

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:	Health , Standard 04/02/2015
Definition:	An indicator of whether a person with cancer has any first degree relatives or second degree relatives who have a current or prior hereditary genetic event, as represented by a code.

Data element concept attributes

Identifying and definitional attributes

Data element concept:	Person with cancer—family history of hereditary genetic events indicator
METEOR identifier:	564462
Registration status:	Health , Standard 04/02/2015
Definition:	An indicator of whether a person with cancer has any first degree relatives or second degree relatives who have a current or prior hereditary genetic event.
Object class:	Person with cancer
Property:	Family history of hereditary genetic events indicator

Source and reference attributes

Submitting organisation:	Cancer Australia
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Value domain attributes

Identifying and definitional attributes

Value domain:	Yes/no/unknown/not stated/inadequately described code N
METEOR identifier:	546669
Registration status:	Health , Standard 11/04/2014 Indigenous , Standard 05/12/2017 Australian Institute of Health and Welfare , Qualified 17/01/2024 Early Childhood , Recorded 16/05/2024
Definition:	A code set representing 'yes', 'no', 'unknown' and 'not stated/inadequately described' responses.

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

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

