National Healthcare Agreement: PI 11-Cervical screening rates, 2011

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National Healthcare Agreement: Pl 11-Cervical screening rates, 2011

Identifying and definitional attributes

Metadata item type: Indicator

Indicator type: Output measure

Short name: PI 11-Cervical screening rates, 2011

METEOR identifier: 421674

Registration status: Health, Superseded 30/10/2011

Description: Rates of cervical screening for women within national target age group

Indicator set: National Healthcare Agreement (2011)

Health, Superseded 31/10/2011

Outcome area: Prevention

Health, Standard 07/07/2010

Data quality statement: National Healthcare Agreement: PI 11-Cervical screening rates, 2011 QS

Health, Superseded 04/12/2012

Collection and usage attributes

Population group age

from:

Women aged 20 years

Population group age to: Women aged 69 years

Computation description: Denominator is adjusted to exclude the estimated number of women who have had

a hysterectomy, using national hysterectomy fractions.

Rates directly age-standardised.

Analysis by remoteness and SEIFA Indicator of Relative Socio-economic Disadvantage (IRSD) is based on postcode of residential address at the time of

screening.

Presented as a percentage.

Computation: 100 × (Numerator ÷ Denominator)

Numerator: Number of women aged 20–69 years who have been screened in a 2 year period.

Numerator data elements:

Data Element / Data Set

Data Element

Women who have undergone cervical cancer screening in a 2 year period

Data Source

ABS 2004-05 National Aboriginal and Torres Strait Islander Health Survey (NATSIHS)

Guide for use

Data source type: Survey For Indigenous women only

Data Element / Data Set-

Data Element

Person—age

Data Source

National Cervical Screening Program

Guide for use

Data source type: Registry

Data Element / Data Set

Data Element

Person—person identifier

Data Source

National Cervical Screening Program

Guide for use

Data source type: Registry

Data Element / Data Set-

Person—age, total years N[NN]

Data Source

ABS 2004-05 National Aboriginal and Torres Strait Islander Health Survey (NATSIHS)

Guide for use

Data source type: Survey For Indigenous women only

Denominator:

Population of women aged 20-69 years

Denominator data elements:

Data Element / Data Set-

Data Element

Person—estimated resident population of Australia

Data Source

ABS Estimated resident population (total population)

Guide for use

Data source type: Census-based plus administrative by-product data

Data Element / Data Set

Data Element

Proportion of women within the target population who have had a hysterectomy

Data Source

ABS 2001 National Health Survey (NHS)

Guide for use

Data source type: Survey

-Data Element / Data Set-

Person—age, total years N[NN]

Data Source

ABS 2004-05 National Aboriginal and Torres Strait Islander Health Survey (NATSIHS)

Guide for use

Data source type: Survey For Indigenous women only

Data Element / Data Set

Person—age, total years N[NN]

Data Source

ABS 2001 National Health Survey (NHS)

Guide for use

Data source type: Survey

Used for hysterectomy fractions

Disaggregation:

2008–2009—State and territory, by:

- remoteness
- SEIFA Index of Relative Socioeconomic Disadvantage (IRSD) quintiles

2008–2009—Nationally, by:

SEIFA IRSD deciles

State and territory, by Indigenous status (no data)

Disaggregations within individual jurisdictions are subject to data quality considerations. Some disaggregations may result in numbers too small for publication.

Disaggregation data elements:

Data Element / Data Set

Data Element

Establishment—Australian state/territory identifier

Data Source

National Cervical Screening Program

Guide for use

Data source type: Administrative by-product data Used for disaggregation by state and territory

Data Element / Data Set-

Data Element

Person (address)—Australian postcode

Data Source

National Cervical Screening Program

Guide for use

Data source type: Administrative by-product data Used for disaggregation by remoteness and SEIFA IRSD

Comments:

Hysterectomy fractions are derived from the 2001 National Health Survey, and were

recently validated using hospitals data.

Screening rates for Indigenous women in 2011 reporting were based on the ABS

2004-05 NATSIHS.

Most recent data available for 2011 CRC report: 2008–2009 (calendar years).

Representational attributes

Representation class: Percentage

Data type: Real
Unit of measure: Person
Format: NN.N

Indicator conceptual framework

Framework and dimensions:

Accessibility

Health behaviours

Data source attributes

Data sources:

Data Source

ABS Estimated resident population (total population)

Frequency

Quarterly

Data quality statement

ABS Estimated resident population (total population), QS

Data custodian

Australian Bureau of Statistics

Data Source

National Cervical Screening Program

Frequency

Annual

Data custodian

Department of Health and Ageing

Data Source

ABS 2001 National Health Survey (NHS)

Frequency

Every three years

Data custodian

Australian Bureau of Statistics

Data Source

ABS 2004-05 National Aboriginal and Torres Strait Islander Health Survey (NATSIHS)

Frequency

Every 6 years

Data custodian

Australian Bureau of Statistics

Accountability attributes

Reporting requirements: National Healthcare Agreement

Organisation responsible

for providing data:

Australian Institute of Health and Welfare

Further data development / Specification: Long-term

collection required:

Relational attributes

Related metadata references:

Supersedes National Healthcare Agreement: P11-Cervical screening rates, 2010 Health, Superseded 08/06/2011

Has been superseded by <u>National Healthcare Agreement: PI 11-Cervical screening rates</u>, 2012

Health, Retired 25/06/2013

See also National Healthcare Agreement: PI 04-Incidence of selected cancers, 2011

Health, Superseded 30/10/2011

See also <u>National Healthcare Agreement: PI 10-Breast cancer screening rates, 2011</u>

Health, Superseded 30/10/2011

See also <u>National Healthcare Agreement: PI 12-Bowel cancer screening rates, 2011</u>

Health, Superseded 30/10/2011

See also National Healthcare Agreement: PI 44-Survival of people diagnosed with cancer, 2011

Health, Superseded 31/10/2011