

Procedure Report – Adverse Events



This report provides valuable information to the National Cancer Screening Register (Register) about your patient, where they are a National Bowel Cancer Screening Program (Program) participant. Your assistance is sought to ensure Program information is complete.

When to use this report

This report is to provide information to the Register where your patient is a Program participant and there has been an adverse outcome in relation to a Colonoscopy, Double Contrast Barium Enema, Flexible Sigmoidoscopy or CT Colonography, or any other relevant procedure relating to diagnostic investigation.

Instructions for using this report

Please use a black pen and write in BLOCK LETTERS in the boxes provided.

Mandatory fields are marked with an asterisk (*).

Preferred fields are marked with a plus (+).

How to lodge the report

The original copy of the report can be lodged with the Register:

via free fax to 1800 115 062; or

post to National Bowel Cancer Screening Program, Reply Paid 90965, Sunshine, VIC 3020

More information

More information about this report can be obtained by contacting the National Bowel Cancer Screening Program Contact Centre on 1800 118 868.

Participant privacy

NBCSP Participant Privacy

In accordance with the relevant requirements of the *Privacy Act 1988 (Cth)*, patients are made aware that healthcare providers may collect and disclose their personal information to the NCSR. You are authorised to collect and disclose your patient's personal information under the *National Cancer Screening Register Act 2016*.

NBCSP Practitioner Privacy

The NCSR is authorised to collect information under the *Privacy Act 1988 (Cth)* and the *National Cancer Screening Register Act 2016.* The NCSR collects information about you and other healthcare providers from the Department of Human Services and others for the purpose of verifying your identity and communicating with you.

The NCSR also collects information directly from you. Your personal information may be disclosed to a range of agencies or organisations, including State and Territory Health Departments, Australian Government agencies and where you have agreed or where it is authorised or required by law or court or tribunal order.

If you require information on the NCSR's privacy policy, please visit www.ncsr.gov.au



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- 1. Mandatory fields are marked with an asterisk (*).
- 2. Preferred fields are marked with a plus (+).

Patient Details																			
Participant ID number									ledica umbe		DVA								
*Family name																			
*Given name																			
*Date of birth (dd/mm/yyyy)		/		/				4	'Gen	der	Ma	ale		Fe	male		(Othe	r
Does the patient ide	ntify a	as Al	orio	gina	ıl or	Tor	res S	Strait	Islaı	nde	r orig	in?	(if kr	now	n)				
Aboriginal Torres Strait Islander			Aboriginal and Torres Strait Is							on digenous			Prefer not to answer				r		
What is the patient's	coun	ntry c	f or	igin	? (ii	f kn	own)												
What is the patient's	; prefe	erred	lan	gua	ge s	spol	ken a	t hor	ne?	(if k	nowr	1)							
Adverse Outcomes Bleeding Infection/ sepsis			Perforation				Reaction to sedation							Death					
Other (p	olease s	specify)																
Delayed discharge	? 1	No		Υe	es						hospi				No			Ye	s
Surgery required	l? 1	No		Υe	es														
Provider details																			
Facility/Hospital provider number																			
*Name of Facility /																			
Hospital				T							nber is			. If c	linicia	n/nr	-000	dura	
Hospital								doe	s not	nav	e a pr	ovide	er nur	nbe				ndat	OI y
Hospital *Clinician/Proceduralist								doe	s not	nav	e a pr	ovide	er nur	nbe				ndat	Ory
*Clinician/Proceduralist provider number Name of Clinician/Proceduralist *Date of procedure		1		/				doe	s not	nav	e a pr	ovide	er nur	nbe				ndat	Ory
*Clinician/Proceduralist provider number Name of Clinician/			ultina	l l	der)			doe	s not	nav	e a pr	ovide	er nur	nbe				ndat	Ory