

# **Person with cancer—personal genetic syndrome indicator, yes/no/unknown/not stated/inadequately described code N**

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# Person with cancer—personal genetic syndrome indicator, yes/no/unknown/not stated/inadequately described code N

## Identifying and definitional attributes

Metadata item type:	Data Element
Short name:	Personal genetic syndrome indicator
METEOR identifier:	457065
Registration status:	<a href="#">Health</a> , Standard 04/02/2015
Definition:	An indicator of whether the person with cancer has been diagnosed with a <a href="#">genetic syndrome</a> , as represented by a code.
Data Element Concept:	<a href="#">Person with cancer—personal genetic syndrome indicator</a>
Value Domain:	<a href="#">Yes/no/unknown/not stated/inadequately described code N</a>

## Value domain attributes

## Representational attributes

<b>Representation class:</b>	Code	
<b>Data type:</b>	Number	
<b>Format:</b>	N	
<b>Maximum character length:</b>	1	
	<b>Value</b>	<b>Meaning</b>
<b>Permissible values:</b>	1	Yes
	2	No
<b>Supplementary values:</b>	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

<b>Guide for use:</b>	<p>Record yes when a person has been diagnosed with any genetic syndrome.</p> <p>This includes, but is not exclusive to:</p> <ul style="list-style-type: none"> <li>• Peutz-Jehgers syndrome</li> <li>• B-Raf mutation</li> <li>• Undescended testes</li> <li>• Von Recklinghausen syndrome</li> <li>• Xeroderma pigmentosa</li> <li>• Bloom syndrome</li> <li>• Ataxia telangiectasia</li> <li>• Beckwith-Wiedemann syndrome</li> <li>• Down syndrome</li> <li>• Klinefelter syndrome</li> <li>• Neurofibromatosis type I (NF-1)</li> <li>• Von Hippel–Lindau disease (VHL)</li> <li>• RET proto-oncogene</li> <li>• Adenomatous polyposis coli (APC)</li> <li>• Hereditary nonpolyposis colorectal cancer (HNPCC or Lynch syndrome)</li> <li>• Breast cancer gene 1 (BRCA1)</li> <li>• Breast cancer gene 2 (BRCA2)</li> <li>• Neurofibromatosis</li> <li>• Li-Fraumeni syndrome</li> <li>• Fanconi anaemia</li> </ul> <p>If Yes is selected for personal genetic syndrome indicator, record the type of specific syndrome/s at <a href="#">Person with cancer—hereditary genetic events type, code N[N]</a> or <a href="#">Person with cancer—hereditary genetic events type, text X[X(39)]</a>.</p>
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**Collection methods:** Collect from patient medical records.

## Source and reference attributes

**Submitting organisation:** Cancer Australia

## Relational attributes

<b>Related metadata references:</b>	<p>See also <a href="#">Person with cancer—personal genetic syndrome type, genetic event type code N[N]</a>  <a href="#">Health</a>, Standard 04/02/2015</p> <p>See also <a href="#">Person with cancer—personal genetic syndrome type, text X[X(39)]</a>  <a href="#">Health</a>, Standard 04/02/2015</p>
<b>Implementation in Data Set Specifications:</b>	<p><a href="#">Adolescent and young adult cancer (clinical) DSS</a>  <a href="#">Health</a>, Superseded 14/05/2015</p> <p><a href="#">Adolescent and young adult cancer (clinical) NBPDS</a>  <a href="#">Health</a>, Standard 14/05/2015</p>