



# National Cardiac Rehabilitation Quality Indicators: Data Dictionary

Version 10.0

June 2021





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- National Heart Foundation of Australia, NSW Agency for Clinical Innovation (ACI) and Cardiovascular Health and Rehabilitation Association NSW/ACT. NSW cardiac rehabilitation quality indicators data dictionary and definition guide 2017.
- International Council of Cardiovascular Prevention and Rehabilitation (ICCPR). International Cardiac Rehabilitation Registry. Data elements definitions 2.0. 2020
- AuSCR Data Dictionary. Version 6.0. 2019 Available online: <u>https://auscr.com.au/health-professionals/variables-program-bundles/</u>

The National Cardiac Rehabilitation Quality Indicators were developed by a Taskforce group originally comprising the following members in October 2019:

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## Preamble

Care *quality* is an increasing focus of funders, providers and consumers of healthcare. Measuring quality and the clinical effectiveness of health services is important for ensuring accountability of healthcare providers, enhancing patient outcomes, minimising adverse events and aligning care with what patients want and the best available evidence.

*Quality indicators* are explicitly defined statements that aim to measure adherence to aspects of evidence-based care that are deemed necessary for reaching optimal patient outcomes and provide a basis for quality improvement projects.<sup>1,2</sup>

The need for Australian cardiac rehabilitation (CR) quality indicators was determined at a Think Tank on improving CR measurement, which was held on 26 September 2018 at the South Australian Advanced Health Research and Translation Centre and was attended by researchers, clinicians, policymakers and consumers with representation from each state and territory. The aim of the Think Tank was to discuss state-based CR quality activities and future national directions and it was agreed that a national set of quality indicators for CR service measurement was required.

A Taskforce, co-chaired by the National Heart Foundation of Australia and the Australian Cardiovascular Health and Rehabilitation Association (ACRA), was established to progress the development of the quality indicators. The purpose of the quality indicators is to set recommendations for what should be collected and reported on at a minimum so that CR programs can collect uniform data.

The Taskforce developed a draft set of 11 quality indicators and disseminated these to ACRA members for feedback, via email and at the 2019 ACRA Annual Scientific Meeting, on their perceived importance to: (i) clinicians, (ii) managers, and (iii) patients, as well as the feasibility of collecting the indicators. Based on feedback, one indicator (waist circumference) was removed. This process has been published in the following editorial:

• Gallagher R, Thomas E, Astley C, Foreman R, Ferry C, Zecchin R, Woodruffe S. Cardiac Rehabilitation Quality in Australia: Proposed National Indicators for Field-Testing. Heart, Lung and Circulation. 2020 Epub Apr 30.

This document provides the proposed 10 quality indicators developed by the Taskforce along with a description of how the indicators can be defined and measured. To collect this information a number of data elements need to be collected at two time points – pre and post-CR.

<sup>&</sup>lt;sup>1</sup> Spertus JA, Eagle KA, Krumholz HM, Mitchell KR, Normand SL and American College of Cardiology/American Heart Association Task Force on Performance M. American College of Cardiology and American Heart Association methodology for the selection and creation of performance measures for quantifying the quality of cardiovascular care. *J Am Coll Cardiol.* 2005; 45: 1147-56.

<sup>&</sup>lt;sup>2</sup> Campbell SM, Braspenning J, Hutchinson A and Marshall M. Research methods used in developing and applying quality indicators in primary care. Quality & Safety in Health Care. 2002; 11: 358-64.





The purpose of this document is to guide CR services in the type of information they should collect on every patient referred to their service. We provide a standardised format for data collection to ensure information is collected consistently across sites and provides a basis for comparison and reporting across sites. The purpose of the national CR quality indicators is not to duplicate effort but to ensure a) services that do not collect data have guidance on what to collect, b) services that are collecting data are doing so in a standardised way and c) services go beyond collecting data to measuring care quality against the quality indicators described within this document.

The National Cardiac Rehabilitation Measurement Taskforce is NOT collecting or collating this information. Services are required to collect and securely save their own data. Each state is at different levels of organisational readiness regarding how they collect and collate this information. The Taskforce will be evaluating the feasibility, acceptability, and usefulness of the quality indicators in the future.





### Using the data dictionary

This data dictionary is divided into 2 sections:

- Section A: the data elements
  - To calculate the quality indicators, each service needs to collect several data elements for each eligible CR patient at two time points: Pre and post the CR program
  - To assist with standardised data collection, each data element contains the following information: reference number, common name, description, codes and values, help notes.
- Section B: the quality indicators (with reference to Section A as needed)
  - Once the data elements have been collected, they can be used to determine the quality indicators over a determined reference period (the recommended minimum period of assessment is annually).

The following terms are frequently used throughout the document and are defined below.

Term	Definition
CR program	A CR program that provides exercise training, health behaviour change and education regarding lifestyle risk factor management, psychosocial health, medical risk factor management, and cardio-protective medications assessment and monitoring.
Eligibility	<ul> <li>Eligible patients for the purpose of measuring the quality indicators include patients hospitalised with any of the following procedures/conditions or interventions listed below:</li> <li>Acute myocardial infarction (STEMI, non-STEMI) (see Appendix A for ICD codes)</li> <li>Percutaneous coronary intervention (see Appendix A for ICD codes)</li> <li>Coronary artery bypass surgery (see Appendix A for ICD codes)</li> <li>Valve surgery (see Appendix A for ICD codes)</li> <li>Atrial fibrillation, heart failure, unstable angina, arrhythmia, angina pectoralis, congenital heart disease, automatic implantable cardioverter defibrillator</li> </ul>
Enrolment	Enrolment is defined as patient registration into the CR program and attendance at the first CR program visit (including an assessment session).
Comprehensive assessment	A comprehensive assessment is defined as the assessment of exercise capacity, lifestyle risk factors (smoking, diet, physical activity), psychosocial health status, clinical risk factors (e.g. blood pressure) and use of cardio- protective medication. Additionally, it should include assessment of the patient's secondary prevention needs.

Table 1. Common terms





	The assessment is finalised when the discussion and agreement of individualised goals to be achieved during CR has been completed, with a written self-management plan and a copy made available to the patient and/or family.
Completion	To complete the CR program a patient must have participated in at least
	some of the CR intervention components (guided by a health professional)
	and have had a documented re-assessment.
Health related	HRQoL is a patient reported outcome measure that takes into account the
quality of life	patient's emotional, social, and physical well-being.
(HRQoL)	
Individualised	An individualised management plan is designed to promote the safe and
management	timely transition of patients between different health care providers and
plan	across care settings.
Reference period	Reference period is defined as the time span of data collection that is
	included in analysis. The recommended minimum period of assessment is
	annually.
Referred	A referral is defined as an official written or electronic communication on
	behalf of any health care provider (including CR staff) to CR.





## Summary – Data elements (Section A)

Overview of data elements			
Pre-program	Pre-program		
Identifiable information	Referral information		
<ol> <li>Patient record ID</li> <li>First name</li> <li>Last name</li> <li>Hospital medical record number</li> <li>Medicare number</li> <li>Date of birth</li> </ol> Socio-demographic information <ol> <li>Age</li> <li>Sex Recorded at Birth</li> <li>Aboriginal and Torres Strait Islander status</li> <li>Interpreter needed</li> <li>Postcode</li> </ol>	<ul> <li>12. Hospital discharge date</li> <li>13. CR referral date</li> <li>14. Principal referral diagnosis</li> <li><i>Initial assessment</i></li> <li>15. Initial assessment date</li> <li>16. Depression screening</li> <li>17. Depression referral</li> <li>18. Smoking status</li> <li>19. Smoking referral</li> <li>20. Medication adherence</li> <li>21. Exercise capacity</li> <li>22. Health-related quality of life</li> </ul>		
Post-program			
Re-assessment Service delivery information			
<ul> <li>23. Re-assessment date</li> <li>24. Re-assessment depression screening</li> <li>25. Re-assessment depression referral</li> <li>26. Re-assessment smoking status</li> <li>27. Re-assessment smoking referral</li> <li>28. Re-assessment medication adherence</li> <li>29. Re-assessment of exercise capacity</li> <li>30. Change in exercise capacity</li> <li>31. Re-assessment of health-related quality of life</li> </ul>	<ol> <li>34. Mode of program delivery</li> <li>35. Frequency of program delivery</li> <li>36. Content of program</li> <li>37. Number of supervised exercise sessions attended</li> <li>38. CR completion</li> <li>39. Reason for CR non-completion</li> </ol>		
<ol> <li>Change in health-related quality of life</li> <li>Care transition</li> </ol>			





# Summary - Australian Cardiac Rehabilitation Quality Indicators (Section B)

The table below provides a summary of the 10 quality indicators for CR. Some indicators aim to evaluate processes of care (process indicators,) while others evaluate the outcomes of CR (outcome indicators). These are colour co-ordinated as per the key below the figure.

#### Overview of cardiac rehabilitation quality indicators

#### QI-1. Referral to CR

Eligible in-patients are referred to cardiac rehabilitation within 3 calendar days after hospital discharge.

#### QI-2. Time to enrolment

Eligible in-patients commence cardiac rehabilitation within 28 calendar days after hospital discharge.

QI-3. Comprehensive assessment

Patients who commence cardiac rehabilitation receive a comprehensive assessment of cardiovascular risk factors.

#### QI-4. Depression screening

Patients who commence cardiac rehabilitation are screened for depression at initial and re-assessment and offered counselling (or referral to counselling) if symptoms are identified.

QI-5. Assessment of smoking

Patients who commence cardiac rehabilitation are assessed for smoking use at initial and re-assessment and offered smoking cessation counselling (or referral to counselling) if they are a current or recent smoker.

QI-6. Assessment of medication adherence

Patients who commence cardiac rehabilitation are assessed for medication adherence at initial and reassessment.

#### QI-7. Assessment of exercise capacity

Patients who commence cardiac rehabilitation have an initial assessment and re-assessment to determine exercise capacity change.

#### QI-8. Assessment of health-related quality of life

Patients who commence cardiac rehabilitation have an initial assessment and re-assessment to determine health-related quality of life change.

#### QI-9. Re-assessment

Patients who participate in cardiac rehabilitation receive a comprehensive re-assessment of their cardiovascular risk factors.

#### QI-10. Care transition

Patients and ongoing care providers are provided with a report which outlines patient risk factors and an individualised ongoing management plan.

Process indicator

Outcome indicator





## **SECTION A**





### PRE-PROGRAM

The following information should be collected for each eligible CR patient and stored securely by each service according to institutional protocols and should not be shared beyond the service-level unless clear data sharing approvals and governance structures are in place.

#### IDENTIFIABLE INFORMATION

Reference number	1
Description	This number represents a unique identifier that indicates which clinical record this patient relates to but does not identify the patient's identity.
Codes and values	Free text
Help notes	Services should ensure patient record IDs are provided in a systematic and consistent way. Each patient record ID is unique for each episode of care. This number is useful to identify records whilst observing confidentiality of patient information. This should not be the patients MRN or URN.

#### Patient record ID

#### First name

Reference number	2
Description	The person's identifying name within the family group or by which the person is
	socially identified, as represented by text.
Codes and values	Free text
Help notes	The format in which it is written should be the same as that indicated by the person
	(e.g. written on a form) or in the same format as that printed on an identification
	card, such as a Medicare card, to ensure consistent collection of name data.

#### Last name

Reference number	3
Description	That part of a name a person usually has in common with some other members of his/her family, as distinguished from his/her first or given names, as represented by text.
Codes and values	Free text
Help notes	The full family name should be recorded. The format in which it is written should be the same as that printed on an identification card, such as a Medicare card, to ensure consistent collection of name data.

#### Hospital medical record number (MRN)

Reference number	4
Description	Person identifier unique within an establishment or agency also known as Unit
	Record Number (UR) and Patient Record Number.
Codes and values	Free text.
Help notes	The MRN is collected to assist in individual patient identification and to identify
	potential duplicates in the database. It is useful for linking the CR patient with their
	hospital record to determine the number of eligible patients that were referred to
	CR.





#### Medicare number (once data-linked)

Reference number	5
Description	Number on the person's Medicare Card used as an Australian Commonwealth
	Government identifier.
Codes and values	Free text
Help notes	The full Medicare number for an individual should be recorded, without including
	the person (individual reference) number. For example, John Smith's full Medicare
	number is 1234567890
	Medicare
	1234 56789 0
	1 JOHN SMITH 2 HELEN SMITH 3 JAMES SMITH 4 JESSICA SMITH
	VAUD TO 11/10
	For overseas visitors or patients without Medicare Card please leave blank

#### Date of birth

Reference number	6
Description	The date of birth of the person.
Codes and values	DD/MM/YYYY
Help notes	If day of birth is unknown, use 01 for the day (01/MM/YYYY). If the day and month of birth are unknown, use 01 for the day and month (01/01/YYYY).

#### SOCIO-DEMOGRAPHIC INFORMATION

#### Age

5	
Reference number	7
Description	The age in completed years at the time of CR referral.
Codes and values	DD/MM/YYYY
Help notes	Subtract current date from DOB

#### Sex Recorded at Birth

Reference number	8
Description	The sex recorded at birth (male, female, non - binary) that the person identifies as.
Codes and values	1 Male
	2 Female
	3 Non-binary sex – people who reported their sex as non – binary sex
	99 Not stated / inadequately described
Help notes	Sex recorded at birth should be captured as it is written in the medical record. If
	there is a conflict, document with the self-identified sex recorded at birth, i.e. sex
	recorded at birth as reported by the person.

#### Aboriginal and Torres Strait Islander status

Reference number	9
Description	Whether a person identifies as being of Australian Indigenous, Aboriginal or Torres
	Strait Islander origin.
Codes and values	1 No
	2 Yes Aboriginal origin
	3 Yes Torres Strait Islander origin





	4 Yes both Aboriginal and Torres Strait Islander origin
	5 Missing/not stated
Help notes	Aboriginal or Torres Strait Islander origin status should be captured as it is written
	in the medical record. If there is a conflict, document with the self-identified origin,
	i.e. origin as reported by the person.

#### Interpreter needed

Reference number	10
Description	Whether an interpreter service is required for the patient to participate in CR.
Codes and values	1. Yes
	2 No
Help notes	Includes whether an approved interpreter service is required. Any form of sign
	language should also be coded as 'Yes'.

#### Post code

Reference number	11
Description	The numeric descriptor for a postal delivery area, aligned with locality, suburb or place for the address of a person. METeOR Identifier: 611398
Codes and values	Free text
Help notes	Leave blank when the locality name or geographic area for a person is not known, or when a person has no fixed address. For person's visiting from overseas, record their local Australian address.

#### **REFERRAL INFORMATION**

Please note that all referred patients' information should be entered (not just those who attend the initial assessment) so that the proportion of referred patients enrolling can be assessed.

#### Hospital discharge date

Reference number	12
Description	The date the patient was discharged from an acute episode of care.
Codes and values	DD/MM/YYYY
Help notes	The patient may have several inpatient separations during a single acute episode of care (i.e. short stay unit to ward to ICU to ward). The final date of discharge from the acute episode of care should be used. May source from patient's medical record, CR referral log.

#### CR referral date

Reference number	13
Description	Date CR referral document was signed by referring healthcare provider
Codes and values	DD/MM/YYYY)
Help notes	May source from patient's medical record, CR referral log, GP letter

#### Principal referral diagnosis/Interventions

Reference number	14a
Description	The referral diagnosis refers to the most recent diagnosis preceding the patient's
	referral to cardiac rehabilitation.
Codes and values	Select from below
	1. STEMI
	2. NSTEMI
	3. CABG





	4. Valve Sx
	5. Elective PCI
	6. Heart Failure (HFrEF, HFpEF)
	7. Arrhythmia
	8. AICD/PPM
	9. Angina Pectoralis
	10. Congenital Heart Disease
	11. Congenital Heart Sx
	12. TAVI
	13. Other
Help notes	Please use ICD codes to guide selection (See Appendix A). There may be more than
	one possible referral diagnosis reported if the second occurred within the same
	hospitalization period. DO NOT report historical diagnoses/interventions. May
	source from patient's medical record, CR referral log. AICD = Automatic implantable
	cardioverter defibrillator; ICD = International Classification of Disease
	PPM=permanent pacemaker
Reference number	14b
Description	Additional diagnosis/interventions
Codes and values	Select from below
	1. CABG Sx
	2. Valve Sx
	3. Primary PCI
	4. Staged PCI
	5. Elective PCI
	6. Heart Failure (HFrEF, HFpEF)
	7. Arrhythmia
	8. AICD/PPM
	9. Congenital Heart Sx
	10. TAVI
	11. Other
	12. Medical Management
Help notes	There may be more than one possible referral diagnosis reported if the second
	occurred within the same hospitalization period. DO NOT report historical
	diagnoses/interventions. May source from patient's medical record, CR referral log.
Reference number	14c
Description	Additional diagnosis/interventions
Codes and values	Select from below
	1. CABG Sx
	2. Valve Sx
	3. Primary PCI
	4. Staged PCI
	5. Elective PCI
	6. Heart Failure (HFrEF, HFpEF)
	7. Arrhythmia
	8. AICD/PPM
	9. Congenital Heart Sx
	10. TAVI
	11. Other
	12. Medical Management
Help notes	There may be more than one possible referral diagnosis reported if the second
	occurred within the same hospitalization period. DO NOT report historical
	diagnoses/interventions. May source from patient's medical record, CR referral log.



#### INITIAL ASSESSMENT



#### Initial assessment date

Reference number	15
Description	Enter the date of the initial assessment took place i.e. the date the patient had
	their initial visit at CR for assessment of risk, history, etc.
Codes and values	DD/MM/YYYY
Help notes	May source from CR administrative database, initial assessment information.

#### Depression screening

Reference number	16
Description	Was the patient screened for depression using a valid and reliable screening tool?
Codes and values	0 Yes
	1 No
	If no provide reason for not screening [open text].
	If yes, please document the depression scores and the depression screening tool
	used.
Help notes	May source from patient's medical record, patient initial assessment.
	Suggested tools include (but are not limited to) the following: the Patient Health
	Questionnaire (PHQ) -2 and PHQ-9, the Hospital Anxiety and Depression Screener
	(HADS), the Cardiac Depression Screener (CDS), the Beck Depression Inventory
	(BDI-II), and the Kessler Psychological Distress Scale (Kessler-10).

#### Depression referral

Reference number	17	
Description	Patients who screen positive for depression were offered/referred for counselling	
Codes and values	0 Yes	
	1 No	
	99 N/A	
If no select reason for not referring for further management [optional]		
	Declined	
	Under current treatment	
	Unknown	
	Other	
Help notes	May source from patient's medical record (i.e. documentation of referral or	
	counselling support provided).	

#### Smoking status

Reference number	18		
Description	Was the CR patient's smoking status assessed?		
Codes and values	0 Yes		
	1 No		
	99 Unknown		
	If no provide reason for not screening.		
	If yes, document smoking status		
	1 Daily smoker (A person who smokes daily)		
	2 Weekly smoker (A person who smokes at least weekly but not daily)		
	3 Irregular smoker (A person who smokes less than weekly)		
	4 Ex-smoker (A person who does not smoke at all now, but has smoked at least		
	100 cigarettes or similar amount of other tobacco products in his/her lifetime)		





	5 Never-smoker (A person who does not smoke now and has smoked fewer than
	100 cigarettes or similar amount of other tobacco products in his/her lifetime
	99 Unknown
Help notes	May source from patient's medical record, patient initial assessment, self-report.

#### Smoking referral

Reference number	19	
Description	Patients who are current or recent smokers were offered smoking cessation	
	counselling (or were referred to counselling).	
Codes and values	0 Yes	
	1 No	
99 Unknown		
If no, select reason for not offering smoking cessation counselling (or a refe		
	counselling)	
	N/A	
	Declined	
	Under current treatment	
	Unknown	
Help notes	May source from patient's medical record (i.e. documentation of referral or	
	smoking cessation support provided).	

#### Medication adherence

Reference number	20
Description	Medication adherence was assessed?
Codes and values	0 Yes 1 No 99 Unknown
	Is the patient adhering to prescribed treatment O Yes 1 No 3 Unknown
	Reason for non-adherence Cost Cognitive issues Side effects/Contraindication Patient choice Unknown
Help notes	<ul> <li>May source from patient's medical record, patient initial assessment, self-report.</li> <li>Suggested tools include the Brief Adherence Self-Report Questionnaire (Zeller, Schroeder et al., 2008); the Voils Measure of Extent and Reasons for Medication Non-Adherence (Voils et al., 2012); Brief Medication Questionnaire (Svarstad et al., 1999) or self-report response to medication adherence questions such as:</li> <li>Are you taking your medications as instructed? (Y/N)</li> <li>If no, reason why (e.g. forgot, cost, side effects, other)</li> <li>What would you do if you missed your regular dose? (Take dose later in the day, call GP or local pharmacist, miss the dose altogether, other)</li> </ul>





#### Exercise capacity

Reference number	21		
Description	Functional exercise capacity was assessed		
Codes and values	0 No		
	1 Yes		
	99 Unknown		
	If Yes – please add number of metres achieved for 6MWT or METs achieved for EST		
	or according to questionnaire results		
Help notes	May source from patient's medical record, patient initial assessment. Suggested		
	tools depending on local CR program requirements/capabilities include:		
	The Six Minute Walk Test (6MWT)		
	Distance walked within 6 minutes can be measured using standardised 6MWT		
	protocols.		
	The Exercise Stress Test (EST)		
	The METs can be estimated from standard equations using speed and grade, or can		
	be calculated from the direct measurement of oxygen consumption using gas		
	analysis.		
	Self-reported exercise capacity		
	The Specific Activity Questionnaire (SAQ) is a 13-item self-report measure of		
	exercise capacity. The scoring method of the tool can be used to estimate METS.		
	The SAQ is publicly available and free of charge and has been validated against		
	exercise stress testing in cardiac patients		

#### Health-related quality of life

Reference number	22	
Description	Health-related quality of life was assessed using a validated tool.	
Codes and values	Yes	
	No	
	Unknown	
	Score for each domain as per validated tool used	
Help notes	May source from patient's medical record, patient initial assessment. Suggested	
	tools include: freely available options: <u>Assessment of Quality of Life</u> (AQOL),	
	EuroQol(EQ)-5D-L and pay-for options: Short Form (SF) -12 and SF-36.	

### POST PROGRAM

#### **RE-ASSESSMENT**

#### Re-assessment date

Reference number	23	
Description	Enter the date the re-assessment took place i.e. the date the patient had their final	
	visit at CR and/or were re-assessed.	
Codes and values	DD/MM/YYYY	
Help notes	May source from CR administrative database, re-assessment information.	

#### Re-assessment depression screening

	•	
Reference number	24	





Description	Was the patient re-screened for depression using a valid and reliable screening	
	tool?	
Codes and values	0 Yes	
	1 No	
	99 N/A	
	If yes, please document the depression scores and the depression screening tool	
	used	
	If no select reason for not screening	
Help notes	May source from patient's medical record, patient re-assessment. Use the same	
	tool as used in initial assessment.	

#### Re-assessment depression referral

Reference number	25	
Description	Patients who screen positive for depression were offered/referred for counselling	
Codes and values	0 Yes	
	1 No	
	99 N/A	
If no select reason for not referring for further management		
Declined		
	Under current treatment	
	Other	
	Unknown	
Help notes	May source from patient's medical record (i.e. documentation of referral or	
	counselling support provided).	

#### *Re-assessment smoking status*

Reference number	26		
Description	Was the CR patient's smoking status re-assessed?		
Codes and values	0 Yes		
	1 No		
	99 Unknown		
	If yes, document smoking status		
	1 Daily smoker (A person who smokes daily)		
	2 Weekly smoker (A person who smokes at least weekly but not daily)		
	3 Irregular smoker (A person who smokes less than weekly)		
	4 Ex-smoker (A person who does not smoke at all now, but has smoked at least		
	100 cigarettes or similar amount of other tobacco products in his/her lifetime)		
	5 Never-smoker (A person who does not smoke now and has smoked fewer than		
	100 cigarettes or similar amount of other tobacco products in his/her lifetime		
	99 Unknown		
Help notes	May source from patient's medical record, patient re-assessment, self-report.		

#### Re-assessment smoking referral

Reference number	27
Description	Patients who are current or recent smokers were offered smoking cessation
Codes and values	0 Yes
	1 No
	99 NA
	If no, select reason for not offering smoking cessation counselling (or a referral to
	counselling)
	Declined





	Under current treatment
	Other
	Unknown
Help notes	May source from patient's medical record (i.e. documentation of referral or
	smoking cessation support provided).

#### *Re-assessment medication adherence*

Reference number	28
Description	Medication adherence was re-assessed?
Codes and values	0 Yes
	1 No
	99 Unknown
	Is the patient adhering to prescribed treatment
	0 Yes
	1 No
	3 Unknown
	Reason for non-adherence
	Cost
	Cognitive issues
	Side effects/Contraindication
	Patient choice
	Unknown
Help notes	May source from patient's medical record, patient re-assessment. Use the same
	tool as used in initial assessment.

#### *Re-assessment of exercise capacity*

Reference number	29
Description	Functional exercise capacity outcome on reassessment
Codes and values	0 No
	1 Yes
	99 Unknown
	If no provide reason for not screening [open text].
Help notes	May source from patient's medical record, patient re-assessment. Use the same
	tool as used in initial assessment.

#### Change in exercise capacity

Reference number	30
Description	The difference between pre and post changes in exercise capacity measured using a
	validated measurement tool.
Codes and values	May be expressed as:
	a) 6MWT
	Absolute delta change in metres (numerical)
	b) EST and SAQ
	Absolute delta change in metabolic equivalences or METS (numerical)
Help notes	The Six Minute Walk Test
	An absolute delta change simply calculates the difference between the two
	distances:
	B1 - A1 = Absolute $\Delta$ , where B1 is the re-assessment 6MWT measurement and A1 is
	the initial 6MWT measurement.





i.e. 285 metres (re-assessment 6MWD measurement) – 224 metres (initial 6MWD measurement) = 61 metres; the patient has increased his/her exercise capacity by
61 metres.
The Exercise Stress Test
An absolute delta simply calculates the difference between two MET values:
B1 - A1 = Absolute $\Delta$ , where B1 is the reassessment MET measurement and A1 is
the initial MET measurement.
Self-reported exercise capacity
An absolute delta simply calculates the difference between two MET values:
B1 - A1 = Absolute $\Delta$ , where B1 is the reassessment MET measurement and A1 is
the initial MET measurement.

#### Re-assessment of health-related quality of life

Reference number	31
Description	Health-related quality of life was assessed same using a validated tool as per entry
	assessment.
Codes and values	0 No
	1 Yes
	99 Unknown
Help notes	May source from patient's medical record, patient re-assessment. Use the same
	tool as used in initial assessment.

#### Change in health-related quality of life

Reference number	32
Description	The difference between pre and post changes in HRQoL measured using a validated
	measurement tool.
Codes and values	Scores for each Domain as per validated tool used
Help notes	An absolute delta simply calculates the difference between two HRQoL values:
	B1 - A1 = Absolute $\Delta$ , where B1 is the reassessment HRQoL measurement and A1 is
	the initial HRQoL measurement.

#### Care transition

Reference number	33
Description	Is there documentation of communication with ongoing care providers?
Codes and values	0 Yes
	1 No
	99 Unknown
Help notes	May source from patient's medical record, patient re-assessment (e.g. documented
	communication with health care provider)

#### SERVICE DELIVERY INFORMATION

#### Mode of program delivery

Reference number	34
Description	The primary modality in which the CR program was delivered
Codes and values	Select all that apply
	1. Centre-based
	2. Community-based
	3. Home-based
	4. Telephone





	<ol> <li>5. Videoconference (e.g. via platform such as Zoom, Coviou, Skype)</li> <li>6. Web</li> </ol>
	7.Hybrid (combination of above)
	8. Other
Help notes	May source from patient's medical record or CR administrative database.

#### Frequency of program delivery

Reference number	35
Description	How often did the patient participate in the CR program?
Codes and values	1. once only
	2. 1 per week
	3. 2 times per week
	4. >2 times per week
Help notes	May source from patient's medical record or CR administrative database.

#### Content of program delivered

Reference number	36
Description	What aspects of the CR program did the patient receive?
Codes and values	Select all that apply
	Exercise and Education
	Exercise Only
	Education only – Individual
	Education only – Group
	Other
Help notes	May source from patient's medical record or CR administrative database.

#### Number of CR exercise sessions attended

Reference number	37
Description	Enter the total number of supervised exercise classes attended by the patient
	during their CR program (excluding pre and post assessments if applicable)
Codes and values	Number
Help notes	May source from patient's medical record or CR administrative database.

#### Number of CR education sessions attended

Reference number	38
Description	Enter the total number of educational classes attended by the patient during their
	CR program
Codes and values	Number
Help notes	May source from patient's medical record or CR administrative database.

#### CR completion

Reference number	38
Description	Patients have completed exercise sessions and reassessment of exercise.
Codes and values	0 Yes
	1 No
	99 Unknown
Help notes	May source from CR administrative database, re-assessment information.





#### Reasons for CR non-completion

Reference number	39
Description	If known, enter the reason for a CR registered patient to have not completed
	exercise sessions and reassessment of exercise using the below options.
Codes and values	1. N/A
	2. Incomplete - documented cardiac reason for noncompletion, such as an
	adverse health event and/or readmission
	3. Incomplete - documented non cardiac reason for noncompletion, such as
	an adverse health event and/or readmission
	4. Incomplete - death
	5. Incomplete - return to work
	6. Incomplete – relocation (referral to CR service closer to patient's home/
	referral to closest CR)
	<ol> <li>Incomplete - – Lack of contact/lost to follow-up</li> </ol>
	8. Incomplete – Transport issues
	9. Incomplete – lack of social supports
	10. Other
Help notes	May source from the patient's CR record.





## **SECTION B**

Version 10, June 2021





## Eligible in-patients are referred to cardiac rehabilitation within 3 calendar days of hospital discharge.

Short name	QI-1 – Referral to CR
Description	The percentage of eligible in-patients who are referred to a CR program within 3
	calendar days of hospital discharge
Individual data	Ref 12: Hospital discharge date
elements required to	Ref 13: CR referral date
be collected	
Method of	This variable will be reported as a percentage.
calculation	Step 1. CR referral date (Ref 13) – Hospital discharge date (Ref 12)
	Step 2. (Numerator/Denominator) * 100
Numerator	The total number of patients who were referred to a CR program within 3 calendar
	days of hospital discharge (Ref 13 - Ref 12) <u>&lt;</u> 3 days.
Denominator	The total number of CR eligible patients in the reference period as per eligibility
	definition (Table 1)
Rationale	CR participation significantly reduces mortality and morbidity. In-patient referral
	prior to discharge facilitates timely, universal access to CR.
Clinical	ACRA Core Component No. 1: All eligible patients must be offered referral to a CR
recommendations	service which best suits their individual needs, as soon as possible after diagnosis or
	before discharge from hospital.





## Eligible in-patients commence cardiac rehabilitation within 28 calendar days after hospital discharge.

Short name	QI-2 Time to enrolment
Description	The percentage of eligible in-patients who are enrolled in a CR program within 28
	calendar days of hospital discharge.
Individual data	Ref 12: Hospital discharge date
elements required to	Ref 15: Initial assessment date
be collected	
Method of	This variable will be reported as a percentage.
calculation	Step 1. Initial assessment date (Ref 15) – Hospital discharge date (Ref 12)
	Step 2. (Numerator/Denominator) * 100
Numerator	The number of patients who had an initial assessment within 28 calendar days of
	hospital discharge (Ref 15 - Ref 12) <u>&lt;</u> 28 days.
Denominator	The number of CR eligible patients in the reference period as per eligibility
	definition (Table 1)
Rationale	Current literature suggests that targeting earlier enrolment in rehabilitation
	improves overall enrolment, participation, and outcomes (Thomas et al., 2018; Pack
	et al., 2013; Collins et al., 2015, Candelaria et al.; 2021). Addressing time from
	discharge to start in rehabilitation is important in that it can influence potential
	processes or barriers at the patient (conflict with return to work), hospital,
	provider, and program (workflow and throughput) levels.
Clinical	ACRA Core Component No. 1: All eligible patients must be offered referral to a CR
recommendations	service which best suits their individual needs, as soon as possible after diagnosis or
	before discharge from hospital care.





## Patients who commence cardiac rehabilitation receive a comprehensive assessment of cardiovascular risk factors.

Short name	QI-3 Comprehensive assessment
Description	Percentage of patients who commence CR who have had a comprehensive
	assessment of cardiovascular risk factors.
Individual data	Ref 16 Depression screening
elements required to	Ref 18 Smoking status
be collected	Ref 20 Medication adherence
	Ref 21 Exercise capacity
Method of	This variable will be reported as a percentage.
calculation	(Numerator/Denominator) * 100
Numerator	The total number of patients enrolled in a CR program who received a
	comprehensive assessment (Ref 16, 18, 20, 21 assessed)
Denominator	The total number of patients enrolled in a CR program
Rationale	All major clinical CR guidelines (e.g. British, Scottish, Canadian, American, Australian
	Core Components) support individualised assessment of CR participants.
Clinical	ACRA Core Component No. 2: All eligible cardiac patients to receive an
recommendations	individualised initial assessment that includes physical, psychological and social
	parameters with referral on to appropriate services (internal or external to the CR
	service) based on patient needs; followed by ongoing review, discharge assessment
	and follow-up.





Patients who commence cardiac rehabilitation are screened for depression at initial and re-assessment and offered counselling (or referral to counselling) if symptoms are identified.

Short name	QI-4 Depression screening
Description	Percentage of patients who commence CR that are screened for depression at
	initial and reassessment using a valid and reliable screening tool and referred for
	counselling if symptoms are identified.
Individual data	Ref 16. Depression screening
elements required to	Ref 17. Depression referral
be collected	Ref 24. Re-assessment depression screening
	Ref 25. Re-assessment depression referral
Method of	Both variables will be reported as a percentage.
calculation	Step 1. Percentage of patients assessed/reassessed:
	(Numerator/Denominator) * 100
	Step 2. Percentage of patients screened positive referred
	(Numerator/Denominator) * 100
Numerator	Initial assessment:
	Step 1. The total number of patients who were screened for depression at initial
	assessment (Ref 16)
	Step 2. The total number of patients who screened positive and were referred (Ref
	17+) – (Ref 17)
	Re-assessment:
	Step 1. The total number of patients who screened for depression at re-assessment (Ref 24)
	Step 2. The total number of patients screened positive who were referred (Ref 25+) – (Ref 25)
Denominator	Step 1. The total number of patients enrolled in a CR program
	Step 2. The total number of patients that screened positive for depression
Rationale	All major clinical CR guidelines (e.g. British, Scottish, Canadian, American, Australian
	Core Components) support individualised assessment of CR participants.
Clinical	ACRA Core Component No. 2: All eligible cardiac patients to receive an
recommendations	individualised initial assessment that includes physical, psychological and social
	parameters with referral on to appropriate services (internal or external to the CR
	service) based on patient needs; followed by ongoing review, discharge assessment
	and follow-up.





Patients who comr	nence cardiac rehabilitation are assessed for smoking use at initial
and re-assessment	and offered smoking cessation counselling (or referral to
counselling) if they	are a current or recent smoker.
Short name	QI-5
Description	Percentage of patients who enrol in a CR program who are assessed for smoking
	use at initial assessment and offered or referred to smoking cessation counselling if
	they are a current or recent smoker.
Individual data	Ref 18 Smoking status
elements required to	Ref 19 Smoking referral
be collected	<ul> <li>Ref 26 Re-assessment smoking status</li> </ul>
	Ref 27 Re-assessment smoking referral
Method of	Both variables will be reported as a percentage.
calculation	Step 1. Percentage of patients assessed/reassessed:
	(Numerator/Denominator) * 100
	Step 2. Percentage of current or recent smokers referred
	(Numerator/Denominator) * 100
Numerator	Initial assessment:
	Step 1. The total number of patients who were assessed for smoking status at initial
	assessment (Ref 18)
	Step 2. The total number of current or recent smokers who were referred (Ref 19+
	– Ref 19)
	Pa accossment
	Stop 1. The total number of nationts who accossed for smoking status at re-
	assessment (Ref 26)
	Step 2. The total number of current or recent smokers who were referred (Ref 26+)
	- (Ref 27)
Denominator	Step 1. The total number of patients enrolled in a CR program
	Step 2. The total number of current or recent smokers
Rationale	Simple smoking cessation advice from health professionals increases the likelihood
	that someone who smokes will quit (Stead et al, 2013). Prompting a person to try to
	quit, brief reiteration of cardiovascular and other health hazards and agreeing on a
	specific plan with a follow-up arrangement are evidence-based interventions.
Clinical	ACRA Core Component No. 2: All eligible cardiac patients to receive an
recommendations	individualised initial assessment that includes physical, psychological and social
	parameters with referral on to appropriate services (internal or external to the CR
	service) based on patient needs; followed by ongoing review, discharge assessment
	and follow-up.





## Patients who commence cardiac rehabilitation are assessed for medication adherence at initial and re-assessment.

Short name	QI-6 Assessment of medication adherence
Description	The percentage of patients who are assessed for medication adherence at initial
	and re-assessment.
Individual data	Ref 20 Medication adherence
elements required to	Ref 28 Re-assessment medication adherence
be collected	
Method of	This variable will be reported as a percentage.
calculation	(Numerator/Denominator) * 100
Numerator	The total number of patients enrolled in a CR program who were assessed for
	medication adherence at initial and re-assessment (Ref 20 & Ref 28)
Denominator	The total number of patients enrolled in a CR program
Rationale	Medication adherence is often suboptimal for many reasons, including
	affordability, treatment complexity and lack of consumer understanding.
	Optimal medication adherence has been associated with a 20% reduction in
	cardiovascular disease risk and a 35% reduction in all-cause mortality (Chowdhury
	et al, 2013). CR is the ideal time to review and optimise cardio-protective therapies
	as per key guidelines (National Heart Foundation of Australia/Cardiac Society of
	Australia and New Zealand acute coronary syndromes guidelines (Chew et al, 2016).
Clinical	ACRA Core Component No. 2: All eligible cardiac patients to receive an
recommendations	individualised initial assessment that includes physical, psychological and social
	parameters with referral on to appropriate services (internal or external to the CR
	service) based on patient needs; followed by ongoing review, discharge assessment
	and follow-up.





## Patients who commence cardiac rehabilitation have an initial and re-assessment to determine exercise capacity change.

Short name	QI-7 Assessment of exercise capacity
Description	Change in exercise capacity from initial assessment to re-assessment.
Individual data	Ref 21 Exercise capacity
elements required to	Ref 29 Re-assessment of exercise capacity
be collected	
Method of	This variable will be reported as a percentage.
calculation	(Numerator/Denominator) * 100
Numerator	The total number of patients enrolled in a CR program who were assessed for
	exercise capacity at initial and re-assessment (Ref 21 & Ref 29)
Denominator	The total number of patients enrolled in a CR program
Rationale	Exercise training decreases mortality and morbidity in CHD patients (Anderson et
	al., 2016). As part of a comprehensive CR program, exercise training can reduce
	hospital admissions and increases health-related quality of life (Anderson et al.,
	2014). Pre and post assessment of functional exercise capacity is a valid way of
	evaluating the exercise component of CR programs. A half MET improvement is
	related to health benefit.
Clinical	ACRA Core Component No. 2: All eligible cardiac patients to receive an
recommendations	individualised initial assessment that includes physical, psychological and social
	parameters with referral on to appropriate services (internal or external to the CR
	service) based on patient needs; followed by ongoing review, discharge assessment
	and follow-up.





## Patients who commence cardiac rehabilitation have an initial and re-assessment to determine health-related quality of life change.

Short name	QI-8 Assessment of health-related quality of life
Description	Change in health-related quality of life from initial assessment to re-assessment.
Individual data	Ref 22 Health-related quality of life
elements required to	Ref 31 Re-assessment of health-related quality of life
be collected	
Method of	This variable will be reported as a percentage.
calculation	(Numerator/Denominator) * 100
Numerator	The total number of patients enrolled in a CR program who were assessed for
	HRQoL at initial and re-assessment (Ref 22 & Ref 31)
Denominator	The total number of patients enrolled in a CR program
Rationale	There is evidence that HRQoL can predict adverse outcomes (e.g. mortality)
	independent of traditional risk factors (Liang et al 2017).
Clinical	ACRA Core Component No. 3: CR services should facilitate patients to return to, or
recommendations	to improve on, baseline everyday functioning, including employment, driving,
	resumption of sexual activity, and other activities of daily living and maintain
	lifelong. When the cardiac condition or other co-morbidities preclude this, the CR
	service should focus on maximising potential and providing coping strategies.





#### Patients who participate in cardiac rehabilitation receive a comprehensive reassessment of their cardiovascular risk factors.

Short name	QI-9 Re-assessment
Description	Percentage of patients who commence CR who have had a comprehensive re-
	assessment of cardiovascular risk factors.
Individual data	Ref 23 Re-assessment date
elements required to	Ref 24 Re-assessment depression screening
be collected	Ref 26 Re-assessment smoking status
	Ref 28 Re-assessment medication adherence
	Ref 29 Re-assessment of exercise capacity
	Ref 37 Number of supervised exercise sessions attended
	Ref 38 CR completion
	Ref 39 Reasons for noncompletion
Method of	This variable will be reported as a percentage.
calculation	(Numerator/Denominator) * 100
Numerator	The total number of patients enrolled in a CR program who completed and received
	a comprehensive re-assessment (Ref 23, 24, 26, 28, 29, 37, 38 assessed)
Denominator	The total number of patients enrolled in a CR program
Rationale	Ideally, at re-assessment participants will have attained their CR goals, have ceased
	smoking, have depression treated, will be meeting exercise and nutrition
	guidelines, will be taking medications as prescribed and will be within
	recommended targets for blood pressure and lipids.
Clinical	ACRA Core Component No. 2: All eligible cardiac patients to receive an
recommendations	individualised assessment at CR discharge that includes physical, psychological and
	social parameters with referral on to appropriate services (internal or external to
	the CR service) based on patient needs.





Patients and ongoing care providers are provided with a report which outlines patient risk factors and an individualised ongoing management plan.

Short name	QI-10 Care transition
Description	Percentage of CR patients with documented communication which outlines patient
	risk factors and an ongoing management plan between the CR program and the
	ongoing care provider (e.g., general practitioner).
Individual data	Ref 33 Care transition
elements required to	
be collected	
Method of	This variable will be reported as a percentage.
calculation	(Numerator/Denominator) * 100
Numerator	The total number of patients who completed a CR program with documented
	communication and ongoing management plan from the CR program to at least
	one ongoing care provider (e.g., general practitioner)
Denominator	The total number of patients who completed the CR program (see Table 1 for
	definitions)
Rationale	Communication between CR programs and other health care providers may
	improve further management of patients' cardiac risk (Redfern & Briffa, 2014). A CR
	program should have strategies in place to communicate CR participants' status at
	entry/exit from the CR program to their general practitioner and other ongoing
	care providers (e.g., cardiologist, Phase III CR). Communication should be sent to
	the ongoing care provider at program exit regardless of patient continuation in a
	maintenance program.
Clinical	BACPR Core Components and Standards, 2012
recommendations	• On programme completion there should be a formal assessment. This should
	be communicated by discharge letter to the referrer and the patient as well as
	those directly involved in the continuation of healthcare provision.
	• There should be communication and collaboration between primary and
	secondary care services to achieve the long-term management plan.





# Appendix A. International classification of disease (ICD) codes eligible for cardiac rehabilitation or secondary prevention referral.

Code id	asci desc	
121.9	acute myocardial infarction	
121.0	ST elevation myocardial infarction	
121.4	non ST elevation myocardial infarction	
Procedures		
Coronary artery surgery		
38497-00	coronary artery bypass, using 1 saphenous vein graft	
38497-01	coronary artery bypass, using 2 saphenous vein graft	
38497-02	coronary artery bypass, using 3 saphenous vein graft	
38497-03	coronary artery bypass, using ≥4 saphenous vein graft	
38497-04	coronary artery bypass, using 1 other venous graft	
38497-05	coronary artery bypass, using 2 other venous grafts	
38497-06	coronary artery bypass, using 3 other venous grafts	
38497-07	coronary artery bypass, using ≥4 other venous grafts	
38500-00	coronary artery bypass, using 1 LIMA graft	
38500-01	coronary artery bypass, using 1 RIMA graft	
38500-02	coronary artery bypass, using 1 radial artery graft	
38500-03	coronary artery bypass, using 1 epigastric graft	
38500-04	coronary artery bypass using 1 other arterial graft	
38500-05	coronary artery bypass, using 1 composite graft	
38503-00	coronary artery bypass, using ≥ 2 LIMA grafts	
38503-01	coronary artery bypass, using ≥ 2 RIMA grafts	
38503-02	coronary artery bypass, using $\geq$ 2 radial artery grafts	
38503-03	coronary artery bypass, using ≥ 2 epigastric grafts	
38503-05	coronary artery bypass, using ≥2 composite grafts	
90201-00	coronary artery bypass using 1 other graft not elsewhere classified	
90201-01	coronary artery bypass using 2 other grafts not elsewhere classified	
90201-02	coronary artery bypass using 3 other grafts not elsewhere classified	
90201-03	coronary artery bypass using ≥4 other grafts not elsewhere classified	
Percutaneous	coronary intervention	
38300-00	percutaneous transluminal balloon angioplasty of 1 coronary artery	
38300-01	open transluminal balloon angioplasty of 1 coronary artery	
38303-00	percutaneous transluminal balloon angioplasty of ≥2 coronary arteries	
38303-01	open transluminal balloon angioplasty of ≥2 coronary arteries	
38306-00	percutaneous insertion of 1 transluminal stent into a single coronary artery	
38306-01	percutaneous insertion of ≥2 transluminal stents into a single coronary artery	
38306-02	percutaneous insertion of ≥2 transluminal stents into multiple coronary arteries	
38306-03	open insertion of 1 transluminal stent into a single coronary artery	
38306-04	open insertion of $\geq$ 2 transluminal stents into a single coronary artery	
38306-05	open insertion of $\geq 2$ transluminal stents into multiple coronary arteries	





<u>Valves</u>	
38475-00	mitral valve annuloplasty
38475-01	tricuspid valve annuloplasty
38475-02	aortic valve annuloplasty
38477-00	mitral valve annuloplasty with ring insertion
38477-01	tricuspid valve annuloplasty with ring insertion
38477-02	aortic valve annuloplasty with ring insertion
38480-00	repair of aortic valve, 1 leaflet
38480-0	repair of mitral valve, 1 leaflet
38480-02	repair of tricuspid valve, 1 leaflet
38481-00	repair of aortic valve, 2 or more leaflets
38481-01	repair of mitral valve, ≥2 leaflet
38481-02	repair of tricuspid valve, ≥2 leaflet
38488-00	replacement of aortic valve with mechanical prosthesis
38488-01	replacement of aortic valve with bioprosthesis
38488-02	replacement of mitral valve with mechanical prosthesis
38488-03	replacement of mitral valve with bioprosthesis
38488-04	replacement of tricuspid valve with mechanical prosthesis
38488-05	replacement of tricuspid valve with bioprosthesis
38488-08	percutaneous replacement of aortic valve with bioprosthesis
38488-09	percutaneous replacement of mitral valve with bioprosthesis
38488-10	percutaneous replacement of tricuspid valve with bioprosthesis
38489-00	replacement of aortic valve with homograft
38489-01	replacement of aortic valve with unstented heterograft
38489-02	replacement of mitral valve with homograft
38489-03	replacement of tricuspid valve with homograft

Citation: Independent Hospital Pricing Authority, Australian Classification of Health Interventions

(ACHI), First Eleventh edition. 2019 11<sup>th</sup> edition.





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