

# Health service event—adverse event grade, code N

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# Health service event—adverse event grade, code N

## Identifying and definitional attributes

<b>Metadata item type:</b>	Data Element
<b>Short name:</b>	Adverse event grade
<b>METEOR identifier:</b>	467449
<b>Registration status:</b>	<a href="#">Health!</a> , Standard 04/02/2015
<b>Definition:</b>	The grade of an <a href="#">adverse event</a> that has occurred during a health service event as defined by the American National Institute of Health and National Cancer Institute's Common Terminology Criteria for Adverse Events (CTCAE, v4.0), as represented by a code.
<b>Data Element Concept:</b>	<a href="#">Health service event—adverse event grade</a>
<b>Value Domain:</b>	<a href="#">Adverse event grade 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	Grade 1
	2	Grade 2
	3	Grade 3
	4	Grade 4
	5	Grade 5
<b>Supplementary values:</b>	7	Not applicable, no adverse events
	8	Unknown
	9	Adverse event, but grade inadequately specified

## Collection and usage attributes

<b>Guide for use:</b>	Grade refers to the severity of the adverse event (AE). The CTCAE displays Grade 1 through to Grade 5 with unique clinical descriptions of severity for each AE based on this general guideline:  CODE 1 Grade 1  Mild; asymptomatic or mild symptoms; clinical or diagnostic observations only intervention not indicated.  CODE 2 Grade 2
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## Source and reference attributes

<b>Submitting organisation:</b>	Cancer Australia
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**Reference documents:** National Cancer Institute 2009. Common Terminology Criteria for Adverse Events (CTCAE) Version 4.0. Viewed 7/11/2011.  
[http://evs.nci.nih.gov/ftp1/CTCAE/CTCAE\\_4.03\\_2010-06-14\\_QuickReference\\_5x7.pdf](http://evs.nci.nih.gov/ftp1/CTCAE/CTCAE_4.03_2010-06-14_QuickReference_5x7.pdf)

## Data element attributes

### Collection and usage attributes

**Guide for use:** Record the grade for any adverse event that has occurred during a health service event.

An **Adverse Event** (AE) is any unfavourable and unintended sign (including an abnormal laboratory finding), symptom, or disease temporally associated with the use of a medical treatment or procedure that may or may not be considered related to the medical treatment or procedure.

**Collection methods:** Collect from patient medical records.

### Source and reference attributes

**Submitting organisation:** Cancer Australia

### Relational attributes

**Related metadata references:** See also [Patient—adverse event indicator, yes/no code N Health!](#), Standard 29/08/2014

See also [Patient—date of adverse event, DDMMYYYY Health!](#), Standard 04/02/2015

**Implementation in Data Set Specifications:** [Adolescent and young adult cancer \(clinical\) DSS Health!](#), Superseded 14/05/2015

**Conditional obligation:** Complete if [Patient—adverse event indicator, yes/no code N](#) equals 'yes'.

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

**Conditional obligation:**

Complete if [Patient—adverse event indicator, yes/no code N](#) equals 'yes'.