

# **Person with cancer—family history of hereditary genetic events indicator, yes/no/unknown/not stated/inadequately described code N**

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# Person with cancer—family history of hereditary genetic events indicator, yes/no/unknown/not stated/inadequately described code N

## Identifying and definitional attributes

Metadata item type:	Data Element
Short name:	Hereditary genetic events family history indicator
METEOR identifier:	457014
Registration status:	<a href="#">Health</a> , Standard 04/02/2015
Definition:	An indicator of whether a person with cancer has any <a href="#">first degree relatives</a> or <a href="#">second degree relatives</a> who have a current or prior hereditary genetic event, as represented by a code.
Data Element Concept:	<a href="#">Person with cancer—family history of hereditary genetic events indicator</a>
Value Domain:	<a href="#">Yes/no/unknown/not stated/inadequately described code N</a>

## Value domain attributes

## Representational attributes

Representation class:	Code	
Data type:	Number	
Format:	N	
Maximum character length:	1	
	Value	Meaning
Permissible values:	1	Yes
	2	No
Supplementary values:	8	Unknown
	9	Not stated/inadequately described

## Source and reference attributes

Submitting organisation:	Australian Institute of Health and Welfare
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## Data element attributes

## Collection and usage attributes

**Guide for use:** Record whether a person has any first degree relatives or second degree relatives who have a current or prior hereditary genetic event.

This includes, but is not exclusive to:

- Peutz-Jehgers syndrome
- B-Raf mutation
- Undescended testes
- Von Recklinghausen syndrome
- Xeroderma pigmentosa
- Bloom syndrome
- Ataxia telangiectasia
- Beckwith-Wiedemann syndrome
- Down syndrome
- Klinefelter's syndrome
- Neurofibromatosis type I (NF-1)
- Von Hippel–Lindau disease (VHL)
- RET proto-oncogene
- Adenomatous polyposis coli (APC)
- Hereditary nonpolyposis colorectal cancer (HNPCC or Lynch syndrome)
- BRCA1
- BRCA2
- Neurofibromatosis
- Li-Fraumeni Syndrome
- Fanconi anaemia
- Other

**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—hereditary genetic events type, code N\[N\] 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 Specifications:** [Adolescent and young adult cancer \(clinical\) DSS Health](#), Superseded 14/05/2015

[Adolescent and young adult cancer \(clinical\) NBPDS Health](#), Standard 14/05/2015