Person with cancer—hereditary genetic events type, code N[N]

Exported from METEOR (AIHW's Metadata Online Registry)		
© Australian Institute of Health and Welfare 2024		
This product, excluding the AlHW logo, Commonwealth Coat of Arms and any material owned by a third party or protected by a trademark, has been released under a Creative Commons BY 4.0 (CC BY 4.0) licence. Excluded material owned by third parties may include, for example, design and layout, images obtained under licence from third parties and signatures. We have made all reasonable efforts to identify and label material owned by third parties.		
You may distribute, remix and build on this website's material but must attribute the AlHW as the copyright holder, in line with our		

meteor@aihw.gov.au.

attribution policy. The full terms and conditions of this licence are available at https://creativecommons.org/licenses/by/4.0/.

Enquiries or comments on the METEOR metadata or download should be directed to the METEOR team at

Enquiries relating to copyright should be addressed to info@aihw.gov.au.

Person with cancer—hereditary genetic events type, code N[N]

Identifying and definitional attributes

Metadata item type: Data Element

Short name: Hereditary genetic events type code

METEOR identifier: 467855

Registration status: Health, Standard 04/02/2015

Definition: The type of hereditary genetic event present in a <u>first degree relative</u> or <u>second</u>

degree relative of a person with cancer, as represented by a code.

Data Element Concept: Person with cancer—hereditary genetic events type

Value Domain: Genetic event type code N[N]

Value domain attributes

Representational attributes

Representation class: Code

Data type: Number

Format: N[N]

Format:	N[N]	
Maximum character length:	2	
	Value	Meaning
Permissible values:	1	Peutz-Jehgers syndrome
	2	B-Raf mutation
	3	Undescended testes
	4	Von Recklinghausen syndrome
	5	Xeroderma pigmentosa
	6	Bloom syndrome
	7	Ataxia telangiectasia
	8	Beckwith-Wiedemann syndrome
	9	Down syndrome
	10	Klinefelter syndrome
	11	Neurofibromatosis type I (NF-1)
	12	Von Hippel–Lindau disease (VHL)
	13	RET proto-oncogene
	14	Adenomatous polyposis coli (APC)
	15	Hereditary nonpolyposis colorectal cancer (HNPCC or Lynch syndrome)
	16	Breast cancer gene 1 (BRCA1)
	17	Breast cancer gene 2 (BRCA2)
	18	Neurofibromatosis
	19	Li-Fraumeni syndrome
	20	Fanconi anaemia

Other genetic events or syndromes

88

Supplementary values: 97 No genetic events or syndromes

> Unknown 98

99 Genetic event or syndrome recorded but type

inadequately described

Source and reference attributes

Submitting organisation: Cancer Australia

Data element attributes

Collection and usage attributes

Guide for use: Record the type of hereditary genetic event or syndrome that is present in any first

degree or second degree relative.

Collection methods: Collect from patient medical records.

Source and reference attributes

Submitting organisation: Cancer Australia

Relational attributes

Related metadata references:

See also Person with cancer—family history of hereditary genetic events indicator,

yes/no/unknown/not stated/inadequately described code N

Health, Standard 04/02/2015

See also Person with cancer—hereditary genetic events type, text X[X(39)]

Health, Standard 04/02/2015

Implementation in Data Set Adolescent and young adult cancer (clinical) DSS

Specifications:

Health, Superseded 14/05/2015

Conditional obligation: Complete if Person with cancer—family history of hereditary genetic events indicator, yes/no/unknown/not stated/inadequately

described code N equals 'yes'.

Adolescent and young adult cancer (clinical) NBPDS

Health, Standard 14/05/2015

Conditional obligation:

Complete if Person with cancer—family history of hereditary genetic events indicator, yes/no/unknown/not stated/inadequately described code N equals 'yes'.