovascular condition. Therefore more than one code may be reported.

Example 1:

If a person takes one of the ACE inhibitors and a Beta blocker, the code recorded would be 13.

Example 2:

If a person takes one of the ACE inhibitors, an Angiotensin II receptor blocker and a Beta blocker, the code recorded would be 123.

Diabetes (clinical) DSS NHIG, Standard 21/09/2005

Information specific to this data set:

A person may be taking one or more of the following medications for a cardiovascular condition. Therefore more than one code may be reported.

Example 1:

If a person takes one of the ACE inhibitors and a Beta blocker, the code recorded would be 13.

Example 2:

If a person takes one of the ACE inhibitors, an Angiotensin II receptor blocker and a Beta blocker, the code recorded would be 123.

Care type

Identifying and definitional attributes

<i>Metadata item type:</i>	Data Element
<i>Technical name:</i>	Hospital service – care type, code N[N].N
<i>METeOR identifier:</i>	270174
<i>Registration status:</i>	NHIG, Standard 01/03/2005
<i>Definition:</i>	The overall nature of a clinical service provided to an admitted patient during an episode of care (admitted care), or the type of service provided by the hospital for boarders or posthumous organ procurement (other care), as represented by a code.

Data element concept attributes

<i>Data element concept:</i>	Hospital service – care type
<i>Definition:</i>	The overall nature of a clinical service provided to an admitted patient during an episode of care (admitted care), or the type of service provided by the hospital for boarders or posthumous organ procurement (other care).
<i>Context:</i>	Admitted patient care and hospital activity: For admitted patients, the type of care received will determine the appropriate casemix classification employed to classify the episode of care.
<i>Object class:</i>	Hospital service
<i>Property:</i>	Care type

Value domain attributes

Representational attributes

<i>Representation class:</i>	Code																				
<i>Data type:</i>	Number																				
<i>Format:</i>	N[N].N																				
<i>Maximum character length:</i>	3																				
<i>Permissible values:</i>	<table><thead><tr><th>Value</th><th>Meaning</th></tr></thead><tbody><tr><td>1.0</td><td>Acute care (Admitted care)</td></tr><tr><td>2.0</td><td>Rehabilitation care (Admitted care)</td></tr><tr><td>2.1</td><td>Rehabilitation care delivered in a designated unit (optional)</td></tr><tr><td>2.2</td><td>Rehabilitation care according to a designated program (optional)</td></tr><tr><td>2.3</td><td>Rehabilitation care is the principal clinical intent (optional)</td></tr><tr><td>3.0</td><td>Palliative care</td></tr><tr><td>3.1</td><td>Palliative care delivered in a designated unit (optional)</td></tr><tr><td>3.2</td><td>Palliative care according to a designated program (optional)</td></tr><tr><td>3.3</td><td>Palliative care is the principal clinical intent</td></tr></tbody></table>	Value	Meaning	1.0	Acute care (Admitted care)	2.0	Rehabilitation care (Admitted care)	2.1	Rehabilitation care delivered in a designated unit (optional)	2.2	Rehabilitation care according to a designated program (optional)	2.3	Rehabilitation care is the principal clinical intent (optional)	3.0	Palliative care	3.1	Palliative care delivered in a designated unit (optional)	3.2	Palliative care according to a designated program (optional)	3.3	Palliative care is the principal clinical intent
Value	Meaning																				
1.0	Acute care (Admitted care)																				
2.0	Rehabilitation care (Admitted care)																				
2.1	Rehabilitation care delivered in a designated unit (optional)																				
2.2	Rehabilitation care according to a designated program (optional)																				
2.3	Rehabilitation care is the principal clinical intent (optional)																				
3.0	Palliative care																				
3.1	Palliative care delivered in a designated unit (optional)																				
3.2	Palliative care according to a designated program (optional)																				
3.3	Palliative care is the principal clinical intent																				

	(optional)
4.0	Geriatric evaluation and management
5.0	Psychogeriatric care
6.0	Maintenance care
7.0	Newborn care
8.0	Other admitted patient care
9.0	Organ procurement - posthumous (Other care)
10.0	Hospital boarder (Other care)

Collection and usage attributes

Guide for use:

Persons with mental illness may receive any one of the care types (except newborn and organ procurement). Classification depends on the principal clinical intent of the care received.

Admitted care can be one of the following:

CODE 1.0 Acute care (Admitted care)

Acute care is care in which the clinical intent or treatment goal is to:

- manage labour (obstetric)
- cure illness or provide definitive treatment of injury
- perform surgery
- relieve symptoms of illness or injury (excluding palliative care)
- reduce severity of an illness or injury
- protect against exacerbation and/or complication of an illness and/or injury which could threaten life or normal function
- perform diagnostic or therapeutic procedures.

CODE 2.0 Rehabilitation care (Admitted care)

Rehabilitation care is care in which the clinical intent or treatment goal is to improve the functional status of a patient with an impairment, disability or handicap. It is usually evidenced by a multi-disciplinary rehabilitation plan comprising negotiated goals and indicative time frames which are evaluated by a periodic assessment using a recognised functional assessment measure. It includes care provided:

- in a designated rehabilitation unit (code 2.1), or
- in a designated rehabilitation program, or in a psychiatric rehabilitation program as designated by the state health authority for public patients in a recognised hospital, for private patients in a public or private hospital as approved by a registered health benefits organisation (code 2.2), or
- under the principal clinical management of a rehabilitation physician or, in the opinion of the treating doctor, when the principal clinical intent of care is rehabilitation (code 2.3).

Optional:

CODE 2.1 Rehabilitation care delivered in a designated unit (optional)

A designated rehabilitation care unit is a dedicated ward or unit (and can be a stand-alone unit) that receives identified funding for rehabilitation care and/or primarily delivers rehabilitation care.

CODE 2.2 Rehabilitation care according to a designated

program (optional)

In a designated rehabilitation care program, care is delivered by a specialised team of staff who provide rehabilitation care to patients in beds that may or may not be dedicated to rehabilitation care. The program may, or may not be funded through identified rehabilitation care funding. Code 2.1 should be used instead of code 2.2 if care is being delivered in a designated rehabilitation care program and a designated rehabilitation care unit.

CODE 2.3 Rehabilitation care is the principal clinical intent (optional)

Rehabilitation as principal clinical intent (code 2.3) occurs when the patient is primarily managed by a medical practitioner who is a specialist in rehabilitation care or when, in the opinion of the treating medical practitioner, the care provided is rehabilitation care even if the doctor is not a rehabilitation care specialist. The exception to this is when the medical practitioner is providing care within a designated unit or a designated program, in which case code 2.1 or 2.2 should be used, respectively.

Code 3.0 Palliative care

Palliative care is care in which the clinical intent or treatment goal is primarily quality of life for a patient with an active, progressive disease with little or no prospect of cure. It is usually evidenced by an interdisciplinary assessment and/or management of the physical, psychological, emotional and spiritual needs of the patient; and a grief and bereavement support service for the patient and their carers/family. It includes care provided:

- in a palliative care unit (code 3.1); or
- in a designated palliative care program (code 3.2); or
- under the principal clinical management of a palliative care physician or, in the opinion of the treating doctor, when the principal clinical intent of care is palliation (code 3.3).

Optional:

CODE 3.1 Palliative care delivered in a designated unit (optional)

A designated palliative care unit is a dedicated ward or unit (and can be a stand-alone unit) that receives identified funding for palliative care and/or primarily delivers palliative care.

CODE 3.2 Palliative care according to a designated program (optional)

In a designated palliative care program, care is delivered by a specialised team of staff who provide palliative care to patients in beds that may or may not be dedicated to palliative care. The program may, or may not be funded through identified palliative care funding. Code 3.1 should be used instead of code 3.2 if care is being delivered in a designated palliative care program and a designated palliative care unit.

CODE 3.3 Palliative care is the principal clinical intent (optional)

Palliative care as principal clinical intent occurs when the patient is primarily managed by a medical practitioner who is a specialist in palliative care or when, in the opinion of the treating medical practitioner, the care provided is palliative care even if the doctor is not a palliative care specialist. The

exception to this is when the medical practitioner is providing care within a designated unit or a designated program, in which case code 3.1 or 3.2 should be used, respectively. For example, code 3.3 would apply to a patient dying of cancer who was being treated in a geriatric ward without specialist input by palliative care staff.

CODE 4.0 Geriatric evaluation and management

Geriatric evaluation and management is care in which the clinical intent or treatment goal is to maximise health status and/or optimise the living arrangements for a patient with multi-dimensional medical conditions associated with disabilities and psychosocial problems, who is usually (but not always) an older patient. This may also include younger adults with clinical conditions generally associated with old age. This care is usually evidenced by multi-disciplinary management and regular assessments against a management plan that is working towards negotiated goals within indicative time frames. Geriatric evaluation and management includes care provided:

- in a geriatric evaluation and management unit; or
- in a designated geriatric evaluation and management program; or
- under the principal clinical management of a geriatric evaluation and management physician or,
- in the opinion of the treating doctor, when the principal clinical intent of care is geriatric evaluation and management.

CODE 5.0 Psychogeriatric care

Psychogeriatric care is care in which the clinical intent or treatment goal is improvement in health, modification of symptoms and enhancement in function, behaviour and/or quality of life for a patient with an age-related organic brain impairment with significant behavioural or late onset psychiatric disturbance or a physical condition accompanied by severe psychiatric or behavioural disturbance. The care is usually evidenced by multi-disciplinary management and regular assessments against a management plan that is working towards negotiated goals within indicative time frames. It includes care provided:

- in a psychogeriatric care unit;
- in a designated psychogeriatric care program; or
- under the principal clinical management of a psychogeriatric physician or,
- in the opinion of the treating doctor, when the principal clinical intent of care is psychogeriatric care.

CODE 6.0 Maintenance care

Maintenance care is care in which the clinical intent or treatment goal is prevention of deterioration in the functional and current health status of a patient with a disability or severe level of functional impairment. Following assessment or treatment the patient does not require further complex assessment or stabilisation, and requires care over an indefinite period. This care includes that provided to a patient who would normally receive care in another setting eg at home, or in a residential aged care service, by a relative or carer, that is

unavailable in the short term.

CODE 7.0 Newborn care

Newborn care is initiated when the patient is born in hospital or is nine days old or less at the time of admission. Newborn care continues until the care type changes or the patient is separated:

- patients who turn 10 days of age and do not require clinical care are separated and, if they remain in the hospital, are designated as boarders
- patients who turn 10 days of age and require clinical care continue in a newborn episode of care until separated
- patients aged less than 10 days and not admitted at birth (eg transferred from another hospital) are admitted with newborn care type
- patients aged greater than 9 days not previously admitted (eg transferred from another hospital) are either boarders or admitted with an acute care type
- within a newborn episode of care, until the baby turns 10 days of age, each day is either a qualified or unqualified day
- a newborn is qualified when it meets at least one of the criteria detailed in **Newborn qualification status**.

Within a newborn episode of care, each day after the baby turns 10 days of age is counted as a qualified patient day. Newborn qualified days are equivalent to acute days and may be denoted as such.

CODE 8.0 Other admitted patient care

Other admitted patient care is care where the principal clinical intent does meet the criteria for any of the above.

Other care can be one of the following:

CODE 9.0 Organ procurement - posthumous (Other care)

Organ procurement - posthumous is the procurement of human tissue for the purpose of transplantation from a donor who has been declared brain dead.

Diagnoses and procedures undertaken during this activity, including mechanical ventilation and tissue procurement, should be recorded in accordance with the relevant ICD-10-AM Australian Coding Standards. These patients are not admitted to the hospital but are registered by the hospital.

CODE 10.0 Hospital boarder (Other care)

Hospital boarder is a person who is receiving food and/or accommodation but for whom the hospital does not accept responsibility for treatment and/or care.

Hospital boarders are not admitted to the hospital. However, a hospital may register a boarder. Babies in hospital at age 9 days of less cannot be boarders. They are admitted patients with each day of stay deemed to be either qualified or unqualified.

Comments:

Unqualified newborn days (and separations consisting entirely of unqualified newborn days are not to be counted under the Australian Health Care Agreements and they are ineligible for health insurance benefit purposes.

Data element attributes

Source and reference attributes

Origin:

National Health Data Committee

Relational attributes

Related metadata references:

Supersedes Care type, version 4, DE, NHDD, NHIMG, Superseded 01/03/2005

Is used in the formation of Episode of care – number of psychiatric care days, total N[NNNN] NHIG, Standard 01/03/2005

Implementation in Data Set Specifications:

Admitted patient care NMDS NHIG, Superseded 07/12/2005

Implementation start date: 01/07/2005

Implementation end date: 30/06/2006

Admitted patient care NMDS 2007-2008 NHIG, Standard 29/11/2006

Implementation start date: 01/07/2007

Admitted patient mental health care NMDS NHIG, Superseded 07/12/2005

Implementation start date: 01/07/2005

Implementation end date: 30/06/2006

Admitted patient mental health care NMDS NHIG, Superseded 23/10/2006

Implementation start date: 01/07/2006

Implementation end date: 30/06/2007

Admitted patient mental health care NMDS 2007-2008 NHIG, Standard 23/10/2006

Implementation start date: 01/07/2007

Admitted patient palliative care NMDS NHIG, Superseded 07/12/2005

Implementation start date: 01/07/2005

Implementation end date: 30/06/2006

Admitted patient palliative care NMDS 2006-2007 NHIG, Superseded 23/10/2006

Implementation start date: 01/07/2006

Implementation end date: 30/06/2007

Admitted patient palliative care NMDS 2007-08 NHIG, Standard 23/10/2006

Implementation start date: 01/07/2007

Carer participation arrangements—carer consultants employed

Identifying and definitional attributes

<i>Metadata item type:</i>	Data Element
<i>Technical name:</i>	Specialised mental health service organisation – carer participation arrangements status (carer consultants employed), code N
<i>METeOR identifier:</i>	288833
<i>Registration status:</i>	NHIG, Standard 08/12/2004
<i>Definition:</i>	Whether a specialised mental health service organisation has carer consultants employed on a paid basis to represent the interests of carers and advocate for their needs, to promote the participation of mental health carers in the planning, delivery and evaluation of the service, as represented by a code.

Data element concept attributes

<i>Data element concept:</i>	Specialised mental health service organisation – carer participation arrangements status (carer consultants employed)
<i>Definition:</i>	Whether a specialised mental health service organisation has carer consultants employed on a paid basis to represent the interests of carers and advocate for their needs, to promote the participation of mental health carers in the planning, delivery and evaluation of the service.
<i>Context:</i>	Specialised mental health services.
<i>Object class:</i>	Specialised mental health service organisation
<i>Property:</i>	Carer participation arrangements

Collection and usage attributes

<i>Guide for use:</i>	A carer is a person whose life is affected by virtue of a family or close relationship and caring role with a mental health consumer.
-----------------------	---

Value domain attributes

Representational attributes

<i>Representation class:</i>	Code								
<i>Data type:</i>	Number								
<i>Format:</i>	N								
<i>Maximum character length:</i>	1								
<i>Permissible values:</i>	<table><thead><tr><th>Value</th><th>Meaning</th></tr></thead><tbody><tr><td>1</td><td>Yes</td></tr><tr><td>2</td><td>No</td></tr><tr><td>9</td><td>Not stated</td></tr></tbody></table>	Value	Meaning	1	Yes	2	No	9	Not stated
Value	Meaning								
1	Yes								
2	No								
9	Not stated								
<i>Supplementary values:</i>									

Source and reference attributes

<i>Submitting organisation:</i>	Australian Institute of Health and Welfare
---------------------------------	--

Data element attributes

Relational attributes

Related metadata references:

See also Specialised mental health service organisation – carer participation arrangements status (formal complaints mechanism), code N NHIG, Standard 08/12/2004

See also Specialised mental health service organisation – carer participation arrangements status (carer satisfaction surveys), code N NHIG, Standard 08/12/2004

See also Specialised mental health service organisation – carer participation arrangements status (formal participation policy), code N NHIG, Standard 08/12/2004

See also Specialised mental health service organisation – carer participation arrangements status (regular discussion groups), code N NHIG, Standard 08/12/2004

Implementation in Data Set Specifications:

Mental health establishments NMDS 2005-2006 NHIG, Superseded 07/12/2005

Implementation start date: 01/07/2005

Information specific to this data set:

Obligation condition: reporting of this data element is optional for non-government residential mental health services and specialised mental health services provided by private hospitals that receive state or territory government funding.

Mental health establishments NMDS 2005-2006 NHIG, Superseded 21/03/2006

Implementation start date: 01/07/2005

Implementation end date: 30/06/2006

Information specific to this data set:

Obligation condition: reporting of this data element is optional for non-government residential mental health services and specialised mental health services provided by private hospitals that receive state or territory government funding.

Mental health establishments NMDS 2006-2007 NHIG, Superseded 23/10/2006

Implementation start date: 01/07/2006

Implementation end date: 30/06/2007

Information specific to this data set:

Obligation condition: reporting of this data element is optional for non-government residential mental health services and specialised mental health services provided by private hospitals that receive state or territory government funding.

Mental health establishments NMDS 2007-2008 NHIG, Standard 23/10/2006

Implementation start date: 01/07/2007

Information specific to this data set:

Obligation condition: reporting of this data element is

optional for non-government residential mental health services and specialised mental health services provided by private hospitals that receive state or territory government funding.

Carer participation arrangements—carer satisfaction surveys

Identifying and definitional attributes

<i>Metadata item type:</i>	Data Element
<i>Technical name:</i>	Specialised mental health service organisation – carer participation arrangements status (carer satisfaction surveys), code N
<i>METeOR identifier:</i>	290367
<i>Registration status:</i>	NHIG, Standard 08/12/2004
<i>Definition:</i>	Whether a specialised mental health service organisation periodically conducts carer satisfaction surveys, to promote the participation of mental health carers in the planning, delivery and evaluation of the service, as represented by a code.

Data element concept attributes

<i>Data element concept:</i>	Specialised mental health service organisation – carer participation arrangements status (carer satisfaction surveys)
<i>Definition:</i>	Whether a specialised mental health service organisation periodically conducts carer satisfaction surveys, to promote the participation of mental health carers in the planning, delivery and evaluation of the service.
<i>Context:</i>	Specialised mental health services.
<i>Object class:</i>	Specialised mental health service organisation
<i>Property:</i>	Carer participation arrangements

Collection and usage attributes

<i>Guide for use:</i>	A carer is a person whose life is affected by virtue of a family or close relationship and caring role with a mental health consumer.
-----------------------	---

Value domain attributes

Representational attributes

<i>Representation class:</i>	Code								
<i>Data type:</i>	Number								
<i>Format:</i>	N								
<i>Maximum character length:</i>	1								
<i>Permissible values:</i>	<table><thead><tr><th>Value</th><th>Meaning</th></tr></thead><tbody><tr><td>1</td><td>Yes</td></tr><tr><td>2</td><td>No</td></tr><tr><td>9</td><td>Not stated</td></tr></tbody></table>	Value	Meaning	1	Yes	2	No	9	Not stated
Value	Meaning								
1	Yes								
2	No								
9	Not stated								
<i>Supplementary values:</i>									

Source and reference attributes

<i>Submitting organisation:</i>	Australian Institute of Health and Welfare
---------------------------------	--

Data element attributes

Relational attributes

Related metadata references:

See also Specialised mental health service organisation – carer participation arrangements status (formal complaints mechanism), code N NHIG, Standard 08/12/2004

See also Specialised mental health service organisation – carer participation arrangements status (formal participation policy), code N NHIG, Standard 08/12/2004

See also Specialised mental health service organisation – carer participation arrangements status (regular discussion groups), code N NHIG, Standard 08/12/2004

See also Specialised mental health service organisation – carer participation arrangements status (carer consultants employed), code N NHIG, Standard 08/12/2004

Implementation in Data Set Specifications:

Mental health establishments NMDS 2005-2006 NHIG, Superseded 07/12/2005

Implementation start date: 01/07/2005

Information specific to this data set:

Obligation condition: reporting of this data element is optional for non-government residential mental health services and specialised mental health services provided by private hospitals that receive state or territory government funding.

Mental health establishments NMDS 2005-2006 NHIG, Superseded 21/03/2006

Implementation start date: 01/07/2005

Implementation end date: 30/06/2006

Information specific to this data set:

Obligation condition: reporting of this data element is optional for non-government residential mental health services and specialised mental health services provided by private hospitals that receive state or territory government funding.

Mental health establishments NMDS 2006-2007 NHIG, Superseded 23/10/2006

Implementation start date: 01/07/2006

Implementation end date: 30/06/2007

Information specific to this data set:

Obligation condition: reporting of this data element is optional for non-government residential mental health services and specialised mental health services provided by private hospitals that receive state or territory government funding.

Mental health establishments NMDS 2007-2008 NHIG, Standard 23/10/2006

Implementation start date: 01/07/2007

Information specific to this data set:

Obligation condition: reporting of this data element is optional for non-government residential mental health services and specialised mental health services provided by private hospitals that receive state or territory

government funding.

Carer participation arrangements—formal complaints mechanism

Identifying and definitional attributes

<i>Metadata item type:</i>	Data Element
<i>Technical name:</i>	Specialised mental health service organisation – carer participation arrangements status (formal complaints mechanism), code N
<i>METeOR identifier:</i>	290370
<i>Registration status:</i>	NHIG, Standard 08/12/2004
<i>Definition:</i>	Whether a specialised mental health service organisation has a formal internal complaints mechanism in which complaints made by carers are regularly reviewed by a committee that includes carers, to promote the participation of mental health carers in the planning, delivery and evaluation of the service, as represented by a code.

Data element concept attributes

<i>Data element concept:</i>	Specialised mental health service organisation – carer participation arrangements status (formal complaints mechanism)
<i>Definition:</i>	Whether a specialised mental health service organisation has a formal internal complaints mechanism in which complaints made by carers are regularly reviewed by a committee that includes carers, to promote the participation of mental health carers in the planning, delivery and evaluation of the service.
<i>Context:</i>	Specialised mental health services.
<i>Object class:</i>	Specialised mental health service organisation
<i>Property:</i>	Carer participation arrangements

Collection and usage attributes

<i>Guide for use:</i>	A carer is a person whose life is affected by virtue of a family or close relationship and caring role with a mental health consumer.
-----------------------	---

Value domain attributes

Representational attributes

<i>Representation class:</i>	Code	
<i>Data type:</i>	Number	
<i>Format:</i>	N	
<i>Maximum character length:</i>	1	
<i>Permissible values:</i>	Value	Meaning
	1	Yes
	2	No
<i>Supplementary values:</i>	9	Not stated

Source and reference attributes

Data element attributes

Relational attributes

Related metadata references:

See also Specialised mental health service organisation – carer participation arrangements status (carer satisfaction surveys), code N NHIG, Standard 08/12/2004

See also Specialised mental health service organisation – carer participation arrangements status (formal participation policy), code N NHIG, Standard 08/12/2004

See also Specialised mental health service organisation – carer participation arrangements status (regular discussion groups), code N NHIG, Standard 08/12/2004

See also Specialised mental health service organisation – carer participation arrangements status (carer consultants employed), code N NHIG, Standard 08/12/2004

Implementation in Data Set Specifications:

Mental health establishments NMDS 2005-2006 NHIG, Superseded 07/12/2005

Implementation start date: 01/07/2005

Information specific to this data set:

Obligation condition: reporting of this data element is optional for non-government residential mental health services and specialised mental health services provided by private hospitals that receive state or territory government funding.

Mental health establishments NMDS 2005-2006 NHIG, Superseded 21/03/2006

Implementation start date: 01/07/2005

Implementation end date: 30/06/2006

Information specific to this data set:

Obligation condition: reporting of this data element is optional for non-government residential mental health services and specialised mental health services provided by private hospitals that receive state or territory government funding.

Mental health establishments NMDS 2006-2007 NHIG, Superseded 23/10/2006

Implementation start date: 01/07/2006

Implementation end date: 30/06/2007

Information specific to this data set:

Obligation condition: reporting of this data element is optional for non-government residential mental health services and specialised mental health services provided by private hospitals that receive state or territory government funding.

Mental health establishments NMDS 2007-2008 NHIG, Standard 23/10/2006

Implementation start date: 01/07/2007

Information specific to this data set:

Obligation condition: reporting of this data element is optional for non-government residential mental health services and specialised mental health services provided by private hospitals that receive state or territory government funding.

Carer participation arrangements—formal participation policy

Identifying and definitional attributes

<i>Metadata item type:</i>	Data Element
<i>Technical name:</i>	Specialised mental health service organisation – carer participation arrangements status (formal participation policy), code N
<i>METeOR identifier:</i>	290365
<i>Registration status:</i>	NHIG, Standard 08/12/2004
<i>Definition:</i>	Whether a specialised mental health service organisation has developed a formal and documented policy on participation by carers, to promote the participation of mental health carers in the planning, delivery and evaluation of the service, as represented by a code.

Data element concept attributes

<i>Data element concept:</i>	Specialised mental health service organisation – carer participation arrangements status (formal participation policy)
<i>Definition:</i>	Whether a specialised mental health service organisation has developed a formal and documented policy on participation by carers, to promote the participation of mental health carers in the planning, delivery and evaluation of the service.
<i>Context:</i>	Specialised mental health services.
<i>Object class:</i>	Specialised mental health service organisation
<i>Property:</i>	Carer participation arrangements

Collection and usage attributes

<i>Guide for use:</i>	A carer is a person whose life is affected by virtue of a family or close relationship and caring role with a mental health consumer.
-----------------------	---

Value domain attributes

Representational attributes

<i>Representation class:</i>	Code						
<i>Data type:</i>	Number						
<i>Format:</i>	N						
<i>Maximum character length:</i>	1						
<i>Permissible values:</i>	<table><thead><tr><th>Value</th><th>Meaning</th></tr></thead><tbody><tr><td>1</td><td>Yes</td></tr><tr><td>2</td><td>No</td></tr></tbody></table>	Value	Meaning	1	Yes	2	No
Value	Meaning						
1	Yes						
2	No						
<i>Supplementary values:</i>	<table><tbody><tr><td>9</td><td>Not stated</td></tr></tbody></table>	9	Not stated				
9	Not stated						

Source and reference attributes

<i>Submitting organisation:</i>	Australian Institute of Health and Welfare
---------------------------------	--

Data element attributes

Relational attributes

Related metadata references:

See also Specialised mental health service organisation – carer participation arrangements status (carer consultants employed), code N NHIG, Standard 08/12/2004

See also Specialised mental health service organisation – carer participation arrangements status (regular discussion groups), code N NHIG, Standard 08/12/2004

See also Specialised mental health service organisation – carer participation arrangements status (carer satisfaction surveys), code N NHIG, Standard 08/12/2004

See also Specialised mental health service organisation – carer participation arrangements status (formal complaints mechanism), code N NHIG, Standard 08/12/2004

Implementation in Data Set Specifications:

Mental health establishments NMDS 2005-2006 NHIG, Superseded 07/12/2005

Implementation start date: 01/07/2005

Information specific to this data set:

Obligation condition: reporting of this data element is optional for non-government residential mental health services and specialised mental health services provided by private hospitals that receive state or territory government funding.

Mental health establishments NMDS 2005-2006 NHIG, Superseded 21/03/2006

Implementation start date: 01/07/2005

Implementation end date: 30/06/2006

Information specific to this data set:

Obligation condition: reporting of this data element is optional for non-government residential mental health services and specialised mental health services provided by private hospitals that receive state or territory government funding.

Mental health establishments NMDS 2006-2007 NHIG, Superseded 23/10/2006

Implementation start date: 01/07/2006

Implementation end date: 30/06/2007

Information specific to this data set:

Obligation condition: reporting of this data element is optional for non-government residential mental health services and specialised mental health services provided by private hospitals that receive state or territory government funding.

Mental health establishments NMDS 2007-2008 NHIG, Standard 23/10/2006

Implementation start date: 01/07/2007

Information specific to this data set:

Obligation condition: reporting of this data element is optional for non-government residential mental health services and specialised mental health services provided by private hospitals that receive state or territory

government funding.

Carer participation arrangements—regular discussion groups

Identifying and definitional attributes

<i>Metadata item type:</i>	Data Element
<i>Technical name:</i>	Specialised mental health service organisation – carer participation arrangements status (regular discussion groups), code N
<i>METeOR identifier:</i>	290359
<i>Registration status:</i>	NHIG, Standard 08/12/2004
<i>Definition:</i>	Whether the service holds regular discussion groups to seek the views of carers about the service, to promote the participation of mental health carers in the planning, delivery and evaluation of the service, as represented by a code.

Data element concept attributes

<i>Data element concept:</i>	Specialised mental health service organisation – carer participation arrangements status (regular discussion groups)
<i>Definition:</i>	Whether the service holds regular discussion groups to seek the views of carers about the service, to promote the participation of mental health carers in the planning, delivery and evaluation of the service.
<i>Context:</i>	Specialised mental health services.
<i>Object class:</i>	Specialised mental health service organisation
<i>Property:</i>	Carer participation arrangements

Collection and usage attributes

<i>Guide for use:</i>	A carer is a person whose life is affected by virtue of a family or close relationship and caring role with a mental health consumer.
-----------------------	---

Value domain attributes

Representational attributes

<i>Representation class:</i>	Code								
<i>Data type:</i>	Number								
<i>Format:</i>	N								
<i>Maximum character length:</i>	1								
<i>Permissible values:</i>	<table><thead><tr><th>Value</th><th>Meaning</th></tr></thead><tbody><tr><td>1</td><td>Yes</td></tr><tr><td>2</td><td>No</td></tr><tr><td>9</td><td>Not stated</td></tr></tbody></table>	Value	Meaning	1	Yes	2	No	9	Not stated
Value	Meaning								
1	Yes								
2	No								
9	Not stated								
<i>Supplementary values:</i>									

Source and reference attributes

<i>Submitting organisation:</i>	Australian Institute of Health and Welfare
---------------------------------	--

Data element attributes

Relational attributes

Related metadata references:

See also Specialised mental health service organisation – carer participation arrangements status (formal complaints mechanism), code N NHIG, Standard 08/12/2004

See also Specialised mental health service organisation – carer participation arrangements status (carer satisfaction surveys), code N NHIG, Standard 08/12/2004

See also Specialised mental health service organisation – carer participation arrangements status (formal participation policy), code N NHIG, Standard 08/12/2004

See also Specialised mental health service organisation – carer participation arrangements status (carer consultants employed), code N NHIG, Standard 08/12/2004

Implementation in Data Set Specifications:

Mental health establishments NMDS 2005-2006 NHIG, Superseded 07/12/2005

Implementation start date: 01/07/2005

Information specific to this data set:

Obligation condition: reporting of this data element is optional for non-government residential mental health services and specialised mental health services provided by private hospitals that receive state or territory government funding.

Mental health establishments NMDS 2005-2006 NHIG, Superseded 21/03/2006

Implementation start date: 01/07/2005

Implementation end date: 30/06/2006

Information specific to this data set:

Obligation condition: reporting of this data element is optional for non-government residential mental health services and specialised mental health services provided by private hospitals that receive state or territory government funding.

Mental health establishments NMDS 2006-2007 NHIG, Superseded 23/10/2006

Implementation start date: 01/07/2006

Implementation end date: 30/06/2007

Information specific to this data set:

Obligation condition: reporting of this data element is optional for non-government residential mental health services and specialised mental health services provided by private hospitals that receive state or territory government funding.

Mental health establishments NMDS 2007-2008 NHIG, Standard 23/10/2006

Implementation start date: 01/07/2007

Information specific to this data set:

Obligation condition: reporting of this data element is optional for non-government residential mental health services and specialised mental health services provided by private hospitals that receive state or territory

government funding.

Cataract - history

Identifying and definitional attributes

<i>Metadata item type:</i>	Data Element
<i>Technical name:</i>	Person – cataract status, code N
<i>METeOR identifier:</i>	270252
<i>Registration status:</i>	NHIG, Standard 01/03/2005
<i>Definition:</i>	Whether the individual has a cataract present in either or both eyes or has had a cataract previously removed from either or both eyes, as represented by a code.

Data element concept attributes

<i>Data element concept:</i>	Person – cataract status
<i>Definition:</i>	Whether the individual has a cataract present in either or both eyes or has had a cataract previously removed from either or both eyes.
<i>Context:</i>	Public health, health care and clinical settings.
<i>Object class:</i>	Person
<i>Property:</i>	Cataract status

Value domain attributes

Representational attributes

<i>Representation class:</i>	Code										
<i>Data type:</i>	Number										
<i>Format:</i>	N										
<i>Maximum character length:</i>	1										
<i>Permissible values:</i>	<table><thead><tr><th>Value</th><th>Meaning</th></tr></thead><tbody><tr><td>1</td><td>Cataract currently present or has been previously removed from the right eye</td></tr><tr><td>2</td><td>Cataract currently present or has been previously removed from the left eye</td></tr><tr><td>3</td><td>Cataract currently present or has been previously removed from both eyes</td></tr><tr><td>4</td><td>No cataract present or has not been previously removed from either eye</td></tr></tbody></table>	Value	Meaning	1	Cataract currently present or has been previously removed from the right eye	2	Cataract currently present or has been previously removed from the left eye	3	Cataract currently present or has been previously removed from both eyes	4	No cataract present or has not been previously removed from either eye
Value	Meaning										
1	Cataract currently present or has been previously removed from the right eye										
2	Cataract currently present or has been previously removed from the left eye										
3	Cataract currently present or has been previously removed from both eyes										
4	No cataract present or has not been previously removed from either eye										
<i>Supplementary values:</i>	<table><tbody><tr><td>9</td><td>Not stated/inadequately described</td></tr></tbody></table>	9	Not stated/inadequately described								
9	Not stated/inadequately described										

Data element attributes

Collection and usage attributes

<i>Collection methods:</i>	Examination of the lens of the eye through a dilated pupil (visible through the pupil by the use of an ophthalmoscope) by an ophthalmologist or optometrist, as a part of the ophthalmological assessment. Ask the individual if he/she has a cataract in either or both eyes or has had a cataract removed from either or both eyes previously. Alternatively obtain information from an
----------------------------	--

ophthalmologist or optometrist or from appropriate documentation.

Comments:

Cataract is a clouding of the lens of the eye or its capsule sufficient to reduce vision. The formation of cataract occurs more rapidly in patients with a history of ocular trauma, uveitis, or diabetes mellitus. Cataract is an associated diabetic eye problem that could lead to blindness.

Regular eye checkups are important for patients suffering from diabetes mellitus. This helps to early detect abnormalities and to avoid or postpone vision-threatening complications. A comprehensive ophthalmological examination includes:

- check visual acuity with Snellen chart -correct with pinhole if indicated
- examine for cataract
- examine fundi with pupils dilated.

Source and reference attributes

Submitting organisation:

National Diabetes Data Working Group

Origin:

National Diabetes Outcomes Quality Review Initiative (NDOQRIN) data dictionary.

Relational attributes

Related metadata references:

Supersedes Cataract - history, version 1, DE, NHDD, NHIMG, Superseded 01/03/2005

Implementation in Data Set Specifications:

Diabetes (clinical) DSS NHIG, Superseded 21/09/2005

Diabetes (clinical) DSS NHIG, Standard 21/09/2005

Category reassignment date

Identifying and definitional attributes

<i>Metadata item type:</i>	Data Element
<i>Technical name:</i>	Elective care waiting list episode – category reassignment date, DDMMYYYY
<i>METeOR identifier:</i>	270010
<i>Registration status:</i>	NHIG, Standard 01/03/2005
<i>Definition:</i>	The date on which a patient awaiting elective hospital care is assigned to a different urgency category as a result of clinical review for the awaited procedure, or is assigned to a different patient listing status category.

Data element concept attributes

<i>Data element concept:</i>	Elective care waiting list episode – category reassignment date
<i>Definition:</i>	The date on which a patient awaiting elective hospital care is assigned to a different urgency category as a result of clinical review for the awaited procedure, or is assigned to a different patient listing status category ('ready for care' or 'not ready for care').
<i>Context:</i>	Elective surgery
<i>Object class:</i>	Elective care waiting list episode
<i>Property:</i>	Category reassignment date

Value domain attributes

Representational attributes

<i>Representation class:</i>	Date
<i>Data type:</i>	Date/Time
<i>Format:</i>	DDMMYYYY
<i>Maximum character length:</i>	8

Data element attributes

Collection and usage attributes

<i>Guide for use:</i>	The date needs to be recorded each time a patient's urgency classification or listing status changes.
<i>Comments:</i>	This date is necessary for the calculation of the waiting time at admission and the waiting time at a census date.

Source and reference attributes

<i>Origin:</i>	National Health Data Committee
----------------	--------------------------------

Relational attributes

<i>Related metadata references:</i>	Supersedes Category reassignment date, version 2, DE, NHDD, NHIMG, Superseded 01/03/2005 Is used in the formation of Elective surgery waiting list episode – waiting time (at removal), total days N[NNN] NHIG,
-------------------------------------	--

Standard 01/03/2005

Is used in the formation of Elective surgery waiting list
episode – waiting time (at a census date), total days N[NNN]
NHIG, Standard 01/03/2005

Census date

Identifying and definitional attributes

<i>Metadata item type:</i>	Data Element
<i>Technical name:</i>	Hospital census (of elective surgery waitlist patients) – census date, DDMMYYYY
<i>METeOR identifier:</i>	270153
<i>Registration status:</i>	NHIG, Standard 01/03/2005
<i>Definition:</i>	Date on which the hospital takes a point in time (census) count of and characterisation of patients on the waiting list.

Data element concept attributes

<i>Data element concept:</i>	Hospital census (of elective surgery waitlist patients) – census date
<i>Definition:</i>	Date on which the hospital takes a point in time (census) count of and characterisation of patients on the waiting list.
<i>Context:</i>	Elective surgery: This metadata item is necessary for the calculation of the waiting time until a census.
<i>Object class:</i>	Hospital census
<i>Property:</i>	Census date

Value domain attributes

Representational attributes

<i>Representation class:</i>	Date
<i>Data type:</i>	Date/Time
<i>Format:</i>	DDMMYYYY
<i>Maximum character length:</i>	8

Data element attributes

Collection and usage attributes

<i>Guide for use:</i>	This date is recorded when a census is done of the patients on a waiting list.
-----------------------	--

Source and reference attributes

<i>Origin:</i>	National Health Data Committee
----------------	--------------------------------

Relational attributes

<i>Related metadata references:</i>	Supersedes Census date, version 2, DE, NHDD, NHIMG, Superseded 01/03/2005 Is used in the formation of Elective surgery waiting list episode – waiting time (at a census date), total days N[NNN] NHIG, Standard 01/03/2005
<i>Implementation in Data Set Specifications:</i>	Elective surgery waiting times (census data) NMDS NHIG, Standard 07/12/2005 <i>Implementation start date:</i> 30/09/2006

Elective surgery waiting times (census data) NMDS NHIG,
Superseded 07/12/2005

Implementation start date: 30/09/2002

Implementation end date: 30/06/2006

Centrelink customer reference number

Identifying and definitional attributes

<i>Metadata item type:</i>	Data Element
<i>Technical name:</i>	Person – government funding identifier, Centrelink customer reference number {N(9)A}
<i>Synonymous names:</i>	CRN
<i>METeOR identifier:</i>	270098
<i>Registration status:</i>	NHIG, Standard 01/03/2005
<i>Definition:</i>	A personal identifier assigned by Centrelink for the purposes of identifying people (and organisations) eligible for specific services, including some public health care services, such as oral health services.

Data element concept attributes

<i>Data element concept:</i>	Person – government funding identifier
<i>Definition:</i>	A personal identifier allocated by a government department for the purpose of identifying those eligible for specific services.
<i>Object class:</i>	Person
<i>Property:</i>	Government funding identifier

Value domain attributes

Representational attributes

<i>Representation class:</i>	Identifier
<i>Data type:</i>	String
<i>Format:</i>	{N(9)A}
<i>Maximum character length:</i>	10

Data element attributes

Collection and usage attributes

<i>Guide for use:</i>	The CRN should only be collected from persons eligible to receive health services that are to be funded by Centrelink. The number may be reported to a Centrelink agency to reconcile payment for the service provided. The data should not be used by private sector organisations for any purpose unless specifically authorised by law. For example, data linkage should not be carried out unless specifically authorised by law.
<i>Collection methods:</i>	The Centrelink Customer Reference Number (CRN) is provided on 'Health Care Cards' and 'Pensioner Concession Cards'.
<i>Comments:</i>	When a person accesses health services on the basis of being a Centrelink customer, collection of the CRN is usually necessary. This data should not be collected and recorded if it is not needed to support the provision of such health services.

Source and reference attributes

<i>Submitting organisation:</i>	Standards Australia
---------------------------------	---------------------

Origin:

AS5017 Health Care Client Identification

Relational attributes

Related metadata references:

Supersedes Centrelink customer reference number, version 1, DE, NHDD, NHIMG, Superseded 01/03/2005

Implementation in Data Set Specifications:

Health care client identification NHIG, Superseded 04/05/2005

Implementation start date: 01/01/2003

Health care client identification DSS NHIG, Standard 04/05/2005

NCSIMG, Standard 03/10/2006

Cerebral stroke due to vascular disease (history)

Identifying and definitional attributes

<i>Metadata item type:</i>	Data Element
<i>Technical name:</i>	Person – cerebral stroke due to vascular disease (history), code N
<i>METeOR identifier:</i>	270355
<i>Registration status:</i>	NHIG, Standard 01/03/2005
<i>Definition:</i>	Whether the individual has had a cerebral stroke due to vascular disease, as represented by a code.

Data element concept attributes

<i>Data element concept:</i>	Person – cerebral stroke due to vascular disease
<i>Definition:</i>	Whether the individual has had a cerebral stroke due to vascular disease.
<i>Context:</i>	Public health, health care and clinical settings.
<i>Object class:</i>	Person
<i>Property:</i>	Cerebral stroke due to vascular disease

Value domain attributes

Representational attributes

<i>Representation class:</i>	Code										
<i>Data type:</i>	Number										
<i>Format:</i>	N										
<i>Maximum character length:</i>	1										
<i>Permissible values:</i>	<table><thead><tr><th>Value</th><th>Meaning</th></tr></thead><tbody><tr><td>1</td><td>Cerebral stroke - occurred in the last 12 months</td></tr><tr><td>2</td><td>Cerebral stroke - occurred prior to the last 12 months</td></tr><tr><td>3</td><td>Cerebral stroke - occurred both in and prior to the last 12 months</td></tr><tr><td>4</td><td>No history of cerebral stroke due to vascular disease</td></tr></tbody></table>	Value	Meaning	1	Cerebral stroke - occurred in the last 12 months	2	Cerebral stroke - occurred prior to the last 12 months	3	Cerebral stroke - occurred both in and prior to the last 12 months	4	No history of cerebral stroke due to vascular disease
Value	Meaning										
1	Cerebral stroke - occurred in the last 12 months										
2	Cerebral stroke - occurred prior to the last 12 months										
3	Cerebral stroke - occurred both in and prior to the last 12 months										
4	No history of cerebral stroke due to vascular disease										
<i>Supplementary values:</i>	9 Not stated/inadequately described										

Data element attributes

Collection and usage attributes

<i>Collection methods:</i>	Obtain this information from appropriate documentation or from the patient.
----------------------------	---

Source and reference attributes

<i>Submitting organisation:</i>	National Diabetes Data Working Group
<i>Origin:</i>	National Diabetes Outcomes Quality Review Initiative (NDOQRIN) data dictionary

Relational attributes

Related metadata references:

Supersedes Cerebral stroke due to vascular disease - history, version 1, DE, NHDD, NHIMG, Superseded 01/03/2005

Implementation in Data Set Specifications:

Diabetes (clinical) DSS NHIG, Superseded 21/09/2005

Information specific to this data set:

Cerebral stroke is a medical emergency condition with a high mortality rate, which is often recognised as a vascular complication of diabetes mellitus.

The risk of stroke in patients with diabetes is at least twice that in non-diabetic patients according to Meigs et al. (Intern Med. 1998). Diabetes may increase actual stroke risk up to fivefold by increasing atheromatous deposits. Patients with diabetes who have a first stroke have 5-year survival rate reduced to 50% in comparison to non-diabetic stroke patients. The duration of diabetes clearly influences the severity of vascular disease. Atherosclerosis is more common and more severe earlier in the course of diabetes. In large arteries, plaque occurs from direct endothelial membrane injury, adverse balance of lipoproteins, and hyperinsulinemia (JAMA 1997). Small vessels are also affected more frequently than they are in non-diabetic stroke, resulting in an increased risk of lacunar stroke.

References:

Meigs J, Nathan D, Wilson P et al. Metabolic risk factors worsen continuously across the spectrum of non-diabetic glucose tolerance. *Ann Intern Med.* 1998; 128:524-533

Gorelick PB, Sacco RL, Smith DB, et al. Prevention of a first stroke: a review of guidelines and a multidisciplinary consensus statement from the National Stroke Association. *JAMA* 1999; 281:1112-1120

Diabetes (clinical) DSS NHIG, Standard 21/09/2005

Information specific to this data set:

Cerebral stroke is a medical emergency condition with a high mortality rate, which is often recognised as a vascular complication of diabetes mellitus.

The risk of stroke in patients with diabetes is at least twice that in non-diabetic patients according to Meigs et al. (Intern Med. 1998). Diabetes may increase actual stroke risk up to fivefold by increasing atheromatous deposits. Patients with diabetes who have a first stroke have 5-year survival rate reduced to 50% in comparison to non-diabetic stroke patients. The duration of diabetes clearly influences the severity of vascular disease. Atherosclerosis is more common and more severe earlier in the course of diabetes. In large arteries, plaque occurs from direct endothelial membrane injury, adverse balance of lipoproteins, and hyperinsulinemia (JAMA 1997). Small vessels are also affected more frequently than they are in non-diabetic stroke, resulting in an increased risk of lacunar stroke.

References:

Meigs J, Nathan D, Wilson P et al. Metabolic risk factors worsen continuously across the spectrum of non-diabetic glucose tolerance. Ann Intern Med. 1998; 128:524-533

Gorelick PB, Sacco RL, Smith DB, et al. Prevention of a first stroke: a review of guidelines and a multidisciplinary consensus statement from the National Stroke Association. JAMA 1999; 281:1112-1120

Change to body structure

Identifying and definitional attributes

<i>Metadata item type:</i>	Data Element
<i>Technical name:</i>	Person – nature of impairment of body structure, code (ICF 2001) N
<i>METeOR identifier:</i>	320171
<i>Registration status:</i>	NHIG, Standard 29/11/2006 NCSIMG, Standard 16/10/2006
<i>Definition:</i>	The qualitative or quantitative change of a person's impairment in a specified body structure, as represented by a code.

Data element concept attributes

<i>Data element concept:</i>	Person – nature of impairment of body structure
<i>Definition:</i>	The qualitative or quantitative change to the characteristics of a person's body structure compared with accepted population standards.
<i>Object class:</i>	Person
<i>Property:</i>	Nature of impairment of body structure

Value domain attributes

Representational attributes

<i>Classification scheme:</i>	International Classification of Functioning, Disability and Health 2001	
<i>Representation class:</i>	Code	
<i>Data type:</i>	Number	
<i>Format:</i>	N	
<i>Maximum character length:</i>	1	
<i>Permissible values:</i>	Value	Meaning
	0	No change in structure
	1	Total absence
	2	Partial absence
	3	Additional part
	4	Aberrant dimensions
	5	Discontinuity
	6	Deviating position
	7	Qualitative changes in structure
<i>Supplementary values:</i>	8	Not specified
	9	Not applicable

Collection and usage attributes

<i>Guide for use:</i>	This metadata item contributes to the definition of the concept ' Disability ' and gives an indication of the experience of disability for a person. <i>Impairments of body structure</i> are problems in body structure such as a loss or significant departure from population
-----------------------	--

standards or averages.

CODE 0 No change in structure

Used when the structure of the body part is within the range of the population standard.

CODE 1 Total absence

Used when the body structure is not present. For example total absence of the structures of the lower leg following a thorough knee amputation.

CODE 2 Partial absence

Used when only part of a body structure is present. For example partial absence of the bones of the lower leg following below knee amputation.

CODE 3 Additional part

Used when a structure, not usually present in the population is present, for example a sixth lumbar vertebra or an sixth digit on one hand.

CODE 4 Aberrant dimensions

Used when the shape and size of a body structure is significantly different from the population standard. For example radial aplasia where the shape and size of the radial bone does not develop.

CODE 5 Discontinuity

Used when parts of a body structure are separated, for example cleft palate or fracture.

CODE 6 Deviating position

Used when the location of a structure is not according to population standard; for example, transposition of the great vessels, where the aorta arises from the right ventricle and the pulmonary vessels from the left ventricle.

CODE 7 Qualitative changes in structure

Used when the structure of a body part is altered from the population standard. This includes accumulation of fluid, changes in bone structure as a result of osteoporosis or Paget's disease.

CODE 8 Not specified

Used when there is a change to a body structure, but the nature of the change is not described.

CODE 9 Not applicable

Used when it is not appropriate to code the nature of the change to a body structure.

Source and reference attributes

<i>Submitting organisation:</i>	Australian Institute of Health and Welfare (AIHW) which is the Australian Collaborating Centre for the World Health Organization Family of International Classifications.
<i>Origin:</i>	WHO 2001. ICF: International Classification of Functioning, Disability and Health. Geneva: WHO AIHW 2003. ICF Australian User Guide Version 1.0. Canberra: AIHW
<i>Reference documents:</i>	Further information on the ICF, including more detailed codes, can be found in the ICF itself and the ICF Australian User Guide (AIHW 2003), at the following websites: <ul style="list-style-type: none">• WHO ICF website

<http://www.who.int/classifications/icf/en/>

- Australian Collaborating Centre ICF website
<http://www.aihw.gov.au/disability/icf/index.html>

Data element attributes

Collection and usage attributes

Guide for use:

This data element is used in conjunction with specified body structures, for example 'partial absence of structures related to movement'. This data element may also be used in conjunction with Person – extent of impairment of body structure, code (ICF 2001) N and Person – location of impairment of body structure, code (ICF 2001) N.

Source and reference attributes

Submitting organisation:

Australian Institute of Health and Welfare (AIHW) which is the Australian Collaborating Centre for the World Health Organization Family of International Classifications.

Relational attributes

Implementation in Data Set Specifications:

Body structures cluster NHIG, Standard 29/11/2006
NCSIMG, Standard 16/10/2006

Chest pain pattern category

Identifying and definitional attributes

<i>Metadata item type:</i>	Data Element
<i>Technical name:</i>	Person – chest pain pattern, code N
<i>METeOR identifier:</i>	284823
<i>Registration status:</i>	NHIG, Standard 04/06/2004
<i>Definition:</i>	The person's chest pain pattern, as represented by a code.

Data element concept attributes

<i>Data element concept:</i>	Person – chest pain pattern
<i>Definition:</i>	Describes the person's chest pain pattern.
<i>Context:</i>	Health care and clinical settings.
<i>Object class:</i>	Person
<i>Property:</i>	Chest pain pattern

Value domain attributes

Representational attributes

<i>Representation class:</i>	Code																
<i>Data type:</i>	Number																
<i>Format:</i>	N																
<i>Maximum character length:</i>	1																
<i>Permissible values:</i>	<table><thead><tr><th>Value</th><th>Meaning</th></tr></thead><tbody><tr><td>1</td><td>Atypical chest pain</td></tr><tr><td>2</td><td>Stable chest pain pattern</td></tr><tr><td>3</td><td>Unstable chest pain pattern: rest &/or prolonged</td></tr><tr><td>4</td><td>Unstable chest pain pattern: new & severe</td></tr><tr><td>5</td><td>Unstable chest pain pattern: accelerated & severe</td></tr><tr><td>8</td><td>No chest pain/discomfort</td></tr><tr><td>9</td><td>Not stated/inadequately described</td></tr></tbody></table>	Value	Meaning	1	Atypical chest pain	2	Stable chest pain pattern	3	Unstable chest pain pattern: rest &/or prolonged	4	Unstable chest pain pattern: new & severe	5	Unstable chest pain pattern: accelerated & severe	8	No chest pain/discomfort	9	Not stated/inadequately described
Value	Meaning																
1	Atypical chest pain																
2	Stable chest pain pattern																
3	Unstable chest pain pattern: rest &/or prolonged																
4	Unstable chest pain pattern: new & severe																
5	Unstable chest pain pattern: accelerated & severe																
8	No chest pain/discomfort																
9	Not stated/inadequately described																
<i>Supplementary values:</i>																	

Collection and usage attributes

<i>Guide for use:</i>	<p>Chest pain or discomfort of myocardial ischaemic origin is usually described as chest pain, discomfort or pressure, jaw pain, arm pain or other equivalent discomfort suggestive of cardiac ischaemia. Ask the person when the symptoms first occurred or obtain this information from appropriate documentation.</p> <p>CODE 1 Atypical chest pain Use this code for pain, pressure, or discomfort in the chest, neck, or arms not clearly exertional or not otherwise consistent with pain or discomfort of myocardial ischaemic origin.</p> <p>CODE 2 Stable chest pain pattern Use this code for chest pain without a change in frequency or</p>
-----------------------	---

pattern for the 6 weeks before this presentation or procedure.
Chest pain is controlled by rest and/or
sublingual/oral/transcutaneous medications.

CODE 3 Unstable chest pain pattern: rest and/or prolonged
Use this code for chest pain that occurred at rest and was
prolonged, usually lasting more than 10 minutes

CODE 4 Unstable chest pain pattern: new and severe.
Use this code for new-onset chest pain that could be described
as at least Canadian Cardiovascular Society (CCS) classification
III severity.

CODE 5 Unstable chest pain pattern: accelerated and severe
Use this code for recent acceleration of chest pain pattern that
could be described by an increase in severity of at least 1 CCS
class to at least CCS class III.

Source and reference attributes

Submitting organisation: Australian Institute of Health and Welfare

Data element attributes

Source and reference attributes

Submitting organisation: Acute coronary syndrome data working group

Steward: The National Heart Foundation of Australia and The Cardiac
Society of Australia and New Zealand

Relational attributes

Related metadata references: Supersedes Chest pain pattern category, version 1, DE, NHDD,
NHIMG, Superseded 01/03/2005

*Implementation in Data Set
Specifications:* Acute coronary syndrome (clinical) DSS NHIG, Standard
07/12/2005

Implementation start date: 07/12/2005

Information specific to this data set:

The Canadian Cardiovascular Society classes of angina can
be used to support categorisation of chest pain patterns.
Canadian Cardiovascular Society (CCS) classes of angina
(Campeau L. Grading of angina pectoris. *Circulation* 1976;
54:522.)

1. Ordinary physical activity (for example, walking
or climbing stairs) does not cause angina; angina
occurs with strenuous or rapid or prolonged
exertion at work or recreation.
2. Slight limitation of ordinary activity (for example,
angina occurs walking or stair climbing after
meals, in cold, in wind, under emotional stress, or
only during the few hours after awakening;
walking more than 2 blocks on the level or
climbing more than 1 flight of ordinary stairs at a
normal pace; and in normal conditions).
3. Marked limitation of ordinary activity (for
example, angina occurs with walking 1 or 2 blocks
on the level or climbing 1 flight of stairs in normal
conditions and at a normal pace).

4. Inability to perform any physical activity without discomfort; angina syndrome may be present at rest.

Acute coronary syndrome (clinical) DSS NHIG, Superseded
07/12/2005

Information specific to this data set:

The Canadian Cardiovascular Society classes of angina can be used to support categorisation of chest pain patterns. Canadian Cardiovascular Society (CCS) classes of angina (Campeau L. Grading of angina pectoris. *Circulation* 1976; 54:522.)

1. Ordinary physical activity (for example, walking or climbing stairs) does not cause angina; angina occurs with strenuous or rapid or prolonged exertion at work or recreation.
2. Slight limitation of ordinary activity (for example, angina occurs walking or stair climbing after meals, in cold, in wind, under emotional stress, or only during the few hours after awakening; walking more than 2 blocks on the level or climbing more than 1 flight of ordinary stairs at a normal pace; and in normal conditions).
3. Marked limitation of ordinary activity (for example, angina occurs with walking 1 or 2 blocks on the level or climbing 1 flight of stairs in normal conditions and at a normal pace).
4. Inability to perform any physical activity without discomfort; angina syndrome may be present at rest.

Cholesterol—HDL (measured)

Identifying and definitional attributes

<i>Metadata item type:</i>	Data Element
<i>Technical name:</i>	Person – high-density lipoprotein cholesterol level (measured), total millimoles per litre [N].NN
<i>METeOR identifier:</i>	270401
<i>Registration status:</i>	NHIG, Standard 01/03/2005
<i>Definition:</i>	A person's high-density lipoprotein cholesterol (HDL-C), measured in mmol/L.

Data element concept attributes

<i>Data element concept:</i>	Person – high-density lipoprotein cholesterol level
<i>Definition:</i>	A person's high-density lipoprotein cholesterol (HDL-C) level.
<i>Context:</i>	Public health, health care and clinical settings
<i>Object class:</i>	Person
<i>Property:</i>	High-density lipoprotein cholesterol level

Value domain attributes

Representational attributes

<i>Representation class:</i>	Total				
<i>Data type:</i>	Number				
<i>Format:</i>	[N].NN				
<i>Maximum character length:</i>	3				
<i>Supplementary values:</i>	<table><thead><tr><th>Value</th><th>Meaning</th></tr></thead><tbody><tr><td>9.99</td><td>Not measured/inadequately described</td></tr></tbody></table>	Value	Meaning	9.99	Not measured/inadequately described
Value	Meaning				
9.99	Not measured/inadequately described				
<i>Unit of measure:</i>	Millimole per litre (mmol/L)				

Data element attributes

Collection and usage attributes

<i>Guide for use:</i>	<p>When reporting, record whether or not the measurement of High-density Lipoprotein Cholesterol (HDL-C) was performed in a fasting specimen.</p> <p>In settings where the monitoring of a person's health is ongoing and where a measure can change over time (such as general practice), the date of assessment should be recorded.</p>
<i>Collection methods:</i>	<p>When reporting, record absolute result of the most recent HDL-Cholesterol measurement in the last 12 months to the nearest 0.01 mmol/L.</p> <p>Measurement of lipid levels should be carried out by laboratories, or practices, which have been accredited to perform these tests by the National Association of Testing Authorities.</p> <ul style="list-style-type: none">To be collected as a single venous blood sample, preferably following a 12-hour fast where only water and medications have been consumed.

- Prolonged tourniquet use can artefactually increase levels by up to 20%.

Source and reference attributes

<i>Submitting organisation:</i>	Cardiovascular Data Working Group National Diabetes Data Working Group
<i>Origin:</i>	National Heart Foundation of Australia and the Cardiac Society of Australia and New Zealand, Lipid Management Guidelines - 2001, MJA 2001; 175: S57-S88.

Relational attributes

<i>Related metadata references:</i>	Supersedes Cholesterol-HDL - measured, version 1, DE, NHDD, NHIMG, Superseded 01/03/2005 Is used in the formation of Person—low-density lipoprotein cholesterol level (calculated), total millimoles per litre N[N].N NHIG, Standard 01/03/2005
<i>Implementation in Data Set Specifications:</i>	Acute coronary syndrome (clinical) DSS NHIG, Standard 07/12/2005 <i>Implementation start date:</i> 07/12/2005 Acute coronary syndrome (clinical) DSS NHIG, Superseded 07/12/2005 Cardiovascular disease (clinical) DSS NHIG, Superseded 15/02/2006

Information specific to this data set:

High-density Lipoprotein Cholesterol (HDL-C) is easily measured and has been shown to be a negative predictor of future coronary events.

An inverse relationship between the level of HDL-C and the risk of developing premature coronary heart disease (CHD) has been a consistent finding in a large number of prospective population studies. In many of these studies, the level of HDL-C has been the single most powerful predictor of future coronary events. Key studies of the relationship between HDLs and CHD include the Framingham Heart Study (Castelli et al. 1986), the PROCAM Study (Assman et al 1998), the Helsinki Heart Study (Manninen et al. 1992) and the MRFIT study (Stamler et al. 1986; Neaton et al 1992).

There are several well-documented functions of HDLs that may explain the ability of these lipoproteins to protect against arteriosclerosis (Barter and Rye 1996). The best recognised of these is the cholesterol efflux from cells promoted by HDLs in a process that may minimise the accumulation of foam cells in the artery wall. The major proteins of HDLs and also other proteins (e.g. paraoxonase) that co-transport with HDLs in plasma have anti-oxidant properties. Thus, HDLs have the ability to inhibit the oxidative modification of LDLs and may therefore reduce the atherogenicity of these lipoproteins.

Overall, it has been concluded from the prospective population studies that for every 0.025 mmol/L increase in HDL-C, the coronary risk is reduced by 2-5%. For a review of the relationship between HDL-C and CHD, see Barter and Rye (1996). A level below 1.0 mmol/L increases risk

approximately 2-fold (Gordon et al. 1989; Assmann et al. 1998). (Lipid Management Guidelines - 2001, MJA 2001; 175: S57-S88.

In settings such as general practice where the monitoring of a person's health is ongoing and where a measure can change over time, the Service contact date should be recorded.

Cardiovascular disease (clinical) DSS NHIG, Standard 15/02/2006

Information specific to this data set:

High-density Lipoprotein Cholesterol (HDL-C) is easily measured and has been shown to be a negative predictor of future coronary events.

An inverse relationship between the level of HDL-C and the risk of developing premature coronary heart disease (CHD) has been a consistent finding in a large number of prospective population studies. In many of these studies, the level of HDL-C has been the single most powerful predictor of future coronary events. Key studies of the relationship between HDLs and CHD include the Framingham Heart Study (Castelli et al. 1986), the PROCAM Study (Assman et al 1998), the Helsinki Heart Study (Manninen et al. 1992) and the MRFIT study (Stamler et al. 1986; Neaton et al 1992).

There are several well-documented functions of HDLs that may explain the ability of these lipoproteins to protect against arteriosclerosis (Barter and Rye 1996). The best recognised of these is the cholesterol efflux from cells promoted by HDLs in a process that may minimise the accumulation of foam cells in the artery wall. The major proteins of HDLs and also other proteins (e.g. paraoxonase) that co-transport with HDLs in plasma have anti-oxidant properties. Thus, HDLs have the ability to inhibit the oxidative modification of LDLs and may therefore reduce the atherogenicity of these lipoproteins.

Overall, it has been concluded from the prospective population studies that for every 0.025 mmol/L increase in HDL-C, the coronary risk is reduced by 2-5%. For a review of the relationship between HDL-C and CHD, see Barter and Rye (1996). A level below 1.0 mmol/L increases risk approximately 2-fold (Gordon et al. 1989; Assmann et al. 1998). (Lipid Management Guidelines - 2001, MJA 2001; 175: S57-S88.

In settings such as general practice where the monitoring of a person's health is ongoing and where a measure can change over time, the Service contact date should be recorded.

Diabetes (clinical) DSS NHIG, Superseded 21/09/2005

Information specific to this data set:

Lowered HDL-Cholesterol, with increased serum triglyceride and increased low-density lipoprotein cholesterol are important risk factors for vascular disease in type 2 diabetes.

In the New South Wales Principles of Care and Guidelines

for the Clinical Management of Diabetes Mellitus, recommendations are that HDL, total cholesterol, triglycerides are to be measured:

- every 1-2 years (if normal)
- every 3-6 months (if abnormal or on treatment)

and the target is:

- to increase HDL Cholesterol to more than or equal to 1.0 mmol/L
- to reduce total Cholesterol to less than 5.5 mmol/L
- to reduce triglyceride levels to less than 2.0 mmol/L.

If pre-existing cardiovascular disease (bypass surgery or myocardial infarction) total cholesterol should be less than 4.5 mmol/L. A level below 1.0 mmol/L increases risk approximately 2-fold (Gordon et al. 1989; Assmann et al, 1998), (Draft NHF Lipid Guidelines Paper 2001). It has been concluded from prospective population studies that for every 0.025 mmol/L increase in HDL-C, the coronary risk is reduced by 2-5%.

In settings such as general practice where the monitoring of a person's health is ongoing and where a measure can change over time, the date of assessment should be recorded.

References:

National Heart Foundation of Australia - Lipid Management Guidelines 2001.

Diabetes (clinical) DSS NHIG, Standard 21/09/2005

Information specific to this data set:

Lowered HDL-Cholesterol, with increased serum triglyceride and increased low-density lipoprotein cholesterol are important risk factors for vascular disease in type 2 diabetes.

In the New South Wales Principles of Care and Guidelines for the Clinical Management of Diabetes Mellitus, recommendations are that HDL, total cholesterol, triglycerides are to be measured:

- every 1-2 years (if normal)
- every 3-6 months (if abnormal or on treatment)

and the target is:

- to increase HDL Cholesterol to more than or equal to 1.0 mmol/L
- to reduce total Cholesterol to less than 5.5 mmol/L
- to reduce triglyceride levels to less than 2.0 mmol/L.

If pre-existing cardiovascular disease (bypass surgery or myocardial infarction) total cholesterol should be less than 4.5 mmol/L. A level below 1.0 mmol/L increases risk approximately 2-fold (Gordon et al. 1989; Assmann et al, 1998), (Draft NHF Lipid Guidelines Paper 2001). It has been concluded from prospective population studies that for every 0.025 mmol/L increase in HDL-C, the coronary risk is reduced by 2-5%.

In settings such as general practice where the monitoring of a person's health is ongoing and where a measure can change over time, the date of assessment should be

recorded.

References:

National Heart Foundation of Australia - Lipid Management Guidelines 2001.

Cholesterol—LDL (calculated)

Identifying and definitional attributes

<i>Metadata item type:</i>	Data Element
<i>Technical name:</i>	Person – low-density lipoprotein cholesterol level (calculated), total millimoles per litre N[N].N
<i>METeOR identifier:</i>	270402
<i>Registration status:</i>	NHIG, Standard 01/03/2005
<i>Definition:</i>	A person's calculated low-density lipoprotein cholesterol (LDL-C).

Data element concept attributes

<i>Data element concept:</i>	Person – low-density lipoprotein cholesterol level
<i>Definition:</i>	A person's low-density lipoprotein cholesterol (LDL-C) level.
<i>Context:</i>	Public health, health care and clinical setting.
<i>Object class:</i>	Person
<i>Property:</i>	Low-density lipoprotein cholesterol level

Value domain attributes

Representational attributes

<i>Representation class:</i>	Total				
<i>Data type:</i>	Number				
<i>Format:</i>	N[N].N				
<i>Maximum character length:</i>	3				
<i>Supplementary values:</i>	<table><thead><tr><th>Value</th><th>Meaning</th></tr></thead><tbody><tr><td>99.9</td><td>Not stated/inadequately described</td></tr></tbody></table>	Value	Meaning	99.9	Not stated/inadequately described
Value	Meaning				
99.9	Not stated/inadequately described				
<i>Unit of measure:</i>	Millimole per litre (mmol/L)				
<i>Unit of measure precision:</i>	1				

Collection and usage attributes

<i>Guide for use:</i>	Measurement in mmol/L to 1 decimal place.
-----------------------	---

Data element attributes

Collection and usage attributes

<i>Guide for use:</i>	Formula: LDL-C = (plasma total cholesterol) - (high density lipoprotein cholesterol) - (fasting plasma triglyceride divided by 2.2).
<i>Collection methods:</i>	The LDL-C is usually calculated from the Friedwald Equation (Friedwald et al. 1972), which depends on knowing the blood levels of the total cholesterol and HDL-C and the fasting level of the triglyceride. Note that the Friedwald equation becomes unreliable when the plasma triglyceride exceeds 4.5 mmol/L. Note also that while cholesterol levels are reliable for the first 24 hours after the onset of acute coronary syndromes, they may be

unreliable for the subsequent 6 weeks after an event.

- Measurement of lipid levels should be carried out by laboratories, or practices, which have been accredited to perform these tests by the National Association of Testing Authorities.
- To be collected as a single venous blood sample, preferably following a 12-hour fast where only water and medications have been consumed.

Comments:

High blood cholesterol is a key factor in heart, stroke and vascular disease, especially coronary heart disease (CHD). Poor nutrition can be a contributing factor to heart, stroke and vascular disease as a population's level of saturated fat intake is the prime determinant of its level of blood cholesterol. The majority of the cholesterol in plasma is transported as a component of LDL-C. Thus, the evidence linking CHD to plasma total cholesterol and LDL-C is essentially the same.

Source and reference attributes

Submitting organisation:

Cardiovascular Data Working Group

Origin:

National Heart Foundation of Australia and the Cardiac Society of Australia and New Zealand, Lipid Management Guidelines - 2001, MJA 2001; 175: S57-S88.

Relational attributes

Related metadata references:

Is formed using Person – cholesterol level (measured), total millimoles per litre N[N].N NHIG, Standard 01/03/2005

Is formed using Person – high-density lipoprotein cholesterol level (measured), total millimoles per litre [N].NN NHIG, Standard 01/03/2005

Is formed using Person – triglyceride level (measured), total millimoles per litre N[N].N NHIG, Standard 01/03/2005

Supersedes Cholesterol-LDL calculated, version 1, Derived DE, NHDD, NHIMG, Superseded 01/03/2005

Is formed using Health service event – fasting indicator, code N NHIG, Standard 21/09/2005

Implementation in Data Set Specifications:

Acute coronary syndrome (clinical) DSS NHIG, Standard 07/12/2005

Implementation start date: 07/12/2005

Acute coronary syndrome (clinical) DSS NHIG, Superseded 07/12/2005

Cardiovascular disease (clinical) DSS NHIG, Superseded 15/02/2006

Information specific to this data set:

Many studies have demonstrated the significance of blood cholesterol components as risk factors for heart, stroke and vascular disease.

Scientific studies have shown a continuous relationship between lipid levels and Coronary Heart Disease (CHD) and overwhelming evidence that lipid lowering interventions reduces CHD progression, morbidity and mortality.

There are many large-scale, prospective population studies

defining the relationship between plasma total (and Low-density Lipoprotein (LDL)) cholesterol and the future risk of developing CHD. The results of prospective population studies are consistent and support several general conclusions:

- the majority of people with CHD do not have markedly elevated levels of plasma total cholesterol or LDL-C,
- there is a continuous positive but curvilinear relationship between the concentration of plasma total (and LDL) cholesterol and the risk of having a coronary event and of dying from CHD,
- there is no evidence that a low level of plasma (or LDL) cholesterol predisposes to an increase in non-coronary mortality.

The excess non-coronary mortality at low cholesterol levels in the Honolulu Heart Study (Yano et al. 1983; Stemmermann et al. 1991) was apparent only in people who smoked and is consistent with a view that smokers may have occult smoking related disease that is responsible for both an increased mortality and a low plasma cholesterol.

It should be emphasised that the prospective studies demonstrate an association between plasma total cholesterol and LDL-C and the risk of developing CHD. (Lipid Management Guidelines - 2001, MJA 2001; 175: S57-S88 and Commonwealth Department of Health & Ageing and Australian Institute of Health and Welfare (1999) National Health Priority Areas Report: Cardiovascular Health 1998. AIHW Cat. No. PHE 9. HEALTH and AIHW, Canberra pgs 14-17).

In settings such as general practice where the monitoring of a person's health is ongoing and where a measure can change over time, the service contact date should be recorded.

Cardiovascular disease (clinical) DSS NHIG, Standard
15/02/2006

Information specific to this data set:

Many studies have demonstrated the significance of blood cholesterol components as risk factors for heart, stroke and vascular disease.

Scientific studies have shown a continuous relationship between lipid levels and Coronary Heart Disease (CHD) and overwhelming evidence that lipid lowering interventions reduces CHD progression, morbidity and mortality.

There are many large-scale, prospective population studies defining the relationship between plasma total (and Low-density Lipoprotein (LDL)) cholesterol and the future risk of developing CHD. The results of prospective population studies are consistent and support several general conclusions:

- the majority of people with CHD do not have markedly elevated levels of plasma total cholesterol or LDL-C,

- there is a continuous positive but curvilinear relationship between the concentration of plasma total (and LDL) cholesterol and the risk of having a coronary event and of dying from CHD,
- there is no evidence that a low level of plasma (or LDL) cholesterol predisposes to an increase in non-coronary mortality.

The excess non-coronary mortality at low cholesterol levels in the Honolulu Heart Study (Yano et al. 1983; Stemmermann et al. 1991) was apparent only in people who smoked and is consistent with a view that smokers may have occult smoking related disease that is responsible for both an increased mortality and a low plasma cholesterol.

It should be emphasised that the prospective studies demonstrate an association between plasma total cholesterol and LDL-C and the risk of developing CHD. (Lipid Management Guidelines - 2001, MJA 2001; 175: S57-S88 and Commonwealth Department of Health & Ageing and Australian Institute of Health and Welfare (1999) National Health Priority Areas Report: Cardiovascular Health 1998. AIHW Cat. No. PHE 9. HEALTH and AIHW, Canberra pgs 14-17).

In settings such as general practice where the monitoring of a person's health is ongoing and where a measure can change over time, the service contact date should be recorded.

Cholesterol—total (measured)

Identifying and definitional attributes

<i>Metadata item type:</i>	Data Element
<i>Technical name:</i>	Person – cholesterol level (measured), total millimoles per litre N[N].N
<i>METeOR identifier:</i>	270403
<i>Registration status:</i>	NHIG, Standard 01/03/2005
<i>Definition:</i>	A person's total cholesterol (TC), measured in mmol/L.

Data element concept attributes

<i>Data element concept:</i>	Person – cholesterol level
<i>Definition:</i>	A person's total cholesterol (TC).
<i>Context:</i>	Public health, health care and clinical settings
<i>Object class:</i>	Person
<i>Property:</i>	Cholesterol level

Value domain attributes

Representational attributes

<i>Representation class:</i>	Total				
<i>Data type:</i>	Number				
<i>Format:</i>	N[N].N				
<i>Maximum character length:</i>	3				
<i>Supplementary values:</i>	<table><thead><tr><th>Value</th><th>Meaning</th></tr></thead><tbody><tr><td>99.9</td><td>Not stated/inadequately described.</td></tr></tbody></table>	Value	Meaning	99.9	Not stated/inadequately described.
Value	Meaning				
99.9	Not stated/inadequately described.				
<i>Unit of measure:</i>	Millimole per litre (mmol/L)				

Data element attributes

Collection and usage attributes

<i>Guide for use:</i>	Measurement in mmol/L to 1 decimal place. Record the absolute result of the total cholesterol measurement. When reporting, record whether or not the measurement of Cholesterol-total - measured was performed in a fasting specimen.
<i>Collection methods:</i>	When reporting, record absolute result of the most recent Cholesterol-total - measured in the last 12 months to the nearest 0.1 mmol/L. Measurement of lipid levels should be carried out by laboratories, or practices, which have been accredited to perform these tests by the National Association of Testing Authorities. <ul style="list-style-type: none">To be collected as a single venous blood sample, preferably following a 12-hour fast where only water and medications have been consumed.Prolonged tourniquet use can artifactually increase levels by up to 20%.

Comments:

In settings where the monitoring of a person's health is ongoing and where a measure can change over time (such as general practice), the Service contact – service contact date, DDMMYYYY should be recorded.

High blood cholesterol is a key factor in heart, stroke and vascular disease, especially coronary heart disease.

Poor nutrition can be a contributing factor to heart, stroke and vascular disease as a population's level of saturated fat intake is the prime determinant of its level of blood cholesterol.

Large clinical trials have shown that people at highest risk of cardiovascular events (e.g. pre-existing ischaemic heart disease) will derive the greatest benefit from lipid lowering drugs. For this group of patients, the optimum threshold plasma lipid concentration for drug treatment is still a matter of research. In May 1999 the PBS threshold total cholesterol concentration, for subsidy of drug treatment, was reduced from 5.5 to 4.0 mmol/L. (Australian Medical Handbook).

Source and reference attributes

Submitting organisation:

Cardiovascular Data Working Group

Origin:

National Heart Foundation of Australia and the Cardiac Society of Australia and New Zealand, Lipid Management Guidelines - 2001, MJA 2001; 175: S57-S88

National Health Priority Areas Report: Cardiovascular Health 1998. AIHW Cat. No. PHE 9. HEALTH and AIHW, Canberra.

The Royal College of Pathologists of Australasia web based Manual of Use and Interpretation of Pathology Tests

Relational attributes

Related metadata references:

Supersedes Cholesterol-total - measured, version 1, DE, NHDD, NHIMG, Superseded 01/03/2005

Is used in the formation of Person – low-density lipoprotein cholesterol level (calculated), total millimoles per litre N[N].N NHIG, Standard 01/03/2005

Implementation in Data Set Specifications:

Acute coronary syndrome (clinical) DSS NHIG, Standard 07/12/2005

Implementation start date: 07/12/2005

Acute coronary syndrome (clinical) DSS NHIG, Superseded 07/12/2005

Cardiovascular disease (clinical) DSS NHIG, Superseded 15/02/2006

Information specific to this data set:

Scientific studies have shown a continuous relationship between lipid levels and coronary heart disease and overwhelming evidence that lipid lowering interventions reduce coronary heart disease progression, morbidity and mortality. Studies show a positive relationship between an individual's total blood cholesterol level and risk of coronary heart disease as well as death (Kannel & Gordon 1970; Pocock et al. 1989).

Many studies have demonstrated the significance of blood cholesterol components as risk factors for heart, stroke and vascular disease.

Several generalisations can be made from these cholesterol lowering trials:

- that the results of the intervention trials are consistent with the prospective population studies in which (excluding possible regression dilution bias) a 1.0 mmol/L reduction in plasma total cholesterol translates into an approximate 20% reduction in the risk of future coronary events.
- It should be emphasised, however, that this conclusion does not necessarily apply beyond the range of cholesterol levels which have been tested in these studies.
- That the benefits of cholesterol lowering are apparent in people with and without coronary artery disease.

There is high level evidence that in patients with existing coronary heart disease, lipid intervention therapy reduces the risk of subsequent stroke

Cardiovascular disease (clinical) DSS NHIG, Standard 15/02/2006

Information specific to this data set:

Scientific studies have shown a continuous relationship between lipid levels and coronary heart disease and overwhelming evidence that lipid lowering interventions reduce coronary heart disease progression, morbidity and mortality. Studies show a positive relationship between an individual's total blood cholesterol level and risk of coronary heart disease as well as death (Kannel & Gordon 1970; Pocock et al. 1989).

Many studies have demonstrated the significance of blood cholesterol components as risk factors for heart, stroke and vascular disease.

Several generalisations can be made from these cholesterol lowering trials:

- that the results of the intervention trials are consistent with the prospective population studies in which (excluding possible regression dilution bias) a 1.0 mmol/L reduction in plasma total cholesterol translates into an approximate 20% reduction in the risk of future coronary events.
- It should be emphasised, however, that this conclusion does not necessarily apply beyond the range of cholesterol levels which have been tested in these studies.
- That the benefits of cholesterol lowering are apparent in people with and without coronary artery disease.

There is high level evidence that in patients with existing coronary heart disease, lipid intervention therapy reduces the risk of subsequent stroke

Diabetes (clinical) DSS NHIG, Superseded 21/09/2005

Information specific to this data set:

The risk of coronary and other macrovascular disorders is 2-5 times higher in people with diabetes than in non-diabetic subjects and increases in parallel with the degree

of dyslipidaemia.

Following Principles of Care and Guidelines for the Clinical Management of Diabetes Mellitus, the targets for lipids management are:

- To reduce total Cholesterols to less than 5.5 mmol/L
- To reduce triglyceride levels to less than 2.0 mmol/L
- To increase high density lipoprotein Cholesterols to more than or equal to 1.0 mmol/L.

If pre-existing cardiovascular disease (bypass surgery or myocardial infarction), total cholesterol should be less than 4.5 mmol/L

Diabetes (clinical) DSS NHIG, Standard 21/09/2005

Information specific to this data set:

The risk of coronary and other macrovascular disorders is 2-5 times higher in people with diabetes than in non-diabetic subjects and increases in parallel with the degree of dyslipidaemia.

Following Principles of Care and Guidelines for the Clinical Management of Diabetes Mellitus, the targets for lipids management are:

- To reduce total Cholesterols to less than 5.5 mmol/L
- To reduce triglyceride levels to less than 2.0 mmol/L
- To increase high density lipoprotein Cholesterols to more than or equal to 1.0 mmol/L.

If pre-existing cardiovascular disease (bypass surgery or myocardial infarction), total cholesterol should be less than 4.5 mmol/L

Classification of health labour force job

Identifying and definitional attributes

<i>Metadata item type:</i>	Data Element
<i>Technical name:</i>	Health professional – occupation, code ANN
<i>METeOR identifier:</i>	270140
<i>Registration status:</i>	NHIG, Standard 01/03/2005
<i>Definition:</i>	The position or job classification of a health professional, as represented by a code.

Data element concept attributes

<i>Data element concept:</i>	Health professional – occupation
<i>Definition:</i>	Position or job classification is a broad description of the roles and levels within a general organisational or industrial structure for health professions, and classifications vary among the professions according to organisational arrangements.
<i>Context:</i>	Health labour force
<i>Object class:</i>	Health professional
<i>Property:</i>	Occupation

Value domain attributes

Representational attributes

<i>Representation class:</i>	Code																										
<i>Data type:</i>	String																										
<i>Format:</i>	ANN																										
<i>Maximum character length:</i>	3																										
<i>Permissible values:</i>	<table><thead><tr><th>Value</th><th>Meaning</th></tr></thead><tbody><tr><td>A01</td><td>Medicine - General practitioner working mainly in general practice</td></tr><tr><td>A02</td><td>Medicine - General practitioner working mainly in a special interest area</td></tr><tr><td>A03</td><td>Medicine - Salaried non-specialist hospital practitioner: Resident medical officer or intern</td></tr><tr><td>A04</td><td>Medicine - Salaried non-specialist hospital practitioner: other hospital career medical officer</td></tr><tr><td>A05</td><td>Medicine - Specialist</td></tr><tr><td>A06</td><td>Medicine - Specialist in training (e.g. registrar)</td></tr><tr><td>B01</td><td>Dentistry (private practice only) - Solo practitioner</td></tr><tr><td>B02</td><td>Dentistry (private practice only) - Solo principal with assistant(s)</td></tr><tr><td>B03</td><td>Dentistry (private practice only) - Partnership</td></tr><tr><td>B04</td><td>Dentistry (private practice only) - Associateship</td></tr><tr><td>B05</td><td>Dentistry (private practice only) - Assistant</td></tr><tr><td>B06</td><td>Dentistry (private practice only) - Locum</td></tr></tbody></table>	Value	Meaning	A01	Medicine - General practitioner working mainly in general practice	A02	Medicine - General practitioner working mainly in a special interest area	A03	Medicine - Salaried non-specialist hospital practitioner: Resident medical officer or intern	A04	Medicine - Salaried non-specialist hospital practitioner: other hospital career medical officer	A05	Medicine - Specialist	A06	Medicine - Specialist in training (e.g. registrar)	B01	Dentistry (private practice only) - Solo practitioner	B02	Dentistry (private practice only) - Solo principal with assistant(s)	B03	Dentistry (private practice only) - Partnership	B04	Dentistry (private practice only) - Associateship	B05	Dentistry (private practice only) - Assistant	B06	Dentistry (private practice only) - Locum
Value	Meaning																										
A01	Medicine - General practitioner working mainly in general practice																										
A02	Medicine - General practitioner working mainly in a special interest area																										
A03	Medicine - Salaried non-specialist hospital practitioner: Resident medical officer or intern																										
A04	Medicine - Salaried non-specialist hospital practitioner: other hospital career medical officer																										
A05	Medicine - Specialist																										
A06	Medicine - Specialist in training (e.g. registrar)																										
B01	Dentistry (private practice only) - Solo practitioner																										
B02	Dentistry (private practice only) - Solo principal with assistant(s)																										
B03	Dentistry (private practice only) - Partnership																										
B04	Dentistry (private practice only) - Associateship																										
B05	Dentistry (private practice only) - Assistant																										
B06	Dentistry (private practice only) - Locum																										

C01	Nursing - Enrolled nurse
C02	Nursing - Registered nurse
C03	Nursing - Clinical nurse
C04	Nursing - Clinical nurse consultant/supervisor
C05	Nursing - Nurse manager
C06	Nursing - Nurse educator
C07	Nursing - Nurse researcher
C08	Nursing - Assistant director of nursing
C09	Nursing - Deputy director of nursing
C10	Nursing - Director of nursing
C11	Nursing - Tutor/lecturer/senior lecturer in nursing (tertiary institution)
C12	Nursing - Associate professor/professor in nursing (tertiary institution)
C98	Nursing - Other (specify)
D01	Pharmacy (community pharmacist) - Sole proprietor
D02	Pharmacy (community pharmacist) - Partner-proprietor
D03	Pharmacy (community pharmacist) - Pharmacist-in-charge
D04	Pharmacy (community pharmacist) - Permanent assistant
D05	Pharmacy (community pharmacist) - Reliever, regular location
D06	Pharmacy (community pharmacist) - Reliever, various locations
E01	Pharmacy (Hospital/clinic pharmacist) - Director/deputy director
E02	Pharmacy (Hospital/clinic pharmacist) - Grade III pharmacist
E03	Pharmacy (Hospital/clinic pharmacist) - Grade II pharmacist
E04	Pharmacy (Hospital/clinic pharmacist) - Grade I pharmacist
E05	Pharmacy (Hospital/clinic pharmacist) - Sole pharmacist
F01	Podiatry - Own practice (or partnership)
F02	Podiatry - Own practice and sessional appointments elsewhere
F03	Podiatry - Own practice and fee-for-service elsewhere
F04	Podiatry - Own practice, sessional and fee-for-service appointments elsewhere
F05	Podiatry - Salaried podiatrist
F06	Podiatry - Locum, regular location
F07	Podiatry - Locum, various locations
F08	Podiatry - Other (specify)

	G01	Physiotherapy - Own practice (or partnership)
	G02	Physiotherapy - Own practice and sessional appointments elsewhere
	G03	Physiotherapy - Own practice and fee-for-service elsewhere
	G04	Physiotherapy - Own practice, sessional and fee-for-service appointments elsewhere
	G05	Physiotherapy - Salaried physiotherapist
	G06	Physiotherapy - Locum, regular location
	G07	Physiotherapy - Locum, various locations
<i>Supplementary values:</i>	C99	Nursing - Unknown/inadequately described/not stated

Data element attributes

Collection and usage attributes

Comments: Position or job classifications are specific to each profession and may differ by state or territory. The classifications above are simplified so that comparable data presentation is possible and possible confounding effects of enterprise specific structures are avoided. For example, for medicine, the job classification collected in the national health labour force collection is very broad. State/territory health authorities have more detailed classifications for salaried medical practitioners in hospitals. These classifications separate interns, the resident medical officer levels, registrar levels, career medical officer positions, and supervisory positions including clinical and medical superintendents. Space restrictions do not at present permit these classes to be included in the National Health Labour Force Collection questionnaire.

Source and reference attributes

Submitting organisation: National Health Labour Force Data Working Group

Relational attributes

Related metadata references: Supersedes Classification of health labour force job, version 1, DE, NHDD, NHIMG, Superseded 01/03/2005

Implementation in Data Set Specifications: Health labour force NMDS NHIG, Standard 01/03/2005

Implementation start date: 01/07/2006

Information specific to this data set:

Distribution of a professional labour force across job classification categories cross-classified with other variables allows analysis of:

- career progression
- age and gender distribution
- imputed salary/wage distribution

Client type (alcohol and other drug treatment services)

Identifying and definitional attributes

<i>Metadata item type:</i>	Data Element
<i>Technical name:</i>	Episode of treatment for alcohol and other drugs – client type, code N
<i>METeOR identifier:</i>	270083
<i>Registration status:</i>	NHIG, Standard 01/03/2005
<i>Definition:</i>	The status of a person in terms of whether the treatment episode concerns their own alcohol and/or other drug use or that of another person, as represented by a code.

Data element concept attributes

<i>Data element concept:</i>	Episode of treatment for alcohol and other drugs – client type
<i>Definition:</i>	The status of a person in terms of whether the treatment episode concerns their own alcohol and/or other drug use or that of another person.
<i>Context:</i>	Alcohol and other drug treatment services
<i>Object class:</i>	Episode of treatment for alcohol and other drugs
<i>Property:</i>	Client type

Value domain attributes

Representational attributes

<i>Representation class:</i>	Code						
<i>Data type:</i>	Number						
<i>Format:</i>	N						
<i>Maximum character length:</i>	1						
<i>Permissible values:</i>	<table><thead><tr><th>Value</th><th>Meaning</th></tr></thead><tbody><tr><td>1</td><td>Own alcohol or other drug use</td></tr><tr><td>2</td><td>Other's alcohol or other drug use</td></tr></tbody></table>	Value	Meaning	1	Own alcohol or other drug use	2	Other's alcohol or other drug use
Value	Meaning						
1	Own alcohol or other drug use						
2	Other's alcohol or other drug use						

Collection and usage attributes

<i>Guide for use:</i>	<p>CODE 1 Own alcohol or other drug use Use this code for a client who receives treatment or assistance concerning their own alcohol and/or other drug use. Use this code where a client is receiving treatment or assistance for both their own alcohol and/or other drug use and the alcohol and/or other drug use of another person.</p> <p>CODE 2 Other's alcohol or other drug use Use this code for a client who receives support and/or assistance in relation to the alcohol and/or other drug use of another person.</p>
<i>Collection methods:</i>	To be collected on commencement of a treatment episode with a service.

Data element attributes

Collection and usage attributes

<i>Guide for use:</i>	Where Code 2 Other's alcohol or other drug use is reported, do not collect the following data elements: Episode of treatment for alcohol and other drugs – drug of concern (principal), code (ASCDC 2000 extended) NNNN; Episode of treatment for alcohol and other drugs – drug of concern (other), code (ASCDC 2000 extended) NNNN; Client – injecting drug use status, code N; and Client – method of drug use (principal drug of concern), code N.
<i>Comments:</i>	Required to differentiate between clients according to whether the treatment episode concerns their own alcohol and/or other drug use or that of another person to provide a basis for description of the people accessing alcohol and other drug treatment services.

Source and reference attributes

<i>Submitting organisation:</i>	Intergovernmental Committee on Drugs National Minimum Data Set Working Group
---------------------------------	--

Relational attributes

<i>Related metadata references:</i>	Supersedes Client type - alcohol and other drug treatment services, version 3, DE, NHDD, NHIMG, Superseded 01/03/2005
<i>Implementation in Data Set Specifications:</i>	Alcohol and other drug treatment services NMDS NHIG, Superseded 21/03/2006 <i>Implementation start date:</i> 01/07/2005 <i>Implementation end date:</i> 30/06/2006 Alcohol and other drug treatment services NMDS NHIG, Superseded 23/10/2006 <i>Implementation start date:</i> 01/07/2006 <i>Implementation end date:</i> 30/06/2007 Alcohol and other drug treatment services NMDS 2007-2008 NHIG, Standard 23/10/2006 <i>Implementation start date:</i> 01/07/2007

Clinical evidence of chronic lung disease (status)

Identifying and definitional attributes

<i>Metadata item type:</i>	Data Element
<i>Technical name:</i>	Person – clinical evidence status (chronic lung disease), code N
<i>METeOR identifier:</i>	285285
<i>Registration status:</i>	NHIG, Standard 04/06/2004
<i>Definition:</i>	The status of evidence for a pre-existing clinical condition of chronic lung disease, as represented by a code.

Data element concept attributes

<i>Data element concept:</i>	Person – clinical evidence status (chronic lung disease)
<i>Definition:</i>	Indicator of the status of evidence for a pre-existing clinical condition of chronic lung disease.
<i>Context:</i>	Acute coronary treatment settings.
<i>Object class:</i>	Person
<i>Property:</i>	Clinical evidence status

Value domain attributes

Representational attributes

<i>Representation class:</i>	Code						
<i>Data type:</i>	Number						
<i>Format:</i>	N						
<i>Maximum character length:</i>	1						
<i>Permissible values:</i>	<table><thead><tr><th>Value</th><th>Meaning</th></tr></thead><tbody><tr><td>1</td><td>Objective evidence</td></tr><tr><td>2</td><td>No objective evidence</td></tr></tbody></table>	Value	Meaning	1	Objective evidence	2	No objective evidence
Value	Meaning						
1	Objective evidence						
2	No objective evidence						

Source and reference attributes

<i>Submitting organisation:</i>	Australian Institute of Health and Welfare
---------------------------------	--

Data element attributes

Collection and usage attributes

<i>Guide for use:</i>	Objective evidence is coded where the diagnosis is supported by current use of chronic lung disease pharmacological therapy (e.g. inhalers, theophylline, aminophylline, or steroids), or a forced expiratory volume in 1 second (FEV1) less than 80% predicted FEV1/forced vital capacity (FVC) less than 0.7 (post bronchodilator). Respiratory failure partial pressure of oxygen (PaO ₂) less than 60 mmHg (8kPa), or partial pressure of carbon dioxide (PaCO ₂) greater than 50 mmHg (6.7 kPa).
<i>Collection methods:</i>	For each Person – concurrent clinical condition (acute coronary syndrome), code NN, the data elements Person – clinical evidence status(chronic lung disease), code N; Person – clinical evidence status(heart failure), code N; Person – clinical evidence status(stroke), code N; Person – clinical evidence

status(peripheral arterial disease), code N; Person— clinical evidence status(sleep apnoea syndrome), code N must also be recorded.

Comments:

The diagnosis rests on the airflow limitation, which is not fully reversible. Consider treating as asthma if airflow limitation is substantially reversible. (The Thoracic Society of Australia & New Zealand and the Australian Lung Foundation, Chronic Obstructive Pulmonary Disease (COPD) Australian & New Zealand Management Guidelines and the COPD Handbook. Version 1, November 2002.)

Source and reference attributes

Submitting organisation:

Acute coronary syndrome data working group

Steward:

The National Heart Foundation of Australia and The Cardiac Society of Australia and New Zealand

Relational attributes

Related metadata references:

Supersedes Clinical evidence status, version 1, DE, NHDD, NHIMG, Superseded 01/03/2005

Implementation in Data Set Specifications:

Acute coronary syndrome (clinical) DSS NHIG, Standard 07/12/2005

Implementation start date: 07/12/2005

Information specific to this data set:

This data element seeks to ensure that patients with self-reported past symptoms pertinent to acute coronary syndrome, have objective evidence supporting reported diagnoses, using current medical practice.

Acute coronary syndrome (clinical) DSS NHIG, Superseded 07/12/2005

Information specific to this data set:

This data element seeks to ensure that patients with self-reported past symptoms pertinent to acute coronary syndrome, have objective evidence supporting reported diagnoses, using current medical practice.

Clinical evidence of heart failure (status)

Identifying and definitional attributes

<i>Metadata item type:</i>	Data Element
<i>Technical name:</i>	Person – clinical evidence status (heart failure), code N
<i>METeOR identifier:</i>	285287
<i>Registration status:</i>	NHIG, Standard 04/06/2004
<i>Definition:</i>	The status of evidence for a pre-existing clinical condition of heart failure, as represented by a code.

Data element concept attributes

<i>Data element concept:</i>	Person – clinical evidence status (heart failure)
<i>Definition:</i>	Indicator of the status of evidence for a pre-existing clinical condition of heart failure.
<i>Context:</i>	Acute coronary treatment settings.
<i>Object class:</i>	Person
<i>Property:</i>	Clinical evidence status

Value domain attributes

Representational attributes

<i>Representation class:</i>	Code						
<i>Data type:</i>	Number						
<i>Format:</i>	N						
<i>Maximum character length:</i>	1						
<i>Permissible values:</i>	<table><thead><tr><th>Value</th><th>Meaning</th></tr></thead><tbody><tr><td>1</td><td>Objective evidence</td></tr><tr><td>2</td><td>No objective evidence</td></tr></tbody></table>	Value	Meaning	1	Objective evidence	2	No objective evidence
Value	Meaning						
1	Objective evidence						
2	No objective evidence						

Source and reference attributes

<i>Submitting organisation:</i>	Australian Institute of Health and Welfare
---------------------------------	--

Data element attributes

Collection and usage attributes

<i>Guide for use:</i>	Objective evidence is coded where a patient has current symptoms of heart failure (typically breathlessness or fatigue), either at rest or during exercise and/or signs of pulmonary or peripheral congestion and objective evidence of cardiac dysfunction at rest. The diagnosis is derived from and substantiated by clinical documentation from testing according to current practices.
<i>Collection methods:</i>	For each Person – concurrent clinical condition (acute coronary syndrome), code NN, the data elements Person – clinical evidence status(chronic lung disease), code N; Person – clinical evidence status(heart failure), code N; Person – clinical evidence status(stroke), code N; Person – clinical evidence status(peripheral arterial disease), code N; Person – clinical

evidence status(sleep apnoea syndrome), code N must also be recorded.

Comments:

The most widely available investigation for documenting left ventricular dysfunction is the transthoracic echocardiogram (TTE).

Other modalities include:

- transoesophageal echocardiography (TOE),
- radionuclide ventriculography (RVG),
- left ventriculogram (LVgram),
- magnetic resonance imaging (MRI).

In the absence of any adjunctive laboratory tests, evidence of supportive clinical signs of ventricular dysfunction. These include:

- third heart sound (S3),
- cardiomegaly,
- elevated jugular venous pressure (JVP),
- chest X-ray evidence of pulmonary congestion.

Source and reference attributes

Submitting organisation:

Acute coronary syndrome data working group

Steward:

The National Heart Foundation of Australia and The Cardiac Society of Australia and New Zealand

Relational attributes

Related metadata references:

Supersedes Clinical evidence status, version 1, DE, NHDD, NHIMG, Superseded 01/03/2005

Implementation in Data Set Specifications:

Acute coronary syndrome (clinical) DSS NHIG, Standard 07/12/2005

Implementation start date: 07/12/2005

Information specific to this data set:

This data element seeks to ensure that patients with self-reported past symptoms pertinent to acute coronary syndrome, have objective evidence supporting reported diagnoses, using current medical practice.

Acute coronary syndrome (clinical) DSS NHIG, Superseded 07/12/2005

Information specific to this data set:

This data element seeks to ensure that patients with self-reported past symptoms pertinent to acute coronary syndrome, have objective evidence supporting reported diagnoses, using current medical practice.

Clinical evidence of peripheral arterial disease (status)

Identifying and definitional attributes

<i>Metadata item type:</i>	Data Element
<i>Technical name:</i>	Person – clinical evidence status (peripheral arterial disease), code N
<i>METeOR identifier:</i>	285289
<i>Registration status:</i>	NHIG, Standard 04/06/2004
<i>Definition:</i>	The status of evidence for a pre-existing clinical condition of peripheral arterial disease, as represented by a code.

Data element concept attributes

<i>Data element concept:</i>	Person – clinical evidence status (peripheral arterial disease)
<i>Definition:</i>	Indicator of the status of evidence for a pre-existing clinical condition of peripheral arterial disease.
<i>Context:</i>	Acute coronary treatment settings.
<i>Object class:</i>	Person
<i>Property:</i>	Clinical evidence status

Value domain attributes

Representational attributes

<i>Representation class:</i>	Code						
<i>Data type:</i>	Number						
<i>Format:</i>	N						
<i>Maximum character length:</i>	1						
<i>Permissible values:</i>	<table><thead><tr><th>Value</th><th>Meaning</th></tr></thead><tbody><tr><td>1</td><td>Objective evidence</td></tr><tr><td>2</td><td>No objective evidence</td></tr></tbody></table>	Value	Meaning	1	Objective evidence	2	No objective evidence
Value	Meaning						
1	Objective evidence						
2	No objective evidence						

Source and reference attributes

<i>Submitting organisation:</i>	Australian Institute of Health and Welfare
---------------------------------	--

Data element attributes

Collection and usage attributes

<i>Guide for use:</i>	<p>For peripheral artery disease, objective evidence is coded where the diagnosis is derived from and substantiated by clinical documentation for a patient with a history of either chronic or acute occlusion or narrowing of the arterial lumen in the aorta or extremities.</p> <p>For aortic aneurysm, objective evidence is coded when the diagnosis of aneurysmal dilatation of the aorta (thoracic and or abdominal) is supported and substantiated by appropriate documentation of objective testing.</p> <p>For renal artery stenosis, objective evidence is coded when the diagnosis of functional stenosis of one or both renal arteries is present and is supported and substantiated by appropriate</p>
-----------------------	---

Collection methods:

documentation of objective testing.

For each Person – concurrent clinical condition (acute coronary syndrome), code NN, the data elements Person – clinical evidence status (chronic lung disease), code N; Person – clinical evidence status (heart failure), code N; Person – clinical evidence status (stroke), code N; Person – clinical evidence status (peripheral arterial disease), code N; Person – clinical evidence status (sleep apnoea syndrome), code N must also be recorded.

Source and reference attributes

Submitting organisation:

Acute coronary syndrome data working group

Steward:

The National Heart Foundation of Australia and The Cardiac Society of Australia and New Zealand

Relational attributes

Related metadata references:

Supersedes Clinical evidence status, version 1, DE, NHDD, NHIMG, Superseded 01/03/2005

Implementation in Data Set Specifications:

Acute coronary syndrome (clinical) DSS NHIG, Standard 07/12/2005

Implementation start date: 07/12/2005

Information specific to this data set:

This data element seeks to ensure that patients with self-reported past symptoms pertinent to acute coronary syndrome, have objective evidence supporting reported diagnoses, using current medical practice.

Acute coronary syndrome (clinical) DSS NHIG, Superseded 07/12/2005

Information specific to this data set:

This data element seeks to ensure that patients with self-reported past symptoms pertinent to acute coronary syndrome, have objective evidence supporting reported diagnoses, using current medical practice.

Clinical evidence of sleep apnoea syndrome (status)

Identifying and definitional attributes

<i>Metadata item type:</i>	Data Element
<i>Technical name:</i>	Person – clinical evidence status (sleep apnoea syndrome), code N
<i>METeOR identifier:</i>	285291
<i>Registration status:</i>	NHIG, Standard 04/06/2004
<i>Definition:</i>	The status of evidence for a pre-existing clinical condition of sleep apnoea syndrome, as represented by a code.

Data element concept attributes

<i>Data element concept:</i>	Person – clinical evidence status (sleep apnoea syndrome)
<i>Definition:</i>	Indicator of the status of evidence for a pre-existing clinical condition of sleep apnoea syndrome.
<i>Context:</i>	Acute coronary treatment settings.
<i>Object class:</i>	Person
<i>Property:</i>	Clinical evidence status

Value domain attributes

Representational attributes

<i>Representation class:</i>	Code						
<i>Data type:</i>	Number						
<i>Format:</i>	N						
<i>Maximum character length:</i>	1						
<i>Permissible values:</i>	<table><thead><tr><th>Value</th><th>Meaning</th></tr></thead><tbody><tr><td>1</td><td>Objective evidence</td></tr><tr><td>2</td><td>No objective evidence</td></tr></tbody></table>	Value	Meaning	1	Objective evidence	2	No objective evidence
Value	Meaning						
1	Objective evidence						
2	No objective evidence						

Source and reference attributes

<i>Submitting organisation:</i>	Australian Institute of Health and Welfare
---------------------------------	--

Data element attributes

Collection and usage attributes

<i>Guide for use:</i>	Objective evidence is coded where the diagnosis is derived from and substantiated by clinical documentation of sleep apnoea syndrome (SAS). SAS has been diagnosed from the results of a sleep study.
<i>Collection methods:</i>	For each Person – concurrent clinical condition (acute coronary syndrome), code NN, the data elements Person – clinical evidence status(chronic lung disease), code N; Person – clinical evidence status(heart failure), code N; Person – clinical evidence status(stroke), code N; Person – clinical evidence status(peripheral arterial disease), code N; Person – clinical evidence status(sleep apnoea syndrome), code N must also be recorded.

Source and reference attributes

Submitting organisation: Acute coronary syndrome data working group
Steward: The National Heart Foundation of Australia and The Cardiac Society of Australia and New Zealand

Relational attributes

Related metadata references: Supersedes Clinical evidence status, version 1, DE, NHDD, NHIMG, Superseded 01/03/2005

Implementation in Data Set Specifications: Acute coronary syndrome (clinical) DSS NHIG, Standard 07/12/2005

Implementation start date: 07/12/2005

Information specific to this data set:

This data element seeks to ensure that patients with self-reported past symptoms pertinent to acute coronary syndrome, have objective evidence supporting reported diagnoses, using current medical practice.

Acute coronary syndrome (clinical) DSS NHIG, Superseded 07/12/2005

Information specific to this data set:

This data element seeks to ensure that patients with self-reported past symptoms pertinent to acute coronary syndrome, have objective evidence supporting reported diagnoses, using current medical practice.

Clinical evidence of stroke (status)

Identifying and definitional attributes

<i>Metadata item type:</i>	Data Element
<i>Technical name:</i>	Person – clinical evidence status (stroke), code N
<i>METeOR identifier:</i>	285293
<i>Registration status:</i>	NHIG, Standard 04/06/2004
<i>Definition:</i>	The status of evidence for a pre-existing clinical condition of stroke, as represented by a code.

Data element concept attributes

<i>Data element concept:</i>	Person – clinical evidence status (stroke)
<i>Definition:</i>	Indicator of the status of evidence for a pre-existing clinical condition of stroke.
<i>Context:</i>	Acute coronary treatment settings.
<i>Object class:</i>	Person
<i>Property:</i>	Clinical evidence status

Value domain attributes

Representational attributes

<i>Representation class:</i>	Code						
<i>Data type:</i>	Number						
<i>Format:</i>	N						
<i>Maximum character length:</i>	1						
<i>Permissible values:</i>	<table><thead><tr><th>Value</th><th>Meaning</th></tr></thead><tbody><tr><td>1</td><td>Objective evidence</td></tr><tr><td>2</td><td>No objective evidence</td></tr></tbody></table>	Value	Meaning	1	Objective evidence	2	No objective evidence
Value	Meaning						
1	Objective evidence						
2	No objective evidence						

Source and reference attributes

<i>Submitting organisation:</i>	Australian Institute of Health and Welfare
---------------------------------	--

Data element attributes

Collection and usage attributes

<i>Guide for use:</i>	For ischaemic: non-haemorrhagic cerebral infarction, objective evidence is coded where the diagnosis is supported by cerebral imaging (CT or MRI), or For haemorrhagic: intracerebral haemorrhage, objective evidence is coded where the diagnosis is supported by cerebral imaging (CT or MRI).
<i>Collection methods:</i>	For each Person – concurrent clinical condition (acute coronary syndrome), code NN, the data elements Person – clinical evidence status (chronic lung disease), code N; Person – clinical evidence status (heart failure), code N; Person – clinical evidence status (stroke), code N; Person – clinical evidence status (peripheral arterial disease), code N; Person – clinical

evidence status (sleep apnoea syndrome), code N must also be recorded.

Source and reference attributes

Submitting organisation: Acute coronary syndrome data working group
Steward: The National Heart Foundation of Australia and The Cardiac Society of Australia and New Zealand

Relational attributes

Related metadata references: Supersedes Clinical evidence status, version 1, DE, NHDD, NHIMG, Superseded 01/03/2005

Implementation in Data Set Specifications: Acute coronary syndrome (clinical) DSS NHIG, Standard 07/12/2005

Implementation start date: 07/12/2005

Information specific to this data set:

This data element seeks to ensure that patients with self-reported past symptoms pertinent to acute coronary syndrome, have objective evidence supporting reported diagnoses, using current medical practice.

Acute coronary syndrome (clinical) DSS NHIG, Superseded 07/12/2005

Information specific to this data set:

This data element seeks to ensure that patients with self-reported past symptoms pertinent to acute coronary syndrome, have objective evidence supporting reported diagnoses, using current medical practice.

Clinical procedure timing (status)

Identifying and definitional attributes

<i>Metadata item type:</i>	Data Element
<i>Technical name:</i>	Person – clinical procedure timing, code N
<i>METeOR identifier:</i>	284863
<i>Registration status:</i>	NHIG, Standard 04/06/2004
<i>Definition:</i>	The timing of the provision of a clinical procedure, as represented by a code.

Data element concept attributes

<i>Data element concept:</i>	Person – clinical procedure timing
<i>Definition:</i>	An indicator of the timing of the provision of a clinical procedure.
<i>Context:</i>	Acute coronary treatment settings.
<i>Object class:</i>	Person
<i>Property:</i>	Clinical procedure timing

Value domain attributes

Representational attributes

<i>Representation class:</i>	Code						
<i>Data type:</i>	Number						
<i>Format:</i>	N						
<i>Maximum character length:</i>	1						
<i>Permissible values:</i>	<table><thead><tr><th>Value</th><th>Meaning</th></tr></thead><tbody><tr><td>1</td><td>Procedure performed prior to an episode of admitted patient care</td></tr><tr><td>2</td><td>Procedure performed during an episode of admitted patient care</td></tr></tbody></table>	Value	Meaning	1	Procedure performed prior to an episode of admitted patient care	2	Procedure performed during an episode of admitted patient care
Value	Meaning						
1	Procedure performed prior to an episode of admitted patient care						
2	Procedure performed during an episode of admitted patient care						

Data element attributes

Collection and usage attributes

<i>Guide for use:</i>	Record only for those procedure codes that apply.
<i>Collection methods:</i>	This data element should be recorded for each type of procedure performed that is pertinent to the treatment of acute coronary syndrome.

Source and reference attributes

<i>Submitting organisation:</i>	Acute coronary syndrome data working group
<i>Steward:</i>	The National Heart Foundation of Australia and The Cardiac Society of Australia and New Zealand

Relational attributes

<i>Related metadata references:</i>	Supersedes Clinical procedure timing status, version 1, DE, NHDD, NHIMG, Superseded 01/03/2005
-------------------------------------	--

Implementation in Data Set Specifications:

Acute coronary syndrome (clinical) DSS NHIG, Standard
07/12/2005

Implementation start date: 07/12/2005

Acute coronary syndrome (clinical) DSS NHIG, Superseded
07/12/2005

Clinical urgency

Identifying and definitional attributes

<i>Metadata item type:</i>	Data Element
<i>Technical name:</i>	Elective surgery waiting list episode—clinical urgency, code N
<i>METeOR identifier:</i>	270008
<i>Registration status:</i>	NHIG, Standard 01/03/2005
<i>Definition:</i>	A clinical assessment of the urgency with which a patient requires elective hospital care, as represented by a code.

Data element concept attributes

<i>Data element concept:</i>	Elective surgery waiting list episode—clinical urgency
<i>Definition:</i>	A clinical assessment of the urgency with which a patient requires elective hospital care.
<i>Context:</i>	<p>Elective surgery:</p> <p>Categorisation of waiting list patients by clinical urgency assists hospital management and clinicians in the prioritisation of their workloads. It gives health consumers a reasonable estimate of the maximum time they should expect to wait for care. Clinical urgency classification allows a meaningful measure of system performance to be calculated, namely the number or proportion of patients who wait for times in excess of the maximum desirable time limit for their urgency category (metadata item the overdue patient status).</p>
<i>Object class:</i>	Elective surgery waiting list episode
<i>Property:</i>	Clinical urgency

Value domain attributes

Representational attributes

<i>Representation class:</i>	Code								
<i>Data type:</i>	Number								
<i>Format:</i>	N								
<i>Maximum character length:</i>	1								
<i>Permissible values:</i>	<table><thead><tr><th>Value</th><th>Meaning</th></tr></thead><tbody><tr><td>1</td><td>Admission within 30 days desirable for a condition that has the potential to deteriorate quickly to the point that it may become an emergency</td></tr><tr><td>2</td><td>Admission within 90 days desirable for a condition causing some pain, dysfunction or disability but which is not likely to deteriorate quickly or become an emergency</td></tr><tr><td>3</td><td>Admission at some time in the future acceptable for a condition causing minimal or no pain, dysfunction or disability, which is unlikely to deteriorate quickly and which does not have the potential to become an emergency</td></tr></tbody></table>	Value	Meaning	1	Admission within 30 days desirable for a condition that has the potential to deteriorate quickly to the point that it may become an emergency	2	Admission within 90 days desirable for a condition causing some pain, dysfunction or disability but which is not likely to deteriorate quickly or become an emergency	3	Admission at some time in the future acceptable for a condition causing minimal or no pain, dysfunction or disability, which is unlikely to deteriorate quickly and which does not have the potential to become an emergency
Value	Meaning								
1	Admission within 30 days desirable for a condition that has the potential to deteriorate quickly to the point that it may become an emergency								
2	Admission within 90 days desirable for a condition causing some pain, dysfunction or disability but which is not likely to deteriorate quickly or become an emergency								
3	Admission at some time in the future acceptable for a condition causing minimal or no pain, dysfunction or disability, which is unlikely to deteriorate quickly and which does not have the potential to become an emergency								

Data element attributes

Collection and usage attributes

Guide for use: The classification employs a system of urgency categorisation based on factors such as the degree of pain, dysfunction and disability caused by the condition and its potential to deteriorate quickly into an emergency. All patients ready for care must be assigned to one of the urgency categories, regardless of how long it is estimated they will need to wait for surgery.

Comments: A patient's classification may change if he or she undergoes **clinical review** during the waiting period. The need for clinical review varies with the patient's condition and is therefore at the discretion of the treating clinician. The waiting list information system should be able to record dates when the classification is changed (metadata item Elective care waiting list episode – category reassignment date, DDMMYYYY).

Source and reference attributes

Origin: National Health Data Committee

Relational attributes

Related metadata references: Supersedes Clinical urgency, version 2, DE, NHDD, NHIMG, Superseded 01/03/2005

See also Elective surgery waiting list episode – overdue patient status, code N NHIG, Standard 01/03/2005

Implementation in Data Set Specifications: Elective surgery waiting times (census data) NMDS NHIG, Standard 07/12/2005

Implementation start date: 30/09/2006

Elective surgery waiting times (census data) NMDS NHIG, Superseded 07/12/2005

Implementation start date: 30/09/2002

Implementation end date: 30/06/2006

Elective surgery waiting times (removals data) NMDS NHIG, Standard 07/12/2005

Implementation start date: 01/07/2006

Elective surgery waiting times (removals data) NMDS NHIG, Superseded 07/12/2005

Implementation start date: 01/07/2002

Implementation end date: 30/06/2006

Clopidogrel therapy status

Identifying and definitional attributes

<i>Metadata item type:</i>	Data Element
<i>Technical name:</i>	Person – clopidogrel therapy status, code NN
<i>METeOR identifier:</i>	284873
<i>Registration status:</i>	NHIG, Standard 04/06/2004
<i>Definition:</i>	The person's clopidogrel therapy status, as represented by a code.

Data element concept attributes

<i>Data element concept:</i>	Person – clopidogrel therapy status
<i>Definition:</i>	Identifies the person's clopidogrel therapy status.
<i>Context:</i>	Health care and clinical settings.
<i>Object class:</i>	Person
<i>Property:</i>	Clopidogrel therapy status

Value domain attributes

Representational attributes

<i>Representation class:</i>	Code																				
<i>Data type:</i>	Number																				
<i>Format:</i>	NN																				
<i>Maximum character length:</i>	2																				
<i>Permissible values:</i>	<table><thead><tr><th>Value</th><th>Meaning</th></tr></thead><tbody><tr><td>10</td><td>Given</td></tr><tr><td>21</td><td>Not given - therapy not indicated</td></tr><tr><td>22</td><td>Not given - patient refusal</td></tr><tr><td>23</td><td>Not given - true allergy to clopidogrel</td></tr><tr><td>24</td><td>Not given - active bleeding</td></tr><tr><td>25</td><td>Not given - bleeding risk</td></tr><tr><td>26</td><td>Not given - thrombocytopenia</td></tr><tr><td>27</td><td>Not given - severe hepatic dysfunction</td></tr><tr><td>29</td><td>Not given - other</td></tr></tbody></table>	Value	Meaning	10	Given	21	Not given - therapy not indicated	22	Not given - patient refusal	23	Not given - true allergy to clopidogrel	24	Not given - active bleeding	25	Not given - bleeding risk	26	Not given - thrombocytopenia	27	Not given - severe hepatic dysfunction	29	Not given - other
Value	Meaning																				
10	Given																				
21	Not given - therapy not indicated																				
22	Not given - patient refusal																				
23	Not given - true allergy to clopidogrel																				
24	Not given - active bleeding																				
25	Not given - bleeding risk																				
26	Not given - thrombocytopenia																				
27	Not given - severe hepatic dysfunction																				
29	Not given - other																				
<i>Supplementary values:</i>	90 Not stated/inadequately described																				

Collection and usage attributes

<i>Guide for use:</i>	CODES 21 - 29 Not given If recording 'Not given', record the principal reason if more than one code applies.
-----------------------	---

Source and reference attributes

<i>Submitting organisation:</i>	Australian Institute of Health and Welfare
---------------------------------	--

Data element attributes

Source and reference attributes

Submitting organisation: Acute coronary syndrome data working group
Steward: The National Heart Foundation of Australia and The Cardiac Society of Australia and New Zealand

Relational attributes

Related metadata references: Supersedes Clopidogrel therapy status, version 1, DE, NHDD, NHIMG, Superseded 01/03/2005

Implementation in Data Set Specifications: Acute coronary syndrome (clinical) DSS NHIG, Standard 07/12/2005

Implementation start date: 07/12/2005

Information specific to this data set:

For Acute coronary syndrome (ACS) reporting, can be collected at any time point during the management of the current event (i.e. at the time of triage, at times during the admission, or at the time of discharge).

Acute coronary syndrome (clinical) DSS NHIG, Superseded 07/12/2005

Information specific to this data set:

For Acute coronary syndrome (ACS) reporting, can be collected at any time point during the management of the current event (i.e. at the time of triage, at times during the admission, or at the time of discharge).

Co-location status of mental health service

Identifying and definitional attributes

<i>Metadata item type:</i>	Data Element
<i>Technical name:</i>	Specialised mental health service—co-location with acute care hospital, code N
<i>METeOR identifier:</i>	286995
<i>Registration status:</i>	NHIG, Standard 08/12/2004
<i>Definition:</i>	Whether a mental health service is co-located with an acute care hospital, as represented by a code.

Data element concept attributes

<i>Data element concept:</i>	Specialised mental health service—co-location with acute care hospital
<i>Definition:</i>	Whether a mental health service is co-located with an acute care hospital.
<i>Context:</i>	Admitted patient mental health services.
<i>Object class:</i>	Specialised mental health service
<i>Property:</i>	Co-location with acute care hospital

Value domain attributes

Representational attributes

<i>Representation class:</i>	Code						
<i>Data type:</i>	Number						
<i>Format:</i>	N						
<i>Maximum character length:</i>	1						
<i>Permissible values:</i>	<table><thead><tr><th>Value</th><th>Meaning</th></tr></thead><tbody><tr><td>1</td><td>Co-located</td></tr><tr><td>2</td><td>Not co-located</td></tr></tbody></table>	Value	Meaning	1	Co-located	2	Not co-located
Value	Meaning						
1	Co-located						
2	Not co-located						

Collection and usage attributes

<i>Guide for use:</i>	<p>CODE 1 Co-located</p> <p>Co-located health services are those that are established physically and organisationally as part of an acute care hospital service. There are two forms of co-location:</p> <ul style="list-style-type: none">• a health service that is built and managed as a ward or unit within an acute care hospital; or• the health service operates in a separate building but is located on, or immediately adjoining, the acute care hospital campus. <p>In the second option, units and wards within a psychiatric hospital may be classified as co-located when all the following criteria apply:</p> <ul style="list-style-type: none">• a single organisational or management structure covers the acute care hospital and the psychiatric hospital;• a single employer covers the staff of the acute care hospital and the psychiatric hospital;
-----------------------	--

- the location of the acute care hospital and psychiatric hospital can be regarded as part of a single overall hospital campus; and
- the patients of the psychiatric hospital are regarded as patients of the single integrated health service.

Source and reference attributes

Submitting organisation: Australian Institute of Health and Welfare

Data element attributes

Collection and usage attributes

Collection methods: To be reported for mental health services that primarily provide overnight admitted patient care. Excludes residential mental health services and ambulatory mental health services.

Relational attributes

Implementation in Data Set Specifications: Mental health establishments NMDS 2005-2006 NHIG, Superseded 07/12/2005

Implementation start date: 01/07/2005

Mental health establishments NMDS 2005-2006 NHIG, Superseded 21/03/2006

Implementation start date: 01/07/2005

Implementation end date: 30/06/2006

Mental health establishments NMDS 2006-2007 NHIG, Superseded 23/10/2006

Implementation start date: 01/07/2006

Implementation end date: 30/06/2007

Mental health establishments NMDS 2007-2008 NHIG, Standard 23/10/2006

Implementation start date: 01/07/2007

Compensable status

Identifying and definitional attributes

<i>Metadata item type:</i>	Data Element
<i>Technical name:</i>	Patient – compensable status, code N
<i>METeOR identifier:</i>	270100
<i>Registration status:</i>	NHIG, Standard 01/03/2005
<i>Definition:</i>	Whether or not a patient is a compensable patient , as represented by a code.

Data element concept attributes

<i>Data element concept:</i>	Patient – compensable status
<i>Definition:</i>	A compensable patient is an individual who is entitled to receive or has received a compensation payment with respect to an injury or disease.
<i>Object class:</i>	Patient
<i>Property:</i>	Compensable status

Value domain attributes

Representational attributes

<i>Representation class:</i>	Code								
<i>Data type:</i>	Number								
<i>Format:</i>	N								
<i>Maximum character length:</i>	1								
<i>Permissible values:</i>	<table><thead><tr><th>Value</th><th>Meaning</th></tr></thead><tbody><tr><td>1</td><td>Compensable</td></tr><tr><td>2</td><td>Non-compensable</td></tr><tr><td>9</td><td>Not stated/not known</td></tr></tbody></table>	Value	Meaning	1	Compensable	2	Non-compensable	9	Not stated/not known
Value	Meaning								
1	Compensable								
2	Non-compensable								
9	Not stated/not known								
<i>Supplementary values:</i>									

Data element attributes

Collection and usage attributes

<i>Guide for use:</i>	This definition of compensable patient excludes eligible beneficiaries (Department of Veterans' Affairs), Defence Force personnel and persons covered by the Motor Accident Compensation Scheme, Northern Territory.
<i>Comments:</i>	To assist in the analyses of utilisation and health care funding.

Source and reference attributes

<i>Origin:</i>	National Health Data Committee
----------------	--------------------------------

Relational attributes

<i>Related metadata references:</i>	Supersedes Compensable status, version 3, DE, NHDD, NHIMG, Superseded 01/03/2005
<i>Implementation in Data Set Specifications:</i>	Non-admitted patient emergency department care NMDS NHIG, Standard 24/03/2006

Implementation start date: 01/07/2006

Non-admitted patient emergency department care NMDS
NHIG, Superseded 07/12/2005

Non-admitted patient emergency department care NMDS
NHIG, Superseded 24/03/2006

Implementation start date: 01/07/2005

Implementation end date: 30/06/2006

Complication of labour and delivery

Identifying and definitional attributes

<i>Metadata item type:</i>	Data Element
<i>Technical name:</i>	Birth event – complication, code (ICD-10-AM 5th edn) ANN{.N[N]}
<i>METeOR identifier:</i>	333818
<i>Registration status:</i>	NHIG, Standard 07/12/2005
<i>Definition:</i>	Medical and obstetric complications (necessitating intervention) arising after the onset of labour and before the completed delivery of the baby and placenta, as represented by a code.

Data element concept attributes

<i>Data element concept:</i>	Birth event – complication
<i>Definition:</i>	Medical and obstetric complications (necessitating intervention) arising after the onset of labour and before the completed delivery of the baby and placenta.
<i>Context:</i>	Perinatal statistics
<i>Object class:</i>	Birth event
<i>Property:</i>	Complication

Value domain attributes

Representational attributes

<i>Classification scheme:</i>	International Statistical Classification of Diseases and Related Health Problems, Tenth Revision, Australian Modification 5th edition
<i>Representation class:</i>	Code
<i>Data type:</i>	String
<i>Format:</i>	ANN{.N[N]}
<i>Maximum character length:</i>	6

Collection and usage attributes

<i>Guide for use:</i>	Complications and conditions should be coded within the Pregnancy, Childbirth, Puerperium chapter 15 of Volume 1, ICD-10-AM.
-----------------------	--

Data element attributes

Collection and usage attributes

<i>Guide for use:</i>	There is no arbitrary limit on the number of conditions specified.
<i>Comments:</i>	Complications of labour and delivery may cause maternal morbidity and may affect the health status of the baby at birth.

Source and reference attributes

<i>Submitting organisation:</i>	National Perinatal Data Development Committee
---------------------------------	---

Relational attributes

Related metadata references:

Supersedes Birth event – complication, code (ICD-10-AM 4th edn) ANN{.N[N]} NHIG, Superseded 07/12/2005

Complications of pregnancy

Identifying and definitional attributes

<i>Metadata item type:</i>	Data Element
<i>Technical name:</i>	Pregnancy (current) – complication, code (ICD-10-AM 5th edn) ANN{.N[N]}
<i>METeOR identifier:</i>	333938
<i>Registration status:</i>	NHIG, Standard 07/12/2005
<i>Definition:</i>	Complications arising up to the period immediately preceding delivery that are directly attributable to the pregnancy and may have significantly affected care during the current pregnancy and/or pregnancy outcome, as represented by a code

Data element concept attributes

<i>Data element concept:</i>	Pregnancy (current) – complication
<i>Definition:</i>	Complications arising up to the period immediately preceding delivery that are directly attributable to the pregnancy and may have significantly affected care during the current pregnancy and/or pregnancy outcome.
<i>Context:</i>	Perinatal statistics
<i>Object class:</i>	Pregnancy
<i>Property:</i>	Complication

Value domain attributes

Representational attributes

<i>Classification scheme:</i>	International Statistical Classification of Diseases and Related Health Problems, Tenth Revision, Australian Modification 5th edition
<i>Representation class:</i>	Code
<i>Data type:</i>	String
<i>Format:</i>	ANN{.N[N]}
<i>Maximum character length:</i>	6

Collection and usage attributes

<i>Guide for use:</i>	Complications and conditions should be coded within the Pregnancy, Childbirth, Puerperium chapter 15 of Volume 1, ICD-10-AM.
-----------------------	--

Data element attributes

Collection and usage attributes

<i>Guide for use:</i>	Examples of these conditions include threatened abortion, antepartum haemorrhage, pregnancy-induced hypertension and gestational diabetes. There is no arbitrary limit on the number of complications specified.
<i>Comments:</i>	Complications often influence the course and outcome of pregnancy, possibly resulting in hospital admissions and/or adverse effects on the fetus and perinatal morbidity.

Source and reference attributes

Submitting organisation: National Perinatal Data Development Committee

Relational attributes

Related metadata references: Supersedes Pregnancy (current) – complication, code (ICD-10-AM 4th edn) ANN{.N[N]} NHIG, Superseded 07/12/2005

Concurrent clinical condition (on presentation)

Identifying and definitional attributes

<i>Metadata item type:</i>	Data Element
<i>Technical name:</i>	Person – acute coronary syndrome concurrent clinical condition, code NN
<i>METeOR identifier:</i>	284891
<i>Registration status:</i>	NHIG, Standard 04/06/2004
<i>Definition:</i>	The concurrent medical conditions, which are pertinent to the risk stratification and treatment of acute coronary syndrome that a person has or has undergone prior to presentation, as represented by a code.

Data element concept attributes

<i>Data element concept:</i>	Person – acute coronary syndrome concurrent clinical condition
<i>Definition:</i>	The concurrent medical conditions, which are pertinent to the risk stratification and treatment of acute coronary syndrome that a person has or has undergone prior to presentation.
<i>Context:</i>	Acute coronary syndrome clinical reporting only.
<i>Object class:</i>	Person
<i>Property:</i>	Acute coronary syndrome concurrent clinical condition

Value domain attributes

Representational attributes

<i>Representation class:</i>	Code																										
<i>Data type:</i>	Number																										
<i>Format:</i>	NN																										
<i>Maximum character length:</i>	2																										
<i>Permissible values:</i>	<table><thead><tr><th>Value</th><th>Meaning</th></tr></thead><tbody><tr><td>11</td><td>Angina for more than last two weeks</td></tr><tr><td>12</td><td>Angina only in the last two weeks</td></tr><tr><td>21</td><td>Chronic lung disease</td></tr><tr><td>31</td><td>Heart failure</td></tr><tr><td>41</td><td>Hypertension</td></tr><tr><td>51</td><td>Ischaemic: non-haemorrhagic cerebral infarction</td></tr><tr><td>52</td><td>Haemorrhagic: intracerebral haemorrhage</td></tr><tr><td>61</td><td>Peripheral artery disease</td></tr><tr><td>62</td><td>Aortic aneurysm</td></tr><tr><td>63</td><td>Renal artery stenosis</td></tr><tr><td>71</td><td>Sleep apnoea</td></tr><tr><td>99</td><td>not stated/inadequately described</td></tr></tbody></table>	Value	Meaning	11	Angina for more than last two weeks	12	Angina only in the last two weeks	21	Chronic lung disease	31	Heart failure	41	Hypertension	51	Ischaemic: non-haemorrhagic cerebral infarction	52	Haemorrhagic: intracerebral haemorrhage	61	Peripheral artery disease	62	Aortic aneurysm	63	Renal artery stenosis	71	Sleep apnoea	99	not stated/inadequately described
Value	Meaning																										
11	Angina for more than last two weeks																										
12	Angina only in the last two weeks																										
21	Chronic lung disease																										
31	Heart failure																										
41	Hypertension																										
51	Ischaemic: non-haemorrhagic cerebral infarction																										
52	Haemorrhagic: intracerebral haemorrhage																										
61	Peripheral artery disease																										
62	Aortic aneurysm																										
63	Renal artery stenosis																										
71	Sleep apnoea																										
99	not stated/inadequately described																										
<i>Supplementary values:</i>																											

Collection and usage attributes

<i>Guide for use:</i>	Angina:
-----------------------	---------

CODE 11 Angina for more than last two weeks

This code is used where there are symptoms, which can be described as chest pain or pressure, jaw pain, arm pain, or other equivalent discomfort suggestive of cardiac ischaemia, for more than the last two weeks.

CODE 12 Angina only in the last two weeks

This code is used where there are symptoms, which can be described as chest pain or pressure, jaw pain, arm pain, or other equivalent discomfort suggestive of cardiac ischaemia, only in the last two weeks.

Chronic lung disease:

CODE 21 Chronic lung disease

This code is used where there is a history or symptoms suggestive of chronic lung disease.

Heart failure:

CODE 31 Heart failure

This code is used where a patient has past or current symptoms of heart failure (typically breathlessness or fatigue), either at rest or during exercise and/or signs of pulmonary or peripheral congestion suggestive of cardiac dysfunction.

Hypertension:

CODE 41 Hypertension

This code is used where there is current use of pharmacotherapy for hypertension and/or clinical evidence of high blood pressure.

Stroke:

CODE 51 Ischaemic: non-haemorrhagic cerebral infarction

This code is used if there is history of stroke or cerebrovascular accident (CVA) resulting from an ischaemic event where the patient suffered a loss of neurological function with residual symptoms remaining for at least 24 hours.

CODE 52 Haemorrhagic: intracerebral haemorrhage

This code is used if there is history of stroke or cerebrovascular accident (CVA) resulting from a haemorrhagic event where the patient suffered a loss of neurological function with residual symptoms remaining for at least 24 hours.

Peripheral arterial disease:

CODE 61 Peripheral artery disease

This code is used where there is history of either chronic or acute occlusion or narrowing of the arterial lumen in the aorta or extremities.

CODE 62 Aortic aneurysm

This code is used where there is a history of aneurysmal dilatation of the aorta (thoracic and or abdominal).

CODE 63 Renal artery stenosis

This code is used where there is history of functional stenosis of one or both renal arteries.

Sleep Apnoea syndrome:

CODE 71 Sleep apnoea

This code is used where there is evidence of sleep apnoea syndrome (SAS) on history.

Source and reference attributes

Data element attributes

Collection and usage attributes

Guide for use:

More than one medical condition may be recorded.

Record only those codes that apply.

Record all codes that apply.

Codes 21, 31, 51, 52, 61, 62, 63 and 71 must be accompanied by a Clinical evidence status code.

Source and reference attributes

Submitting organisation:

Acute coronary syndrome data working group

Steward:

The National Heart Foundation of Australia and The Cardiac Society of Australia and New Zealand

Relational attributes

Related metadata references:

Supersedes Concurrent clinical condition - on presentation, version 1, DE, NHDD, NHIMG, Superseded 01/03/2005

Implementation in Data Set Specifications:

Acute coronary syndrome (clinical) DSS NHIG, Standard 07/12/2005

Implementation start date: 07/12/2005

Acute coronary syndrome (clinical) DSS NHIG, Superseded 07/12/2005

Congenital malformations

Identifying and definitional attributes

<i>Metadata item type:</i>	Data Element
<i>Technical name:</i>	Person – congenital malformation, code (ICD-10-AM 5th edn) ANN{.N[N]}
<i>METeOR identifier:</i>	333934
<i>Registration status:</i>	NHIG, Standard 07/12/2005
<i>Definition:</i>	Structural abnormalities (including deformations) that are present at birth and diagnosed prior to separation from care, as represented by an ICD-10-AM code.
<i>Context:</i>	Admitted patient care

Data element concept attributes

<i>Data element concept:</i>	Person – congenital malformation
<i>Definition:</i>	Structural abnormalities (including deformations) that are present at birth and diagnosed prior to separation from care.
<i>Object class:</i>	Person
<i>Property:</i>	Congenital malformation

Value domain attributes

Representational attributes

<i>Classification scheme:</i>	International Statistical Classification of Diseases and Related Health Problems, Tenth Revision, Australian Modification 5th edition
<i>Representation class:</i>	Code
<i>Data type:</i>	String
<i>Format:</i>	ANN{.N[N]}
<i>Maximum character length:</i>	6

Source and reference attributes

<i>Origin:</i>	International Classification of Diseases - 10th Revision, Australian Modification (5th Edition 2004) National Centre for Classification in Health, Sydney.
----------------	--

Data element attributes

Collection and usage attributes

<i>Guide for use:</i>	Coding to the disease classification of ICD-10-AM is the preferred method of coding admitted patients. However, for the perinatal data collection, the use of BPA is preferred as this is more detailed (see the metadata item Person – congenital malformations, code (BPA 1979) ANN.N[N]).
<i>Comments:</i>	Required to monitor trends in the reported incidence of congenital malformations, to detect new drug and environmental teratogens, to analyse possible causes in epidemiological studies, and to determine survival rates and

the utilisation of paediatric services.

Source and reference attributes

Submitting organisation: National Perinatal Data Development Committee

Relational attributes

Related metadata references: Supersedes Person—congenital malformation, code (ICD-10-AM 4th edn) ANN{.N[N]} NHIG, Superseded 07/12/2005

Congenital malformations—BPA code

Identifying and definitional attributes

<i>Metadata item type:</i>	Data Element
<i>Technical name:</i>	Person—congenital malformation, code (BPA 1979) ANN.N[N]
<i>METeOR identifier:</i>	270408
<i>Registration status:</i>	NHIG, Standard 01/03/2005
<i>Definition:</i>	Structural abnormalities (including deformations) that are present at birth and diagnosed prior to separation from care, as represented by a BPA code.
<i>Context:</i>	Perinatal statistics

Data element concept attributes

<i>Data element concept:</i>	Person—congenital malformation
<i>Definition:</i>	Structural abnormalities (including deformations) that are present at birth and diagnosed prior to separation from care.
<i>Object class:</i>	Person
<i>Property:</i>	Congenital malformation

Value domain attributes

Representational attributes

<i>Classification scheme:</i>	British Paediatric Association Classification of Diseases 1979
<i>Representation class:</i>	Code
<i>Data type:</i>	String
<i>Format:</i>	ANN.N[N]
<i>Maximum character length:</i>	5

Data element attributes

Collection and usage attributes

<i>Guide for use:</i>	Coding to the disease classification of ICD-10-AM is the preferred method of coding admitted patients. For perinatal data collections, the use of British Paediatric Association (BPA) Classification of Diseases is preferred as this is more detailed.
<i>Comments:</i>	There is no arbitrary limit on the number of conditions specified. Most perinatal data groups and birth defects registers in the states and territories have used the 5-digit BPA Classification of Diseases to code congenital malformations since the early 1980s.

Source and reference attributes

<i>Submitting organisation:</i>	National Perinatal Data Development Committee
<i>Origin:</i>	British Paediatric Association Classification of Diseases (1979)

Relational attributes

<i>Related metadata references:</i>	Supersedes Congenital malformations - BPA code, version 1, DE, NHDD, NHIMG, Superseded 01/03/2005
-------------------------------------	---

Consumer committee representation arrangements

Identifying and definitional attributes

<i>Metadata item type:</i>	Data Element
<i>Technical name:</i>	Specialised mental health service organisation – consumer committee representation arrangements, code N
<i>METeOR identifier:</i>	288855
<i>Registration status:</i>	NHIG, Standard 08/12/2004
<i>Definition:</i>	Extent to which a specialised mental health service organisation has formal committee mechanisms in place to promote the participation of mental health consumers in the planning, delivery and evaluation of the service, as represented by a code.

Data element concept attributes

<i>Data element concept:</i>	Specialised mental health service organisation – consumer committee representation arrangements
<i>Definition:</i>	Extent to which a specialised mental health service organisation has formal committee mechanisms in place to promote the participation of mental health consumers in the planning, delivery and evaluation of the service.
<i>Context:</i>	Specialised mental health services.
<i>Object class:</i>	Specialised mental health service organisation
<i>Property:</i>	Consumer committee representation arrangements

Collection and usage attributes

<i>Guide for use:</i>	<p>A consumer is a person who is currently utilising, or has previously utilised, a mental health service.</p> <p>Mental health service consumers include persons receiving care for their own, or another persons', mental illness or psychiatric disability.</p> <p>To be regarded as having a formal position on a management or advisory committee the consumer representative needs to be a voting member.</p>
-----------------------	---

Value domain attributes

Representational attributes

<i>Representation class:</i>	Code						
<i>Data type:</i>	Number						
<i>Format:</i>	N						
<i>Maximum character length:</i>	1						
<i>Permissible values:</i>	<table><thead><tr><th>Value</th><th>Meaning</th></tr></thead><tbody><tr><td>1</td><td>Formal position(s) for consumers exist on the organisation's management committee for the appointment of person(s) to represent the interests of consumers</td></tr><tr><td>2</td><td>Specific consumer advisory committee(s) exists to advise on all relevant mental health services</td></tr></tbody></table>	Value	Meaning	1	Formal position(s) for consumers exist on the organisation's management committee for the appointment of person(s) to represent the interests of consumers	2	Specific consumer advisory committee(s) exists to advise on all relevant mental health services
Value	Meaning						
1	Formal position(s) for consumers exist on the organisation's management committee for the appointment of person(s) to represent the interests of consumers						
2	Specific consumer advisory committee(s) exists to advise on all relevant mental health services						

- managed by the organisation
- 3 Specific consumer advisory committee(s) exists to advise on some but not all relevant mental health services managed by the organisation
 - 4 Consumers participate on a broadly based advisory committee which include a mixture of organisations and groups representing a wide range of interests
 - 5 Consumers are not represented on any advisory committee but are encouraged to meet with senior representatives of the organisation as required
 - 6 No specific arrangements exist for consumer participation in planning and evaluation of services

Collection and usage attributes

Guide for use:

Select the option above that best describes the type of formal committee mechanisms within your organisation for ensuring participation by mental health consumers in the planning and evaluation of services.

Data element attributes

Collection and usage attributes

Guide for use:

Select the option above that best describes the type of formal committee mechanisms with in your organisation for ensuring participation by mental health consumers in the planning and evaluation of services.

Relational attributes

Implementation in Data Set Specifications:

Mental health establishments NMDS 2005-2006 NHIG,
Superseded 07/12/2005

Implementation start date: 01/07/2005

Information specific to this data set:

Obligation condition: reporting of this data element is optional for non-government residential mental health services and specialised mental health services provided by private hospitals that receive state or territory government funding.

Mental health establishments NMDS 2005-2006 NHIG,
Superseded 21/03/2006

Implementation start date: 01/07/2005

Implementation end date: 30/06/2006

Information specific to this data set:

Obligation condition: reporting of this data element is optional for non-government residential mental health services and specialised mental health services provided by private hospitals that receive state or territory government funding.

Mental health establishments NMDS 2006-2007 NHIG,
Superseded 23/10/2006

Implementation start date: 01/07/2006

Implementation end date: 30/06/2007

Information specific to this data set:

Obligation condition: reporting of this data element is optional for non-government residential mental health services and specialised mental health services provided by private hospitals that receive state or territory government funding.

Mental health establishments NMDS 2007-2008 NHIG,
Standard 23/10/2006

Implementation start date: 01/07/2007

Information specific to this data set:

Obligation condition: reporting of this data element is optional for non-government residential mental health services and specialised mental health services provided by private hospitals that receive state or territory government funding.

Consumer participation arrangements—consumer consultants employed

Identifying and definitional attributes

<i>Metadata item type:</i>	Data Element
<i>Technical name:</i>	Specialised mental health service organisation – consumer participation arrangements (consumer consultants employed), code N
<i>METeOR identifier:</i>	288866
<i>Registration status:</i>	NHIG, Standard 08/12/2004
<i>Definition:</i>	Whether the service employs consumer consultants on a paid basis to represent the interests of consumers and advocate for their needs, in order to promote the participation of mental health consumers in the planning, delivery and evaluation of the service, as represented by a code.

Data element concept attributes

<i>Data element concept:</i>	Specialised mental health service organisation – consumer participation arrangements (consumer consultants employed)
<i>Definition:</i>	Whether the service employs consumer consultants on a paid basis to represent the interests of consumers and advocate for their needs, in order to promote the participation of mental health consumers in the planning, delivery and evaluation of the service.
<i>Context:</i>	Specialised mental health services.
<i>Object class:</i>	Specialised mental health service organisation
<i>Property:</i>	Consumer participation arrangements

Collection and usage attributes

<i>Guide for use:</i>	<p>A consumer is a person who is currently utilising, or has previously utilised, a mental health service.</p> <p>Mental health service consumers include persons receiving care for their own, or another persons', mental illness or psychiatric disability.</p> <p>The term mental health consumer consultant refers to a person who is employed (or engaged via contract) on a part-time or full-time basis to act as a consumer consultant. This implies the person received a salary or contract fee on a regular basis. It does not refer to arrangements where the consumer only received reimbursements of expenses or occasional sitting fees for attendance at meetings.</p>
-----------------------	---

Value domain attributes

Representational attributes

<i>Representation class:</i>	Code
<i>Data type:</i>	Boolean
<i>Format:</i>	N
<i>Maximum character length:</i>	1

<i>Permissible values:</i>	Value	Meaning
	1	Yes
	2	No
<i>Supplementary values:</i>	9	Don't know

Data element attributes

Relational attributes

<i>Related metadata references:</i>	See also Specialised mental health service organisation – consumer participation arrangements (consumer satisfaction surveys), code N NHIG, Standard 08/12/2004
	See also Specialised mental health service organisation – consumer participation arrangements (formal complaints mechanism), code N NHIG, Standard 08/12/2004
	See also Specialised mental health service organisation – consumer participation arrangements (formal participation policy), code N NHIG, Standard 08/12/2004
	See also Specialised mental health service organisation – consumer participation arrangements (regular discussion groups), code N NHIG, Standard 08/12/2004
<i>Implementation in Data Set Specifications:</i>	Mental health establishments NMDS 2005-2006 NHIG, Superseded 07/12/2005
	<i>Implementation start date:</i> 01/07/2005
	<i>Information specific to this data set:</i>
	Obligation condition: reporting of this data element is optional for non-government residential mental health services and specialised mental health services provided by private hospitals that receive state or territory government funding.
	Mental health establishments NMDS 2005-2006 NHIG, Superseded 21/03/2006
	<i>Implementation start date:</i> 01/07/2005
	<i>Implementation end date:</i> 30/06/2006
	<i>Information specific to this data set:</i>
	Obligation condition: reporting of this data element is optional for non-government residential mental health services and specialised mental health services provided by private hospitals that receive state or territory government funding.
	Mental health establishments NMDS 2006-2007 NHIG, Superseded 23/10/2006
	<i>Implementation start date:</i> 01/07/2006
	<i>Implementation end date:</i> 30/06/2007
	<i>Information specific to this data set:</i>
	Obligation condition: reporting of this data element is optional for non-government residential mental health services and specialised mental health services provided by private hospitals that receive state or territory government funding.
	Mental health establishments NMDS 2007-2008 NHIG,

Standard 23/10/2006

Implementation start date: 01/07/2007

Information specific to this data set:

Obligation condition: reporting of this data element is optional for non-government residential mental health services and specialised mental health services provided by private hospitals that receive state or territory government funding.

Consumer participation arrangements—consumer satisfaction surveys

Identifying and definitional attributes

<i>Metadata item type:</i>	Data Element
<i>Technical name:</i>	Specialised mental health service organisation – consumer participation arrangements (consumer satisfaction surveys), code N
<i>METeOR identifier:</i>	290418
<i>Registration status:</i>	NHIG, Standard 08/12/2004
<i>Definition:</i>	Whether the service conducts consumer satisfaction surveys, in order to promote the participation of mental health consumers in the planning, delivery and evaluation of the service, as represented by a code.

Data element concept attributes

<i>Data element concept:</i>	Specialised mental health service organisation – consumer participation arrangements (consumer satisfaction surveys)
<i>Definition:</i>	Whether the service conducts consumer satisfaction surveys, in order to promote the participation of mental health consumers in the planning, delivery and evaluation of the service.
<i>Context:</i>	Specialised mental health services.
<i>Object class:</i>	Specialised mental health service organisation
<i>Property:</i>	Consumer participation arrangements

Collection and usage attributes

<i>Guide for use:</i>	A consumer is a person who is currently utilising, or has previously utilised, a mental health service. Mental health service consumers include persons receiving care for their own, or another persons', mental illness or psychiatric disability.
-----------------------	---

Value domain attributes

Representational attributes

<i>Representation class:</i>	Code								
<i>Data type:</i>	Boolean								
<i>Format:</i>	N								
<i>Maximum character length:</i>	1								
<i>Permissible values:</i>	<table><thead><tr><th>Value</th><th>Meaning</th></tr></thead><tbody><tr><td>1</td><td>Yes</td></tr><tr><td>2</td><td>No</td></tr><tr><td>9</td><td>Don't know</td></tr></tbody></table>	Value	Meaning	1	Yes	2	No	9	Don't know
Value	Meaning								
1	Yes								
2	No								
9	Don't know								
<i>Supplementary values:</i>									

Data element attributes

Relational attributes

Related metadata references:

See also Specialised mental health service organisation – consumer participation arrangements (consumer consultants employed), code N NHIG, Standard 08/12/2004

See also Specialised mental health service organisation – consumer participation arrangements (formal complaints mechanism), code N NHIG, Standard 08/12/2004

See also Specialised mental health service organisation – consumer participation arrangements (formal participation policy), code N NHIG, Standard 08/12/2004

See also Specialised mental health service organisation – consumer participation arrangements (regular discussion groups), code N NHIG, Standard 08/12/2004

Implementation in Data Set Specifications:

Mental health establishments NMDS 2005-2006 NHIG, Superseded 07/12/2005

Implementation start date: 01/07/2005

Information specific to this data set:

Obligation condition: reporting of this data element is optional for non-government residential mental health services and specialised mental health services provided by private hospitals that receive state or territory government funding.

Mental health establishments NMDS 2005-2006 NHIG, Superseded 21/03/2006

Implementation start date: 01/07/2005

Implementation end date: 30/06/2006

Information specific to this data set:

Obligation condition: reporting of this data element is optional for non-government residential mental health services and specialised mental health services provided by private hospitals that receive state or territory government funding.

Mental health establishments NMDS 2006-2007 NHIG, Superseded 23/10/2006

Implementation start date: 01/07/2006

Implementation end date: 30/06/2007

Information specific to this data set:

Obligation condition: reporting of this data element is optional for non-government residential mental health services and specialised mental health services provided by private hospitals that receive state or territory government funding.

Mental health establishments NMDS 2007-2008 NHIG, Standard 23/10/2006

Implementation start date: 01/07/2007

Information specific to this data set:

Obligation condition: reporting of this data element is optional for non-government residential mental health services and specialised mental health services provided by private hospitals that receive state or territory government funding.

Consumer participation arrangements—formal complaints mechanism

Identifying and definitional attributes

<i>Metadata item type:</i>	Data Element
<i>Technical name:</i>	Specialised mental health service organisation – consumer participation arrangements (formal complaints mechanism), code N
<i>METeOR identifier:</i>	290415
<i>Registration status:</i>	NHIG, Standard 08/12/2004
<i>Definition:</i>	Whether the service has developed a formal internal complaints mechanism in which complaints can be made by consumers and are regularly reviewed by a committee that includes consumers, in order to promote the participation of mental health consumers in the planning, delivery and evaluation of the service, as represented by a code.

Data element concept attributes

<i>Data element concept:</i>	Specialised mental health service organisation – consumer participation arrangements (formal internal complaints mechanism)
<i>Definition:</i>	Whether the service has developed a formal internal complaints mechanism in which complaints can be made by consumers and are regularly reviewed by a committee that includes consumers, in order to promote the participation of mental health consumers in the planning, delivery and evaluation of the service.
<i>Context:</i>	Specialised mental health services.
<i>Object class:</i>	Specialised mental health service organisation
<i>Property:</i>	Consumer participation arrangements

Collection and usage attributes

<i>Guide for use:</i>	A consumer is a person who is currently utilising, or has previously utilised, a mental health service. Mental health service consumers include persons receiving care for their own, or another persons', mental illness or psychiatric disability.
-----------------------	---

Value domain attributes

Representational attributes

<i>Representation class:</i>	Code						
<i>Data type:</i>	Boolean						
<i>Format:</i>	N						
<i>Maximum character length:</i>	1						
<i>Permissible values:</i>	<table><thead><tr><th>Value</th><th>Meaning</th></tr></thead><tbody><tr><td>1</td><td>Yes</td></tr><tr><td>2</td><td>No</td></tr></tbody></table>	Value	Meaning	1	Yes	2	No
Value	Meaning						
1	Yes						
2	No						

Data element attributes

Relational attributes

Related metadata references:

See also Specialised mental health service organisation – consumer participation arrangements (regular discussion groups), code N NHIG, Standard 08/12/2004

See also Specialised mental health service organisation – consumer participation arrangements (formal participation policy), code N NHIG, Standard 08/12/2004

See also Specialised mental health service organisation – consumer participation arrangements (consumer satisfaction surveys), code N NHIG, Standard 08/12/2004

See also Specialised mental health service organisation – consumer participation arrangements (consumer consultants employed), code N NHIG, Standard 08/12/2004

Implementation in Data Set Specifications:

Mental health establishments NMDS 2005-2006 NHIG, Superseded 07/12/2005

Implementation start date: 01/07/2005

Information specific to this data set:

Obligation condition: reporting of this data element is optional for non-government residential mental health services and specialised mental health services provided by private hospitals that receive state or territory government funding.

Mental health establishments NMDS 2005-2006 NHIG, Superseded 21/03/2006

Implementation start date: 01/07/2005

Implementation end date: 30/06/2006

Information specific to this data set:

Obligation condition: reporting of this data element is optional for non-government residential mental health services and specialised mental health services provided by private hospitals that receive state or territory government funding.

Mental health establishments NMDS 2006-2007 NHIG, Superseded 23/10/2006

Implementation start date: 01/07/2006

Implementation end date: 30/06/2007

Information specific to this data set:

Obligation condition: reporting of this data element is optional for non-government residential mental health services and specialised mental health services provided by private hospitals that receive state or territory government funding.

Mental health establishments NMDS 2007-2008 NHIG, Standard 23/10/2006

Implementation start date: 01/07/2007

Information specific to this data set:

Obligation condition: reporting of this data element is optional for non-government residential mental health services and specialised mental health services provided by private hospitals that receive state or territory government funding.

Consumer participation arrangements—formal participation policy

Identifying and definitional attributes

<i>Metadata item type:</i>	Data Element
<i>Technical name:</i>	Specialised mental health service organisation – consumer participation arrangements (formal participation policy), code N
<i>METeOR identifier:</i>	290410
<i>Registration status:</i>	NHIG, Standard 08/12/2004
<i>Definition:</i>	Whether the service has developed a formal and documented policy on participation by consumers, in order to promote the participation of mental health consumers in the planning, delivery and evaluation of the service, as represented by a code.

Data element concept attributes

<i>Data element concept:</i>	Specialised mental health service organisation – consumer participation arrangements (formal participation policy)
<i>Definition:</i>	Whether the service has developed a formal and documented policy on participation by consumers, in order to promote the participation of mental health consumers in the planning, delivery and evaluation of the service.
<i>Context:</i>	Specialised mental health services.
<i>Object class:</i>	Specialised mental health service organisation
<i>Property:</i>	Consumer participation arrangements

Collection and usage attributes

<i>Guide for use:</i>	A consumer is a person who is currently utilising, or has previously utilised, a mental health service. Mental health service consumers include persons receiving care for their own, or another persons', mental illness or psychiatric disability.
-----------------------	---

Value domain attributes

Representational attributes

<i>Representation class:</i>	Code								
<i>Data type:</i>	Boolean								
<i>Format:</i>	N								
<i>Maximum character length:</i>	1								
<i>Permissible values:</i>	<table><thead><tr><th>Value</th><th>Meaning</th></tr></thead><tbody><tr><td>1</td><td>Yes</td></tr><tr><td>2</td><td>No</td></tr><tr><td>9</td><td>Don't know</td></tr></tbody></table>	Value	Meaning	1	Yes	2	No	9	Don't know
Value	Meaning								
1	Yes								
2	No								
9	Don't know								
<i>Supplementary values:</i>									

Data element attributes

Relational attributes

Related metadata references:

See also Specialised mental health service organisation – consumer participation arrangements (consumer consultants employed), code N NHIG, Standard 08/12/2004

See also Specialised mental health service organisation – consumer participation arrangements (consumer satisfaction surveys), code N NHIG, Standard 08/12/2004

See also Specialised mental health service organisation – consumer participation arrangements (formal complaints mechanism), code N NHIG, Standard 08/12/2004

See also Specialised mental health service organisation – consumer participation arrangements (regular discussion groups), code N NHIG, Standard 08/12/2004

Implementation in Data Set Specifications:

Mental health establishments NMDS 2005-2006 NHIG, Superseded 07/12/2005

Implementation start date: 01/07/2005

Information specific to this data set:

Obligation condition: reporting of this data element is optional for non-government residential mental health services and specialised mental health services provided by private hospitals that receive state or territory government funding.

Mental health establishments NMDS 2005-2006 NHIG, Superseded 21/03/2006

Implementation start date: 01/07/2005

Implementation end date: 30/06/2006

Information specific to this data set:

Obligation condition: reporting of this data element is optional for non-government residential mental health services and specialised mental health services provided by private hospitals that receive state or territory government funding.

Mental health establishments NMDS 2006-2007 NHIG, Superseded 23/10/2006

Implementation start date: 01/07/2006

Implementation end date: 30/06/2007

Information specific to this data set:

Obligation condition: reporting of this data element is optional for non-government residential mental health services and specialised mental health services provided by private hospitals that receive state or territory government funding.

Mental health establishments NMDS 2007-2008 NHIG, Standard 23/10/2006

Implementation start date: 01/07/2007

Information specific to this data set:

Obligation condition: reporting of this data element is optional for non-government residential mental health services and specialised mental health services provided by private hospitals that receive state or territory government funding.

Consumer participation arrangements—regular discussion groups

Identifying and definitional attributes

<i>Metadata item type:</i>	Data Element
<i>Technical name:</i>	Specialised mental health service organisation – consumer participation arrangements (regular discussion groups), code N
<i>METeOR identifier:</i>	290408
<i>Registration status:</i>	NHIG, Standard 08/12/2004

Data element concept attributes

<i>Data element concept:</i>	Specialised mental health service organisation – consumer participation arrangements (regular discussion groups)
<i>Definition:</i>	Whether the service holds regular discussion groups to seek the views of consumers about the service in order to promote the participation of mental health consumers in the planning, delivery and evaluation of the service.
<i>Context:</i>	Specialised mental health services.
<i>Object class:</i>	Specialised mental health service organisation
<i>Property:</i>	Consumer participation arrangements

Collection and usage attributes

<i>Guide for use:</i>	A consumer is a person who is currently utilising, or has previously utilised, a mental health service. Mental health service consumers include persons receiving care for their own, or another persons', mental illness or psychiatric disability.
-----------------------	---

Value domain attributes

Representational attributes

<i>Representation class:</i>	Code						
<i>Data type:</i>	Boolean						
<i>Format:</i>	N						
<i>Maximum character length:</i>	1						
<i>Permissible values:</i>	<table><thead><tr><th>Value</th><th>Meaning</th></tr></thead><tbody><tr><td>1</td><td>Yes</td></tr><tr><td>2</td><td>No</td></tr></tbody></table>	Value	Meaning	1	Yes	2	No
Value	Meaning						
1	Yes						
2	No						
<i>Supplementary values:</i>	<table><tbody><tr><td>9</td><td>Don't know</td></tr></tbody></table>	9	Don't know				
9	Don't know						

Data element attributes

Relational attributes

<i>Related metadata references:</i>	See also Specialised mental health service organisation – consumer participation arrangements (formal participation policy), code N NHIG, Standard 08/12/2004 See also Specialised mental health service organisation – consumer participation arrangements (formal complaints)
-------------------------------------	--

Implementation in Data Set Specifications:

mechanism), code N NHIG, Standard 08/12/2004

See also Specialised mental health service organisation – consumer participation arrangements (consumer satisfaction surveys), code N NHIG, Standard 08/12/2004

See also Specialised mental health service organisation – consumer participation arrangements (consumer consultants employed), code N NHIG, Standard 08/12/2004

Mental health establishments NMDS 2005-2006 NHIG, Superseded 07/12/2005

Implementation start date: 01/07/2005

Information specific to this data set:

Obligation condition: reporting of this data element is optional for non-government residential mental health services and specialised mental health services provided by private hospitals that receive state or territory government funding.

Mental health establishments NMDS 2005-2006 NHIG, Superseded 21/03/2006

Implementation start date: 01/07/2005

Implementation end date: 30/06/2006

Information specific to this data set:

Obligation condition: reporting of this data element is optional for non-government residential mental health services and specialised mental health services provided by private hospitals that receive state or territory government funding.

Mental health establishments NMDS 2006-2007 NHIG, Superseded 23/10/2006

Implementation start date: 01/07/2006

Implementation end date: 30/06/2007

Information specific to this data set:

Obligation condition: reporting of this data element is optional for non-government residential mental health services and specialised mental health services provided by private hospitals that receive state or territory government funding.

Mental health establishments NMDS 2007-2008 NHIG, Standard 23/10/2006

Implementation start date: 01/07/2007

Information specific to this data set:

Obligation condition: reporting of this data element is optional for non-government residential mental health services and specialised mental health services provided by private hospitals that receive state or territory government funding.

Contract establishment identifier

Identifying and definitional attributes

<i>Metadata item type:</i>	Data Element
<i>Technical name:</i>	Contracted hospital care – organisation identifier, NNX[X]NNNNN
<i>METeOR identifier:</i>	270013
<i>Registration status:</i>	NHIG, Standard 01/03/2005
<i>Definition:</i>	The unique establishment identifier of the other hospital involved in the contracted care.

Data element concept attributes

<i>Data element concept:</i>	Contracted hospital care – organisation identifier
<i>Definition:</i>	The establishment identifier of the other hospital involved in the contracted care.
<i>Context:</i>	Admitted patient care and public hospital establishments
<i>Object class:</i>	Contracted hospital care
<i>Property:</i>	Organisation identifier

Value domain attributes

Representational attributes

<i>Representation class:</i>	Identifier
<i>Data type:</i>	String
<i>Format:</i>	NNX[X]NNNNN
<i>Maximum character length:</i>	9

Data element attributes

Collection and usage attributes

<i>Guide for use:</i>	The contracted hospital will record the establishment identifier of the contracting hospital. The contracting hospital will record the establishment identifier of the contracted hospital.
-----------------------	--

Relational attributes

<i>Related metadata references:</i>	Supersedes Contract establishment identifier, version 1, DE, NHDD, NHIMG, Superseded 01/03/2005
-------------------------------------	---

Contract procedure flag

Identifying and definitional attributes

<i>Metadata item type:</i>	Data Element
<i>Technical name:</i>	Episode of care (procedure)—contracted procedure flag, code N
<i>METeOR identifier:</i>	270473
<i>Registration status:</i>	NHIG, Standard 01/03/2005
<i>Definition:</i>	Designation that a procedure was not performed in this hospital but was performed by another hospital as a contracted service, as represented by a code.

Data element concept attributes

<i>Data element concept:</i>	Episode of care (procedure)—contracted procedure flag
<i>Definition:</i>	Designation that a procedure was not performed in this hospital but was performed by another hospital as a contracted service.
<i>Context:</i>	Admitted patient care.
<i>Object class:</i>	Episode of care
<i>Property:</i>	Contracted procedure flag

Value domain attributes

Representational attributes

<i>Representation class:</i>	Code						
<i>Data type:</i>	Number						
<i>Format:</i>	[N]						
<i>Maximum character length:</i>	1						
<i>Permissible values:</i>	<table><thead><tr><th>Value</th><th>Meaning</th></tr></thead><tbody><tr><td>1</td><td>Contracted admitted procedure</td></tr><tr><td>2</td><td>Contracted non-admitted procedure</td></tr></tbody></table>	Value	Meaning	1	Contracted admitted procedure	2	Contracted non-admitted procedure
Value	Meaning						
1	Contracted admitted procedure						
2	Contracted non-admitted procedure						

Data element attributes

Collection and usage attributes

<i>Guide for use:</i>	<p>Procedures performed at another hospital under contract (Hospital B) are recorded by both hospitals, but flagged by the contracting hospital only (Hospital A). This flag is to be used by the contracting hospital to indicate a procedure performed by a contracted hospital. It also indicates whether the procedure was performed as an admitted or non-admitted service.</p> <p>Allocation of procedure codes should not be affected by the contract status of an episode: the Australian Coding Standards should be applied when coding all episodes. In particular, procedures which would not otherwise be coded should not be coded solely because they were performed at another hospital under contract.</p> <p>Procedures performed by a health care service (i.e. not a recognised hospital) should be coded if appropriate. Some jurisdictions may require these to be separately identified and</p>
-----------------------	---

they could be distinguished from contracted hospital procedures through the use of an additional code in the contract procedure flag data item.

Relational attributes

Related metadata references:

Supersedes Contract procedure flag, version 1, DE, NHDD, NHIMG, Superseded 01/03/2005

Contract role

Identifying and definitional attributes

<i>Metadata item type:</i>	Data Element
<i>Technical name:</i>	Hospital – contract role, code A
<i>METeOR identifier:</i>	270114
<i>Registration status:</i>	NHIG, Standard 01/03/2005
<i>Definition:</i>	Whether the hospital is the purchaser of hospital care or the provider of an admitted or non-admitted service, as represented by a code.

Data element concept attributes

<i>Data element concept:</i>	Hospital – contract role
<i>Definition:</i>	Identifies whether the hospital is the purchaser of hospital care (contracting hospital) or the provider of an admitted or non-admitted service (contracted hospital).
<i>Context:</i>	Admitted patient care and public hospital establishments.
<i>Object class:</i>	Hospital
<i>Property:</i>	Contract role

Value domain attributes

Representational attributes

<i>Representation class:</i>	Code						
<i>Data type:</i>	String						
<i>Format:</i>	A						
<i>Maximum character length:</i>	1						
<i>Permissible values:</i>	<table><thead><tr><th>Value</th><th>Meaning</th></tr></thead><tbody><tr><td>A</td><td>Hospital A</td></tr><tr><td>B</td><td>Hospital B</td></tr></tbody></table>	Value	Meaning	A	Hospital A	B	Hospital B
Value	Meaning						
A	Hospital A						
B	Hospital B						

Collection and usage attributes

<i>Guide for use:</i>	CODE A Hospital A Hospital A is the contracting hospital (purchaser). CODE B Hospital B Hospital B is the contracted hospital (provider).
-----------------------	--

Data element attributes

Relational attributes

<i>Related metadata references:</i>	Supersedes Contract role, version 1, DE, NHDD, NHIMG, Superseded 01/03/2005 Is used in the formation of Episode of admitted patient care – inter-hospital contracted patient status, code N NHIG, Standard 01/03/2005
-------------------------------------	--

Contract type

Identifying and definitional attributes

<i>Metadata item type:</i>	Data Element
<i>Technical name:</i>	Hospital – contract type, code N
<i>METeOR identifier:</i>	270475
<i>Registration status:</i>	NHIG, Standard 01/03/2005
<i>Definition:</i>	The type of contract arrangement between contractor and the contracted hospital, as represented by a code.

Data element concept attributes

<i>Data element concept:</i>	Hospital – contract type
<i>Definition:</i>	The contract arrangement between the contractor and the contracted hospital. Contract types are distinguished by the physical movement of the patient between the contracting (where applicable) and contracted hospitals.
<i>Context:</i>	Admitted patient care and public hospital establishments.
<i>Object class:</i>	Hospital
<i>Property:</i>	Contract type

Value domain attributes

Representational attributes

<i>Representation class:</i>	Code												
<i>Data type:</i>	Number												
<i>Format:</i>	N												
<i>Maximum character length:</i>	1												
<i>Permissible values:</i>	<table><thead><tr><th>Value</th><th>Meaning</th></tr></thead><tbody><tr><td>1</td><td>Contract type B</td></tr><tr><td>2</td><td>Contract type ABA</td></tr><tr><td>3</td><td>Contract type AB</td></tr><tr><td>4</td><td>Contract type (A)B</td></tr><tr><td>5</td><td>Contract type BA</td></tr></tbody></table>	Value	Meaning	1	Contract type B	2	Contract type ABA	3	Contract type AB	4	Contract type (A)B	5	Contract type BA
Value	Meaning												
1	Contract type B												
2	Contract type ABA												
3	Contract type AB												
4	Contract type (A)B												
5	Contract type BA												

Collection and usage attributes

<i>Guide for use:</i>	<p>The contracting hospital (purchaser) is termed Hospital A. The contracted hospital (provider) is termed Hospital B.</p> <p>CODE 1 Contract Type B A health authority / other external purchaser contracts hospital B for admitted service which is funded outside the standard funding arrangements.</p> <p>CODE 2 Contract Type ABA Patient admitted by Hospital A. Hospital A contracts Hospital B for admitted or non-admitted patient service. Patient returns to Hospital A on completion of service by Hospital B.</p> <p>For example, a patient has a hip replacement at Hospital A, then receives aftercare at Hospital B, under contract to Hospital A. Complications arise and the patient returns to Hospital A for</p>
-----------------------	---

the remainder of care.

CODE 3 Contract Type AB

Patient admitted by Hospital A. Hospital A contracts Hospital B for admitted or non-admitted patient service. Patient does not return to Hospital A on completion of service by Hospital B.

For example, a patient has a hip replacement at Hospital A and then receives aftercare at Hospital B, under contract to Hospital A. Patient is separated from Hospital B.

CODE 4 Contract Type (A)B

This contract type occurs where a Hospital A contracts Hospital B for the whole episode of care. The patient does not attend Hospital A. For example, a patient is admitted for endoscopy at Hospital B under contract to Hospital A.

CODE 5 Contract Type BA

Hospital A contracts Hospital B for an admitted patient service following which the patient moves to Hospital A for remainder of care. For example, a patient is admitted to Hospital B for a gastric resection procedure under contract to Hospital A and Hospital A provides after care.

Data element attributes

Relational attributes

Related metadata references:

Supersedes Contract type, version 1, DE, NHDD, NHIMG,
Superseded 01/03/2005

Is used in the formation of Episode of admitted patient care –
inter-hospital contracted patient status, code N NHIG, Standard
01/03/2005

Contracted care commencement date

Identifying and definitional attributes

<i>Metadata item type:</i>	Data Element
<i>Technical name:</i>	Contracted hospital care – contracted care commencement date, DDMMYYYY
<i>METeOR identifier:</i>	270105
<i>Registration status:</i>	NHIG, Standard 01/03/2005
<i>Definition:</i>	The date the period of contracted care commenced.

Data element concept attributes

<i>Data element concept:</i>	Contracted hospital care – contracted care commencement date
<i>Definition:</i>	The date the period of contracted care commenced.
<i>Context:</i>	Admitted patient care
<i>Object class:</i>	Contracted hospital care
<i>Property:</i>	Contracted care commencement date

Value domain attributes

Representational attributes

<i>Representation class:</i>	Date
<i>Data type:</i>	Date/Time
<i>Format:</i>	DDMMYYYY
<i>Maximum character length:</i>	8

Data element attributes

Collection and usage attributes

<i>Guide for use:</i>	This metadata item is to be used by the contracting hospital to record the commencement date of the contracted hospital care and will be the admission date for the contracted hospital.
-----------------------	--

Relational attributes

<i>Related metadata references:</i>	Supersedes Contracted care commencement date, version 1, DE, NHDD, NHIMG, Superseded 01/03/2005
-------------------------------------	---

Contracted care completion date

Identifying and definitional attributes

<i>Metadata item type:</i>	Data Element
<i>Technical name:</i>	Contracted hospital care – contracted care completed date, DDMMYYYY
<i>METeOR identifier:</i>	270106
<i>Registration status:</i>	NHIG, Standard 01/03/2005
<i>Definition:</i>	The date the period of contracted care is completed.

Data element concept attributes

<i>Data element concept:</i>	Contracted hospital care – contracted care completed date
<i>Definition:</i>	The date the period of contracted care is completed.
<i>Context:</i>	Admitted patient care
<i>Object class:</i>	Contracted hospital care
<i>Property:</i>	Contracted care completed date

Value domain attributes

Representational attributes

<i>Representation class:</i>	Date
<i>Data type:</i>	Date/Time
<i>Format:</i>	DDMMYYYY
<i>Maximum character length:</i>	8

Data element attributes

Collection and usage attributes

<i>Guide for use:</i>	This metadata item is to be used by the contracting hospital to record the date of completion of the contracted hospital care and will be the separation date for the contracted hospital.
-----------------------	--

Relational attributes

<i>Related metadata references:</i>	Supersedes Contracted care completion date, version 1, DE, NHDD, NHIMG, Superseded 01/03/2005
-------------------------------------	---

Coronary artery disease—history of intervention or procedure

Identifying and definitional attributes

<i>Metadata item type:</i>	Data Element
<i>Technical name:</i>	Person – coronary artery disease intervention (history), code N
<i>METeOR identifier:</i>	270227
<i>Registration status:</i>	NHIG, Standard 01/03/2005
<i>Definition:</i>	Whether the individual has undergone a coronary artery by-pass grafting (CABG), angioplasty or stent, as represented by a code.

Data element concept attributes

<i>Data element concept:</i>	Person – coronary artery disease intervention
<i>Definition:</i>	Whether the individual has undergone a coronary artery by-pass grafting (CABG), angioplasty or stent.
<i>Context:</i>	Public health, health care and clinical settings.
<i>Object class:</i>	Person
<i>Property:</i>	Coronary artery disease intervention

Value domain attributes

Representational attributes

<i>Representation class:</i>	Code												
<i>Data type:</i>	Number												
<i>Format:</i>	N												
<i>Maximum character length:</i>	1												
<i>Permissible values:</i>	<table><thead><tr><th>Value</th><th>Meaning</th></tr></thead><tbody><tr><td>1</td><td>CABG, angioplasty or stent - undertaken in last 12 months</td></tr><tr><td>2</td><td>CABG, angioplasty or stent - undertaken prior to the last 12 months</td></tr><tr><td>3</td><td>CABG, angioplasty or stent - both within and prior to the last 12 months</td></tr><tr><td>4</td><td>No CABG, angioplasty or stent undertaken</td></tr><tr><td>9</td><td>Not stated/inadequately described</td></tr></tbody></table>	Value	Meaning	1	CABG, angioplasty or stent - undertaken in last 12 months	2	CABG, angioplasty or stent - undertaken prior to the last 12 months	3	CABG, angioplasty or stent - both within and prior to the last 12 months	4	No CABG, angioplasty or stent undertaken	9	Not stated/inadequately described
Value	Meaning												
1	CABG, angioplasty or stent - undertaken in last 12 months												
2	CABG, angioplasty or stent - undertaken prior to the last 12 months												
3	CABG, angioplasty or stent - both within and prior to the last 12 months												
4	No CABG, angioplasty or stent undertaken												
9	Not stated/inadequately described												
<i>Supplementary values:</i>													

Collection and usage attributes

<i>Comments:</i>	<p>CABG is known as 'bypass surgery' when a piece of vein (taken from the leg) or of an artery (taken from the chest or wrist) is used to form a connection between the aorta and the coronary artery distal to the obstructive lesion, making a bypass around the blockage. Angioplasty is an elective surgery technique of blood vessels reconstruction.</p> <p>Stenting is a non-surgical treatment used with balloon angioplasty or after, to treat coronary artery disease to widen a coronary artery. A stent is a small, expandable wire mesh tube that is inserted. The purpose of the stent is to help hold the</p>
------------------	---

newly treated artery open, reducing the risk of the artery re-closing (re-stenosis) over time.

Angioplasty with stenting typically leaves less than 10% of the original blockage in the artery (Heart Center Online).

These three procedures are commonly used to improve blood flow to the heart muscle when the heart's arteries are narrowed or blocked.

The sooner procedures are done, the greater the chances of saving heart muscle.

Data element attributes

Collection and usage attributes

Collection methods: Ask the individual if he/she has had a CABG, angioplasty or coronary stent. If so determine when it was undertaken within or prior to the last 12 months (or both).

Source and reference attributes

Submitting organisation: National Diabetes Data Working Group

Origin: National Diabetes Outcomes Quality Review Initiative (NDOQRIN) data dictionary.

Relational attributes

Related metadata references: Supersedes Coronary artery disease - history of intervention or procedure, version 1, DE, NHDD, NHIMG, Superseded 01/03/2005

Implementation in Data Set Specifications: Diabetes (clinical) DSS NHIG, Superseded 21/09/2005
Diabetes (clinical) DSS NHIG, Standard 21/09/2005

Country of birth

Identifying and definitional attributes

<i>Metadata item type:</i>	Data Element
<i>Technical name:</i>	Person – country of birth, code (SACC 1998) NNNN
<i>METeOR identifier:</i>	270277
<i>Registration status:</i>	NHIG, Standard 01/03/2005 NCSIMG, Standard 01/03/2005 NHDAMG, Standard 20/06/2005
<i>Definition:</i>	The country in which the person was born, as represented by a code.

Data element concept attributes

<i>Data element concept:</i>	Person – country of birth
<i>Definition:</i>	The country in which the person was born.
<i>Context:</i>	Country of birth is important in the study of access to services by different population sub-groups. Country of birth is the most easily collected and consistently reported of a range of possible data items that may indicate cultural or language diversity. Country of birth may be used in conjunction with other data such as period of residence in Australia, etc., to derive more sophisticated measures of access to (or need for) services by different population sub-groups.
<i>Object class:</i>	Person
<i>Property:</i>	Country of birth

Value domain attributes

Representational attributes

<i>Classification scheme:</i>	Standard Australian Classification of Countries 1998
<i>Representation class:</i>	Code
<i>Data type:</i>	Number
<i>Format:</i>	NNNN
<i>Maximum character length:</i>	4

Collection and usage attributes

<i>Guide for use:</i>	<p>The Standard Australian Classification of Countries 1998 (SACC) is a four-digit, three-level hierarchical structure specifying major group, minor group and country.</p> <p>A country, even if it comprises other discrete political entities such as states, is treated as a single unit for all data domain purposes. Parts of a political entity are not included in different groups. Thus, Hawaii is included in Northern America (as part of the identified country United States of America), despite being geographically close to and having similar social and cultural characteristics as the units classified to Polynesia.</p>
-----------------------	--

Data element attributes

Collection and usage attributes

Collection methods:

Some data collections ask respondents to specify their country of birth. In others, a pre-determined set of countries is specified as part of the question, usually accompanied by an 'other (please specify)' category.

Recommended questions are:

In which country were you/was the person/was (name) born?

Australia

Other (please specify)

Alternatively, a list of countries may be used based on, for example common Census responses.

In which country were you/was the person/was (name) born?

Australia

England

New Zealand

Italy

Viet Nam

Scotland

Greece

Germany

Philippines

India

Netherlands

Other (please specify)

In either case coding of data should conform to the SACC.

Sometimes respondents are simply asked to specify whether they were born in either 'English speaking' or 'non-English speaking' countries but this question is of limited use and this method of collection is not recommended.

Comments:

This metadata item is consistent with that used in the Australian Census of Population and Housing and is recommended for use whenever there is a requirement for comparison with Census data.

Source and reference attributes

Origin:

National Health Data Committee

National Community Services Data Committee

Reference documents:

Australian Bureau of Statistics 1998. Standard Australian Classification of Countries (SACC) (Cat. no. 1269.0), Canberra. Viewed 3 August 2005.

Relational attributes

Related metadata references:

Supersedes Country of birth, version 4, DE, Int. NCSDD & NHDD, NCSIMG & NHIMG, Superseded 01/03/2005

Implementation in Data Set Specifications:

Acute coronary syndrome (clinical) DSS NHIG, Standard 07/12/2005

Implementation start date: 07/12/2005

Acute coronary syndrome (clinical) DSS NHIG, Superseded 07/12/2005

Admitted patient care NMDS NHIG, Superseded 07/12/2005

Implementation start date: 01/07/2005
Implementation end date: 30/06/2006
Admitted patient care NMDS 2007-2008 NHIG, Standard
29/11/2006
Implementation start date: 01/07/2007
Admitted patient mental health care NMDS NHIG, Superseded
07/12/2005
Implementation start date: 01/07/2005
Implementation end date: 30/06/2006
Admitted patient mental health care NMDS NHIG, Superseded
23/10/2006
Implementation start date: 01/07/2006
Implementation end date: 30/06/2007
Admitted patient mental health care NMDS 2007-2008 NHIG,
Standard 23/10/2006
Implementation start date: 01/07/2007
Admitted patient palliative care NMDS NHIG, Superseded
07/12/2005
Implementation start date: 01/07/2005
Implementation end date: 30/06/2006
Admitted patient palliative care NMDS 2006-2007 NHIG,
Superseded 23/10/2006
Implementation start date: 01/07/2006
Implementation end date: 30/06/2007
Admitted patient palliative care NMDS 2007-08 NHIG,
Standard 23/10/2006
Implementation start date: 01/07/2007
Alcohol and other drug treatment services NMDS NHIG,
Superseded 21/03/2006
Implementation start date: 01/07/2005
Implementation end date: 30/06/2006
Alcohol and other drug treatment services NMDS NHIG,
Superseded 23/10/2006
Implementation start date: 01/07/2006
Implementation end date: 30/06/2007
Alcohol and other drug treatment services NMDS 2007-2008
NHIG, Standard 23/10/2006
Implementation start date: 01/07/2007
Cardiovascular disease (clinical) DSS NHIG, Superseded
15/02/2006
Cardiovascular disease (clinical) DSS NHIG, Standard
15/02/2006
Community mental health care 2004-2005 NHIG, Superseded
08/12/2004
Implementation start date: 01/07/2004
Implementation end date: 30/06/2005
Community mental health care NMDS 2005-2006 NHIG,
Superseded 07/12/2005

Implementation start date: 01/07/2005
Implementation end date: 30/06/2006
 Community mental health care NMDS 2006-2007 NHIG,
 Superseded 23/10/2006
Implementation start date: 01/07/2006
Implementation end date: 30/06/2007
 Community mental health care NMDS 2007-2008 NHIG,
 Standard 23/10/2006
Implementation start date: 01/07/2007
 Computer Assisted Telephone Interview demographic module
 DSS NHIG, Standard 04/05/2005
 Health care client identification NHIG, Superseded 04/05/2005
Implementation start date: 01/01/2003
Information specific to this data set:
 County of birth for newborn babies should be 'Australia'.
 Health care client identification DSS NHIG, Standard
 04/05/2005
 NCSIMG, Standard 03/10/2006
Information specific to this data set:
 Country of birth for newborn babies should be 'Australia'.
 Non-admitted patient emergency department care NMDS
 NHIG, Standard 24/03/2006
Implementation start date: 01/07/2006
 Non-admitted patient emergency department care NMDS
 NHIG, Superseded 07/12/2005
 Non-admitted patient emergency department care NMDS
 NHIG, Superseded 24/03/2006
Implementation start date: 01/07/2005
Implementation end date: 30/06/2006
 Perinatal NMDS NHIG, Superseded 07/12/2005
Implementation start date: 01/07/2005
Implementation end date: 30/06/2006
 Perinatal NMDS NHIG, Superseded 06/09/2006
Implementation start date: 01/07/2006
Implementation end date: 30/06/2007
 Perinatal NMDS 2007-2008 NHIG, Standard 06/09/2006
Implementation start date: 01/07/2007
 Residential mental health care NMDS 2005-2006 NHIG,
 Superseded 07/12/2005
Implementation start date: 01/07/2005
Implementation end date: 30/06/2006
 Residential mental health care NMDS 2006-2007 NHIG,
 Superseded 23/10/2006
Implementation start date: 01/07/2006
Implementation end date: 30/06/2007
 Residential mental health care NMDS 2007-2008 NHIG,
 Standard 23/10/2006

Implementation start date: 01/07/2007

Creatine kinase MB isoenzyme level (index code)

Identifying and definitional attributes

<i>Metadata item type:</i>	Data Element
<i>Technical name:</i>	Person – creatine kinase myocardial band isoenzyme level (measured), index code X[XXX]
<i>Synonymous names:</i>	Creatine kinase MB isoenzyme (CK-MB) - measured
<i>METeOR identifier:</i>	284903
<i>Registration status:</i>	NHIG, Standard 04/06/2004
<i>Definition:</i>	A person's measured creatine kinase myocardial band (CK-MB) isoenzyme level, as represented by an index.

Data element concept attributes

<i>Data element concept:</i>	Person – creatine kinase-myocardial band isoenzyme level
<i>Definition:</i>	A person's creatine kinase-myocardial band (CK-MB) isoenzyme level.
<i>Context:</i>	Health care and clinical settings.
<i>Object class:</i>	Person
<i>Property:</i>	Creatine kinase-myocardial band isoenzyme level

Value domain attributes

Representational attributes

<i>Representation class:</i>	Code				
<i>Data type:</i>	Number				
<i>Format:</i>	X[XXX]				
<i>Maximum character length:</i>	4				
<i>Supplementary values:</i>	<table><thead><tr><th>Value</th><th>Meaning</th></tr></thead><tbody><tr><td>9999</td><td>Not stated/inadequately described</td></tr></tbody></table>	Value	Meaning	9999	Not stated/inadequately described
Value	Meaning				
9999	Not stated/inadequately described				

Data element attributes

Collection and usage attributes

<i>Guide for use:</i>	CODE 88888 if test for CK-MB was not done on this admission. Measured in different units dependent upon laboratory methodology. When only one CK-MB level is recorded, this should be the peak level during admission.
-----------------------	--

Source and reference attributes

<i>Submitting organisation:</i>	Australian Institute of Health and Welfare
<i>Steward:</i>	The National Heart Foundation of Australia and The Cardiac Society of Australia and New Zealand

Relational attributes

<i>Related metadata references:</i>	Supersedes Creatine kinase MB isoenzyme (CK-MB) - units, version 1, DE, NHDD, NHIMG, Superseded 01/03/2005
-------------------------------------	--

Supersedes Creatine kinase MB isoenzyme (CK-MB) -
measured, version 1, DE, NHDD, NHIMG, Superseded
01/03/2005

See also Laboratory standard – upper limit of normal range for
creatinase kinase myocardial band isoenzyme, index code X[XXX]
NHIG, Standard 04/06/2004

*Implementation in Data Set
Specifications:*

Acute coronary syndrome (clinical) DSS NHIG, Standard
07/12/2005

Implementation start date: 07/12/2005

Information specific to this data set:

For Acute coronary syndrome (ACS) reporting, can be
used to determine diagnostic strata.

Acute coronary syndrome (clinical) DSS NHIG, Superseded
07/12/2005

Information specific to this data set:

For Acute coronary syndrome (ACS) reporting, can be
used to determine diagnostic strata.

Creatine kinase MB isoenzyme level (international units)

Identifying and definitional attributes

<i>Metadata item type:</i>	Data Element
<i>Technical name:</i>	Person – creatine kinase-myocardial band isoenzyme level (measured), total international units N[NNN]
<i>Synonymous names:</i>	Creatine kinase MB isoenzyme (CK-MB) - measured
<i>METeOR identifier:</i>	284905
<i>Registration status:</i>	NHIG, Standard 04/06/2004
<i>Definition:</i>	A person's measured creatine kinase-myocardial band (CK-MB) isoenzyme level in international units.

Data element concept attributes

<i>Data element concept:</i>	Person – creatine kinase-myocardial band isoenzyme level
<i>Definition:</i>	A person's creatine kinase-myocardial band (CK-MB) isoenzyme level.
<i>Context:</i>	Health care and clinical settings.
<i>Object class:</i>	Person
<i>Property:</i>	Creatine kinase-myocardial band isoenzyme level

Value domain attributes

Representational attributes

<i>Representation class:</i>	Total						
<i>Data type:</i>	Number						
<i>Format:</i>	N[NNN]						
<i>Maximum character length:</i>	4						
<i>Supplementary values:</i>	<table><thead><tr><th>Value</th><th>Meaning</th></tr></thead><tbody><tr><td>8888</td><td>Not measured</td></tr><tr><td>9999</td><td>Not stated/inadequately described</td></tr></tbody></table>	Value	Meaning	8888	Not measured	9999	Not stated/inadequately described
Value	Meaning						
8888	Not measured						
9999	Not stated/inadequately described						

Source and reference attributes

<i>Submitting organisation:</i>	Australian Institute of Health and Welfare
---------------------------------	--

Data element attributes

Collection and usage attributes

<i>Guide for use:</i>	CODE 8888 if test for CK-MB was not done on this admission. Measured in different units dependent upon laboratory methodology. When only one CK-MB level is recorded, this should be the peak level during admission.
-----------------------	---

Source and reference attributes

<i>Submitting organisation:</i>	Australian Institute of Health and Welfare
<i>Steward:</i>	The National Heart Foundation of Australia and The Cardiac Society of Australia and New Zealand

Relational attributes

Related metadata references:

See also Laboratory standard – upper limit of normal range for creatine kinase myocardial band isoenzyme, total international units N[NNN] NHIG, Standard 04/06/2004

Supersedes Creatine kinase MB isoenzyme (CK-MB) - measured, version 1, DE, NHDD, NHIMG, Superseded 01/03/2005

Supersedes Creatine kinase MB isoenzyme (CK-MB) - units, version 1, DE, NHDD, NHIMG, Superseded 01/03/2005

Implementation in Data Set Specifications:

Acute coronary syndrome (clinical) DSS NHIG, Standard 07/12/2005

Implementation start date: 07/12/2005

Information specific to this data set:

For Acute coronary syndrome (ACS) reporting, can be used to determine diagnostic strata.

Acute coronary syndrome (clinical) DSS NHIG, Superseded 07/12/2005

Information specific to this data set:

For Acute coronary syndrome (ACS) reporting, can be used to determine diagnostic strata.

Creatine kinase MB isoenzyme level (kCat per litre)

Identifying and definitional attributes

<i>Metadata item type:</i>	Data Element
<i>Technical name:</i>	Person – creatine kinase myocardial band isoenzyme level (measured), total kCat per litre N[NNN]
<i>Synonymous names:</i>	Creatine kinase MB isoenzyme (CK-MB) - measured
<i>METeOR identifier:</i>	284915
<i>Registration status:</i>	NHIG, Standard 04/06/2004
<i>Definition:</i>	A person's measured creatine kinase myocardial band (CK-MB) isoenzyme in kCat per litre.

Data element concept attributes

<i>Data element concept:</i>	Person – creatine kinase-myocardial band isoenzyme level
<i>Definition:</i>	A person's creatine kinase-myocardial band (CK-MB) isoenzyme level.
<i>Context:</i>	Health care and clinical settings.
<i>Object class:</i>	Person
<i>Property:</i>	Creatine kinase-myocardial band isoenzyme level

Value domain attributes

Representational attributes

<i>Representation class:</i>	Total						
<i>Data type:</i>	Number						
<i>Format:</i>	N[NNN]						
<i>Maximum character length:</i>	4						
<i>Supplementary values:</i>	<table><thead><tr><th>Value</th><th>Meaning</th></tr></thead><tbody><tr><td>8888</td><td>Not measured</td></tr><tr><td>9999</td><td>Not stated/inadequately described</td></tr></tbody></table>	Value	Meaning	8888	Not measured	9999	Not stated/inadequately described
Value	Meaning						
8888	Not measured						
9999	Not stated/inadequately described						
<i>Proposed unit of measure:</i>	kCat/L						

Source and reference attributes

<i>Submitting organisation:</i>	Australian Institute of Health and Welfare
---------------------------------	--

Data element attributes

Collection and usage attributes

<i>Guide for use:</i>	CODE 8888 if test for CK-MB was not done on this admission. Measured in different units dependent upon laboratory methodology. When only one CK-MB level is recorded, this should be the peak level during admission.
-----------------------	---

Source and reference attributes

<i>Submitting organisation:</i>	Australian Institute of Health and Welfare
<i>Steward:</i>	The National Heart Foundation of Australia and The Cardiac

Relational attributes

Related metadata references:

See also Laboratory standard – upper limit of normal range for creatine kinase myocardial band isoenzyme, total kCat per litre N[NNN] NHIG, Standard 04/06/2004

Supersedes Creatine kinase MB isoenzyme (CK-MB) - units, version 1, DE, NHDD, NHIMG, Superseded 01/03/2005

Supersedes Creatine kinase MB isoenzyme (CK-MB) - measured, version 1, DE, NHDD, NHIMG, Superseded 01/03/2005

Implementation in Data Set Specifications:

Acute coronary syndrome (clinical) DSS NHIG, Standard 07/12/2005

Implementation start date: 07/12/2005

Information specific to this data set:

For Acute coronary syndrome (ACS) reporting, can be used to determine diagnostic strata.

Acute coronary syndrome (clinical) DSS NHIG, Superseded 07/12/2005

Information specific to this data set:

For Acute coronary syndrome (ACS) reporting, can be used to determine diagnostic strata.

Creatine kinase MB isoenzyme level (micrograms per litre)

Identifying and definitional attributes

<i>Metadata item type:</i>	Data Element
<i>Technical name:</i>	Person – creatine kinase-myocardial band isoenzyme level (measured), total micrograms per litre N[NNNN]
<i>METeOR identifier:</i>	284921
<i>Registration status:</i>	NHIG, Standard 04/06/2004
<i>Definition:</i>	A person's measured creatine kinase-myocardial band (CK-MB) isoenzyme level in micrograms per litre.

Data element concept attributes

<i>Data element concept:</i>	Person – creatine kinase-myocardial band isoenzyme level
<i>Definition:</i>	A person's creatine kinase-myocardial band (CK-MB) isoenzyme level.
<i>Context:</i>	Health care and clinical settings.
<i>Object class:</i>	Person
<i>Property:</i>	Creatine kinase-myocardial band isoenzyme level

Value domain attributes

Representational attributes

<i>Representation class:</i>	Total						
<i>Data type:</i>	Number						
<i>Format:</i>	N[NNN]						
<i>Maximum character length:</i>	4						
<i>Supplementary values:</i>	<table><thead><tr><th>Value</th><th>Meaning</th></tr></thead><tbody><tr><td>9999</td><td>Not stated/inadequately described</td></tr><tr><td>8888</td><td>Not measured</td></tr></tbody></table>	Value	Meaning	9999	Not stated/inadequately described	8888	Not measured
Value	Meaning						
9999	Not stated/inadequately described						
8888	Not measured						
<i>Unit of measure:</i>	Microgram per litre (µg/L)						

Source and reference attributes

<i>Submitting organisation:</i>	Australian Institute of Health and Welfare
---------------------------------	--

Data element attributes

Collection and usage attributes

<i>Guide for use:</i>	CODE 8888 if test for CK-MB was not done on this admission. Measured in different units dependent upon laboratory methodology. When only one CK-MB level is recorded, this should be the peak level during admission.
-----------------------	---

Source and reference attributes

<i>Submitting organisation:</i>	Australian Institute of Health and Welfare
<i>Steward:</i>	The National Heart Foundation of Australia and The Cardiac

Relational attributes

Related metadata references:

Supersedes Creatine kinase MB isoenzyme (CK-MB) - measured, version 1, DE, NHDD, NHIMG, Superseded 01/03/2005

Supersedes Creatine kinase MB isoenzyme (CK-MB) - units, version 1, DE, NHDD, NHIMG, Superseded 01/03/2005

See also Laboratory standard – upper limit of normal range for creatine kinase myocardial band isoenzyme, total micrograms per litre N[NNN] NHIG, Standard 04/06/2004

Implementation in Data Set Specifications:

Acute coronary syndrome (clinical) DSS NHIG, Standard 07/12/2005

Implementation start date: 07/12/2005

Information specific to this data set:

For Acute coronary syndrome (ACS) reporting, can be used to determine diagnostic strata.

Acute coronary syndrome (clinical) DSS NHIG, Superseded 07/12/2005

Information specific to this data set:

For Acute coronary syndrome (ACS) reporting, can be used to determine diagnostic strata.

Creatine kinase MB isoenzyme level (nanograms per decilitre)

Identifying and definitional attributes

<i>Metadata item type:</i>	Data Element
<i>Technical name:</i>	Person – creatine kinase myocardial band isoenzyme level (measured), total nanograms per decilitre N[NNN]
<i>Synonymous names:</i>	Creatine kinase MB isoenzyme (CK-MB) - measured
<i>METeOR identifier:</i>	284923
<i>Registration status:</i>	NHIG, Standard 04/06/2004
<i>Definition:</i>	A person's measured creatine kinase myocardial band (CK-MB) isoenzyme in nanograms per decilitre.

Data element concept attributes

<i>Data element concept:</i>	Person – creatine kinase-myocardial band isoenzyme level
<i>Definition:</i>	A person's creatine kinase-myocardial band (CK-MB) isoenzyme level.
<i>Context:</i>	Health care and clinical settings.
<i>Object class:</i>	Person
<i>Property:</i>	Creatine kinase-myocardial band isoenzyme level

Value domain attributes

Representational attributes

<i>Representation class:</i>	Total						
<i>Data type:</i>	Number						
<i>Format:</i>	N[NNN]						
<i>Maximum character length:</i>	4						
<i>Supplementary values:</i>	<table><thead><tr><th>Value</th><th>Meaning</th></tr></thead><tbody><tr><td>8888</td><td>Not measured</td></tr><tr><td>9999</td><td>Not stated/inadequately described</td></tr></tbody></table>	Value	Meaning	8888	Not measured	9999	Not stated/inadequately described
Value	Meaning						
8888	Not measured						
9999	Not stated/inadequately described						
<i>Unit of measure:</i>	Nanogram per decilitre (ng/dl)						

Data element attributes

Collection and usage attributes

<i>Guide for use:</i>	CODE 8888 if test for CK-MB was not done on this admission. Measured in different units dependent upon laboratory methodology. When only one CK-MB level is recorded, this should be the peak level during admission.
-----------------------	---

Source and reference attributes

<i>Submitting organisation:</i>	Australian Institute of Health and Welfare
<i>Steward:</i>	The National Heart Foundation of Australia and The Cardiac Society of Australia and New Zealand

Relational attributes

Related metadata references:

Supersedes Creatine kinase MB isoenzyme (CK-MB) - units, version 1, DE, NHDD, NHIMG, Superseded 01/03/2005

Supersedes Creatine kinase MB isoenzyme (CK-MB) - measured, version 1, DE, NHDD, NHIMG, Superseded 01/03/2005

Implementation in Data Set Specifications:

Acute coronary syndrome (clinical) DSS NHIG, Standard 07/12/2005

Implementation start date: 07/12/2005

Information specific to this data set:

For Acute coronary syndrome (ACS) reporting, can be used to determine diagnostic strata.

Acute coronary syndrome (clinical) DSS NHIG, Superseded 07/12/2005

Information specific to this data set:

For Acute coronary syndrome (ACS) reporting, can be used to determine diagnostic strata.

Creatine kinase MB isoenzyme level (percentage)

Identifying and definitional attributes

<i>Metadata item type:</i>	Data Element
<i>Technical name:</i>	Person – creatine kinase myocardial band isoenzyme level (measured), percentage N[NNN]
<i>Synonymous names:</i>	Creatine kinase MB isoenzyme (CK-MB) - measured
<i>METeOR identifier:</i>	284913
<i>Registration status:</i>	NHIG, Standard 04/06/2004
<i>Definition:</i>	A person's measured creatine kinase myocardial band (CK-MB) isoenzyme as a percentage.

Data element concept attributes

<i>Data element concept:</i>	Person – creatine kinase-myocardial band isoenzyme level
<i>Definition:</i>	A person's creatine kinase-myocardial band (CK-MB) isoenzyme level.
<i>Context:</i>	Health care and clinical settings.
<i>Object class:</i>	Person
<i>Property:</i>	Creatine kinase-myocardial band isoenzyme level

Value domain attributes

Representational attributes

<i>Representation class:</i>	Percentage						
<i>Data type:</i>	Number						
<i>Format:</i>	N[NNN]						
<i>Maximum character length:</i>	4						
<i>Supplementary values:</i>	<table><thead><tr><th>Value</th><th>Meaning</th></tr></thead><tbody><tr><td>8888</td><td>Not measured</td></tr><tr><td>9999</td><td>Not stated/inadequately described</td></tr></tbody></table>	Value	Meaning	8888	Not measured	9999	Not stated/inadequately described
Value	Meaning						
8888	Not measured						
9999	Not stated/inadequately described						

Source and reference attributes

<i>Submitting organisation:</i>	Australian Institute of Health and Welfare
---------------------------------	--

Data element attributes

Collection and usage attributes

<i>Guide for use:</i>	CODE 8888 if test for CK-MB was not done on this admission. Measured in different units dependent upon laboratory methodology. When only one CK-MB level is recorded, this should be the peak level during admission.
-----------------------	---

Source and reference attributes

<i>Submitting organisation:</i>	Australian Institute of Health and Welfare
<i>Steward:</i>	The National Heart Foundation of Australia and The Cardiac Society of Australia and New Zealand

Relational attributes

Related metadata references:

Supersedes Creatine kinase MB isoenzyme (CK-MB) - units, version 1, DE, NHDD, NHIMG, Superseded 01/03/2005

Supersedes Creatine kinase MB isoenzyme (CK-MB) - measured, version 1, DE, NHDD, NHIMG, Superseded 01/03/2005

See also Laboratory standard – upper limit of normal range for creatine kinase myocardial band isoenzyme, percentage N[NNN] NHIG, Standard 04/06/2004

Implementation in Data Set Specifications:

Acute coronary syndrome (clinical) DSS NHIG, Standard 07/12/2005

Implementation start date: 07/12/2005

Information specific to this data set:

For Acute coronary syndrome (ACS) reporting, can be used to determine diagnostic strata.

Acute coronary syndrome (clinical) DSS NHIG, Superseded 07/12/2005

Information specific to this data set:

For Acute coronary syndrome (ACS) reporting, can be used to determine diagnostic strata.

Creatine kinase MB isoenzyme—upper limit of normal range (index code)

Identifying and definitional attributes

<i>Metadata item type:</i>	Data Element
<i>Technical name:</i>	Laboratory standard – upper limit of normal range for creatine kinase myocardial band isoenzyme, index code X[XXX]
<i>Synonymous names:</i>	Creatine kinase MB isoenzyme (CK-MB) - units
<i>METeOR identifier:</i>	284931
<i>Registration status:</i>	NHIG, Standard 04/06/2004
<i>Definition:</i>	Laboratory standard for the value of creatine kinase myocardial band (CK-MB) isoenzyme measured as an index that is the upper boundary of the normal reference range.

Data element concept attributes

<i>Data element concept:</i>	Laboratory standard – upper limit of normal range for creatine kinase myocardial band isoenzyme
<i>Definition:</i>	Laboratory standard for the value of creatine kinase myocardial band (CK-MB) isoenzyme that is the upper boundary of the normal reference range.
<i>Context:</i>	Health care and clinical settings.
<i>Object class:</i>	Laboratory standard
<i>Property:</i>	Upper limit of normal range for creatine kinase myocardial band isoenzyme

Value domain attributes

Representational attributes

<i>Representation class:</i>	Code				
<i>Data type:</i>	Number				
<i>Format:</i>	X[XXX]				
<i>Maximum character length:</i>	4				
<i>Supplementary values:</i>	<table><thead><tr><th>Value</th><th>Meaning</th></tr></thead><tbody><tr><td>9999</td><td>Not stated/inadequately described</td></tr></tbody></table>	Value	Meaning	9999	Not stated/inadequately described
Value	Meaning				
9999	Not stated/inadequately described				

Data element attributes

Collection and usage attributes

<i>Guide for use:</i>	Record the upper limit of the creatine kinase myocardial band (CK-MB) normal reference range for the testing laboratory.
-----------------------	--

Source and reference attributes

<i>Submitting organisation:</i>	Acute coronary syndrome data working group.
<i>Steward:</i>	The National Heart Foundation of Australia and The Cardiac Society of Australia and New Zealand

Relational attributes

Related metadata references:

Supersedes Creatine kinase MB isoenzyme (CK-MB) - upper limit of normal range, version 1, DE, NHDD, NHIMG, Superseded 01/03/2005

Implementation in Data Set Specifications:

Acute coronary syndrome (clinical) DSS NHIG, Standard 07/12/2005

Implementation start date: 07/12/2005

Acute coronary syndrome (clinical) DSS NHIG, Superseded 07/12/2005

Creatine kinase MB isoenzyme—upper limit of normal range (international units)

Identifying and definitional attributes

<i>Metadata item type:</i>	Data Element
<i>Technical name:</i>	Laboratory standard – upper limit of normal range for creatine kinase myocardial band isoenzyme, total international units N[NNN]
<i>METeOR identifier:</i>	284959
<i>Registration status:</i>	NHIG, Standard 04/06/2004
<i>Definition:</i>	Laboratory standard for the value of creatine kinase myocardial band (CK-MB) isoenzyme measured in international units (IU) that is the upper boundary of the normal reference range.

Data element concept attributes

<i>Data element concept:</i>	Laboratory standard – upper limit of normal range for creatine kinase myocardial band isoenzyme
<i>Definition:</i>	Laboratory standard for the value of creatine kinase myocardial band (CK-MB) isoenzyme that is the upper boundary of the normal reference range.
<i>Context:</i>	Health care and clinical settings.
<i>Object class:</i>	Laboratory standard
<i>Property:</i>	Upper limit of normal range for creatine kinase myocardial band isoenzyme

Value domain attributes

Representational attributes

<i>Representation class:</i>	Total						
<i>Data type:</i>	Number						
<i>Format:</i>	N[NNN]						
<i>Maximum character length:</i>	4						
<i>Supplementary values:</i>	<table><thead><tr><th>Value</th><th>Meaning</th></tr></thead><tbody><tr><td>8888</td><td>Not measured</td></tr><tr><td>9999</td><td>Not stated/inadequately described</td></tr></tbody></table>	Value	Meaning	8888	Not measured	9999	Not stated/inadequately described
Value	Meaning						
8888	Not measured						
9999	Not stated/inadequately described						

Source and reference attributes

<i>Submitting organisation:</i>	Australian Institute of Health and Welfare
---------------------------------	--

Data element attributes

Collection and usage attributes

<i>Guide for use:</i>	Record the upper limit of the creatine kinase myocardial band (CK-MB) normal reference range for the testing laboratory.
-----------------------	--

Source and reference attributes

<i>Submitting organisation:</i>	Acute coronary syndrome data working group.
---------------------------------	---

Steward:

The National Heart Foundation of Australia and The Cardiac Society of Australia and New Zealand

Relational attributes

Related metadata references:

Supersedes Creatine kinase MB isoenzyme (CK-MB) - upper limit of normal range, version 1, DE, NHDD, NHIMG, Superseded 01/03/2005

Implementation in Data Set Specifications:

Acute coronary syndrome (clinical) DSS NHIG, Standard 07/12/2005

Implementation start date: 07/12/2005

Acute coronary syndrome (clinical) DSS NHIG, Superseded 07/12/2005

Creatine kinase MB isoenzyme—upper limit of normal range (kCat per litre)

Identifying and definitional attributes

<i>Metadata item type:</i>	Data Element
<i>Technical name:</i>	Laboratory standard – upper limit of normal range for creatine kinase myocardial band isoenzyme, total kCat per litre N[NNN]
<i>METeOR identifier:</i>	284963
<i>Registration status:</i>	NHIG, Standard 04/06/2004
<i>Definition:</i>	Laboratory standard for the value of creatine kinase myocardial band (CK-MB) isoenzyme in kCat per litre that is the upper boundary of the normal reference range.

Data element concept attributes

<i>Data element concept:</i>	Laboratory standard – upper limit of normal range for creatine kinase myocardial band isoenzyme
<i>Definition:</i>	Laboratory standard for the value of creatine kinase myocardial band (CK-MB) isoenzyme that is the upper boundary of the normal reference range.
<i>Context:</i>	Health care and clinical settings.
<i>Object class:</i>	Laboratory standard
<i>Property:</i>	Upper limit of normal range for creatine kinase myocardial band isoenzyme

Value domain attributes

Representational attributes

<i>Representation class:</i>	Total						
<i>Data type:</i>	Number						
<i>Format:</i>	N[NNN]						
<i>Maximum character length:</i>	4						
<i>Supplementary values:</i>	<table><thead><tr><th>Value</th><th>Meaning</th></tr></thead><tbody><tr><td>8888</td><td>Not measured</td></tr><tr><td>9999</td><td>Not stated/inadequately described</td></tr></tbody></table>	Value	Meaning	8888	Not measured	9999	Not stated/inadequately described
Value	Meaning						
8888	Not measured						
9999	Not stated/inadequately described						
<i>Proposed unit of measure:</i>	kCat/L						

Source and reference attributes

<i>Submitting organisation:</i>	Australian Institute of Health and Welfare
---------------------------------	--

Data element attributes

Collection and usage attributes

<i>Guide for use:</i>	Record the upper limit of the creatine kinase myocardial band (CK-MB) normal reference range for the testing laboratory.
-----------------------	--

Source and reference attributes

Submitting organisation:

Acute coronary syndrome data working group.

Steward:

The National Heart Foundation of Australia and The Cardiac Society of Australia and New Zealand

Relational attributes

Related metadata references:

Supersedes Creatine kinase MB isoenzyme (CK-MB) - upper limit of normal range, version 1, DE, NHDD, NHIMG, Superseded 01/03/2005

Implementation in Data Set Specifications:

Acute coronary syndrome (clinical) DSS NHIG, Standard 07/12/2005

Implementation start date: 07/12/2005

Acute coronary syndrome (clinical) DSS NHIG, Superseded 07/12/2005

Creatine kinase MB isoenzyme—upper limit of normal range (micrograms per litre)

Identifying and definitional attributes

<i>Metadata item type:</i>	Data Element
<i>Technical name:</i>	Laboratory standard – upper limit of normal range for creatine kinase myocardial band isoenzyme, total micrograms per litre N[NNN]
<i>METeOR identifier:</i>	284965
<i>Registration status:</i>	NHIG, Standard 04/06/2004
<i>Definition:</i>	Laboratory standard for the value of creatine kinase myocardial band (CK-MB) isoenzyme measured in microgram per litre that is the upper boundary of the normal reference range.

Data element concept attributes

<i>Data element concept:</i>	Laboratory standard – upper limit of normal range for creatine kinase myocardial band isoenzyme
<i>Definition:</i>	Laboratory standard for the value of creatine kinase myocardial band (CK-MB) isoenzyme that is the upper boundary of the normal reference range.
<i>Context:</i>	Health care and clinical settings.
<i>Object class:</i>	Laboratory standard
<i>Property:</i>	Upper limit of normal range for creatine kinase myocardial band isoenzyme

Value domain attributes

Representational attributes

<i>Representation class:</i>	Total						
<i>Data type:</i>	Number						
<i>Format:</i>	N[NNN]						
<i>Maximum character length:</i>	4						
<i>Supplementary values:</i>	<table><thead><tr><th>Value</th><th>Meaning</th></tr></thead><tbody><tr><td>9999</td><td>Not stated/inadequately described</td></tr><tr><td>8888</td><td>Not measured</td></tr></tbody></table>	Value	Meaning	9999	Not stated/inadequately described	8888	Not measured
Value	Meaning						
9999	Not stated/inadequately described						
8888	Not measured						
<i>Unit of measure:</i>	Microgram per litre (µg/L)						

Source and reference attributes

<i>Submitting organisation:</i>	Australian Institute of Health and Welfare
---------------------------------	--

Data element attributes

Collection and usage attributes

<i>Guide for use:</i>	Record the upper limit of the creatine kinase myocardial band (CK-MB) normal reference range for the testing laboratory.
-----------------------	--

Source and reference attributes

Submitting organisation:

Acute coronary syndrome data working group.

Steward:

The National Heart Foundation of Australia and The Cardiac Society of Australia and New Zealand

Relational attributes

Related metadata references:

Supersedes Creatine kinase MB isoenzyme (CK-MB) - upper limit of normal range, version 1, DE, NHDD, NHIMG, Superseded 01/03/2005

Implementation in Data Set Specifications:

Acute coronary syndrome (clinical) DSS NHIG, Standard 07/12/2005

Implementation start date: 07/12/2005

Acute coronary syndrome (clinical) DSS NHIG, Superseded 07/12/2005

Creatine kinase MB isoenzyme—upper limit of normal range (nanograms per decilitre)

Identifying and definitional attributes

<i>Metadata item type:</i>	Data Element
<i>Technical name:</i>	Laboratory standard – upper limit of normal range for creatine kinase myocardial band isoenzyme, total nanograms per decilitre N[NNN]
<i>METeOR identifier:</i>	285957
<i>Registration status:</i>	NHIG, Standard 04/06/2004
<i>Definition:</i>	Laboratory standard for the value of creatine kinase myocardial band (CK-MB) isoenzyme measured in nanograms per decilitre that is the upper boundary of the normal reference range.

Data element concept attributes

<i>Data element concept:</i>	Laboratory standard – upper limit of normal range for creatine kinase myocardial band isoenzyme
<i>Definition:</i>	Laboratory standard for the value of creatine kinase myocardial band (CK-MB) isoenzyme that is the upper boundary of the normal reference range.
<i>Context:</i>	Health care and clinical settings.
<i>Object class:</i>	Laboratory standard
<i>Property:</i>	Upper limit of normal range for creatine kinase myocardial band isoenzyme

Value domain attributes

Representational attributes

<i>Representation class:</i>	Total						
<i>Data type:</i>	Number						
<i>Format:</i>	N[NNN]						
<i>Maximum character length:</i>	4						
<i>Supplementary values:</i>	<table><thead><tr><th>Value</th><th>Meaning</th></tr></thead><tbody><tr><td>8888</td><td>Not measured</td></tr><tr><td>9999</td><td>Not stated/inadequately described</td></tr></tbody></table>	Value	Meaning	8888	Not measured	9999	Not stated/inadequately described
Value	Meaning						
8888	Not measured						
9999	Not stated/inadequately described						
<i>Unit of measure:</i>	Nanogram per decilitre (ng/dl)						

Data element attributes

Collection and usage attributes

<i>Guide for use:</i>	Record the upper limit of the creatine kinase myocardial band (CK-MB) normal reference range for the testing laboratory.
-----------------------	--

Source and reference attributes

<i>Submitting organisation:</i>	Acute coronary syndrome data working group.
<i>Steward:</i>	The National Heart Foundation of Australia and The Cardiac

Society of Australia and New Zealand

Relational attributes

Related metadata references:

Supersedes Creatine kinase MB isoenzyme (CK-MB) - upper limit of normal range, version 1, DE, NHDD, NHIMG, Superseded 01/03/2005

Supersedes Creatine kinase MB isoenzyme (CK-MB) - units, version 1, DE, NHDD, NHIMG, Superseded 01/03/2005

Implementation in Data Set Specifications:

Acute coronary syndrome (clinical) DSS NHIG, Standard 07/12/2005

Implementation start date: 07/12/2005

Acute coronary syndrome (clinical) DSS NHIG, Superseded 07/12/2005

Creatine kinase MB isoenzyme—upper limit of normal range (percentage)

Identifying and definitional attributes

<i>Metadata item type:</i>	Data Element
<i>Technical name:</i>	Laboratory standard – upper limit of normal range for creatine kinase myocardial band isoenzyme, percentage N[NNN]
<i>METeOR identifier:</i>	284961
<i>Registration status:</i>	NHIG, Standard 04/06/2004
<i>Definition:</i>	Laboratory standard for the value of creatine kinase myocardial band (CK-MB) isoenzyme measured as a percentage that is the upper boundary of the normal reference range.

Data element concept attributes

<i>Data element concept:</i>	Laboratory standard – upper limit of normal range for creatine kinase myocardial band isoenzyme
<i>Definition:</i>	Laboratory standard for the value of creatine kinase myocardial band (CK-MB) isoenzyme that is the upper boundary of the normal reference range.
<i>Context:</i>	Health care and clinical settings.
<i>Object class:</i>	Laboratory standard
<i>Property:</i>	Upper limit of normal range for creatine kinase myocardial band isoenzyme

Value domain attributes

Representational attributes

<i>Representation class:</i>	Percentage	
<i>Data type:</i>	Number	
<i>Format:</i>	N[NNN]	
<i>Maximum character length:</i>	4	
<i>Supplementary values:</i>	Value	Meaning
	8888	Not measured
	9999	Not stated/inadequately described

Source and reference attributes

<i>Submitting organisation:</i>	Australian Institute of Health and Welfare
---------------------------------	--

Data element attributes

Collection and usage attributes

<i>Guide for use:</i>	Record the upper limit of the creatine kinase myocardial band (CK-MB) normal reference range for the testing laboratory.
-----------------------	--

Source and reference attributes

<i>Submitting organisation:</i>	Acute coronary syndrome data working group.
---------------------------------	---

Steward:

The National Heart Foundation of Australia and The Cardiac Society of Australia and New Zealand

Relational attributes

Related metadata references:

Supersedes Creatine kinase MB isoenzyme (CK-MB) - upper limit of normal range, version 1, DE, NHDD, NHIMG, Superseded 01/03/2005

Implementation in Data Set Specifications:

Acute coronary syndrome (clinical) DSS NHIG, Standard 07/12/2005

Implementation start date: 07/12/2005

Acute coronary syndrome (clinical) DSS NHIG, Superseded 07/12/2005

Creatinine serum level (measured)

Identifying and definitional attributes

<i>Metadata item type:</i>	Data Element
<i>Technical name:</i>	Person – creatinine serum level, micromoles per litre NN[NN]
<i>METeOR identifier:</i>	270392
<i>Registration status:</i>	NHIG, Standard 01/03/2005
<i>Definition:</i>	A person's serum creatinine level measured in micromoles per litre ($\mu\text{mol/L}$).

Data element concept attributes

<i>Data element concept:</i>	Person – creatinine serum level
<i>Definition:</i>	A person's serum creatinine level.
<i>Context:</i>	Clinical settings and population survey: Serum creatinine can be used to help determine renal function. Serum creatinine by itself is an insensitive measure of renal function because it does not increase until more than 50% of renal function has been lost.
<i>Object class:</i>	Person
<i>Property:</i>	Creatinine serum level

Value domain attributes

Representational attributes

<i>Representation class:</i>	Total
<i>Data type:</i>	String
<i>Format:</i>	NN[NN]
<i>Maximum character length:</i>	4
<i>Unit of measure:</i>	Micromole per litre ($\mu\text{mol/L}$)

Data element attributes

Collection and usage attributes

<i>Guide for use:</i>	There is no agreed standard as to which units serum creatinine should be recorded in. Note: If the measurement is obtained in mmol/L it is to be multiplied by 1000.
<i>Collection methods:</i>	Measurement of creatinine should be carried out by laboratories, or practices, which have been accredited to perform these tests by the National Association of Testing Authority. <ul style="list-style-type: none">• Single venous blood test taken at the time of other screening blood tests.• Fasting not required.
<i>Comments:</i>	Serum creatinine together with a patient's age, weight and sex can be used to calculate glomerular filtration rate (GFR), which is an indicator of renal status/ function. The calculation uses the Cockcroft-Gault formula.

Creatinine is normally produced in fairly constant amounts in the muscles, as a result the breakdown of phosphocreatine. It passes into the blood and is excreted in the urine. Serum creatinine can be used to help determine renal function. The elevation in the creatinine level in the blood indicates disturbance in kidney function.

GFR decreases with age, but serum creatinine remains relatively stable. When serum creatinine is measured, renal function in the elderly tends to be overestimated, and GFR should be used to assess renal function, according to the Cockcroft-Gault formula:

$$\text{GFR (ml/min)} = \frac{(140 - \text{age [yrs]}) \times \text{body wt (kg)}}{814 \times \text{serum creatinine (mmol/l)}} \quad [\times 0.85 \text{ (for women)}]$$

To determine chronic renal impairment

GFR > 90ml/min - normal

GFR >60 - 90ml/min - mild renal impairment

GFR >30 - 60ml/min - moderate renal impairment

GFR 0 - 30 ml/min - severe renal impairment

Note: The above GFR measurement should be for a period greater than 3 months. GFR may also be assessed by 24-hour creatinine clearance adjusted for body surface area.

In general, patients with GFR

Patients should be assessed for the complications of chronic renal impairment including anaemia, hyperparathyroidism and be referred for specialist management if required.

Patients with rapidly declining renal function or clinical features to suggest that residual renal function may decline rapidly (ie. hypertensive, proteinuric (>1g/24hours), significant comorbid illness) should be considered for referral to a nephrologist well before function declines to less than 30ml/min. (Draft CARI Guidelines 2002. Australian Kidney Foundation). Patients in whom the cause of renal impairment is uncertain should be referred to a nephrologist for assessment.

Source and reference attributes

Submitting organisation:

Cardiovascular Data Working Group

National Diabetes Data Working Group

Origin:

Caring for Australians with Renal Impairment (CARI) Guidelines. Australian Kidney Foundation

Relational attributes

Related metadata references:

Supersedes Creatinine serum - measured, version 1, DE, NHDD, NHIMG, Superseded 01/03/2005

Implementation in Data Set Specifications:

Acute coronary syndrome (clinical) DSS NHIG, Standard 07/12/2005

Implementation start date: 07/12/2005

Information specific to this data set:

In settings where the monitoring of a person's health is ongoing and where a measure can change over time (such as general practice), the Service contact – service contact

date, DDMMYYYY should be recorded.

Record absolute result of the most recent serum creatinine measurement in the last 12 months to the nearest $\mu\text{mol/L}$ (micromoles per litre).

Acute coronary syndrome (clinical) DSS NHIG, Superseded
07/12/2005

Cardiovascular disease (clinical) DSS NHIG, Superseded
15/02/2006

Information specific to this data set:

In settings where the monitoring of a person's health is ongoing and where a measure can change over time (such as general practice), the Service contact – service contact date, DDMMYYYY should be recorded.

Record absolute result of the most recent serum creatinine measurement in the last 12 months to the nearest $\mu\text{mol/L}$ (micromoles per litre).

Cardiovascular disease (clinical) DSS NHIG, Standard
15/02/2006

Information specific to this data set:

In settings where the monitoring of a person's health is ongoing and where a measure can change over time (such as general practice), the Service contact – service contact date, DDMMYYYY should be recorded.

Record absolute result of the most recent serum creatinine measurement in the last 12 months to the nearest $\mu\text{mol/L}$ (micromoles per litre).

Diabetes (clinical) DSS NHIG, Superseded 21/09/2005

Diabetes (clinical) DSS NHIG, Standard 21/09/2005

Information specific to this data set:

In settings where the monitoring of a person's health is ongoing and where a measure can change over time (such as general practice), the Service contact – service contact date, DDMMYYYY should be recorded.

Record absolute result of the most recent serum creatinine measurement in the last 12 months to the nearest $\mu\text{mol/L}$ (micromoles per litre).