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

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| Identifying and definitional attributes | |
| Metadata item type: | Data Element |
| Short name: | Hereditary genetic events family history indicator |
| METEOR identifier: | 457014 |
| Registration status: | [Health](https://meteor.aihw.gov.au/RegistrationAuthority/12), Standard 04/02/2015 |
| Definition: | An indicator of whether a person with cancer has any [**first degree relatives**](https://meteor.aihw.gov.au/content/494465) or  [**second degree relatives**](https://meteor.aihw.gov.au/content/494469) who have a current or prior hereditary genetic event, as represented by a code. |
| Data Element Concept: | [Person with cancer—family history of hereditary genetic events indicator](https://meteor.aihw.gov.au/content/564462) |
| Value Domain: | [Yes/no/unknown/not stated/inadequately described code N](https://meteor.aihw.gov.au/content/546669) |

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| 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 |

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| --- | --- |
| 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]](https://meteor.aihw.gov.au/content/467855)  [Health](https://meteor.aihw.gov.au/RegistrationAuthority/12), Standard 04/02/2015  See also [Person with cancer—hereditary genetic events type, text X[X(39)]](https://meteor.aihw.gov.au/content/467963)  [Health](https://meteor.aihw.gov.au/RegistrationAuthority/12), Standard 04/02/2015 |
| Implementation in Data Set Specifications: | [Adolescent and young adult cancer (clinical) DSS](https://meteor.aihw.gov.au/content/432097)  [Health](https://meteor.aihw.gov.au/RegistrationAuthority/12), Superseded 14/05/2015  [Adolescent and young adult cancer (clinical) NBPDS](https://meteor.aihw.gov.au/content/599629)  [Health](https://meteor.aihw.gov.au/RegistrationAuthority/12), Standard 14/05/2015 |