# Factors that influence whether patients with acute coronary syndromes undergo cardiac catheterisation

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**The known**: The management of patients in Australia presenting to hospital with acute coronary syndrome is reported to differ according to whether the hospital of first presentation can provide cardiac catheterisation.

**The new**: For Australian hospitals enrolled in the CONCORDANCE clinical quality registry, differences in catheterisation rates and major outcomes between patients who initially present to catheterisation-capable or non-capable hospitals are not as large as found by earlier studies.

**The implications**: Improved access to angiographic facilities across the Australian health care system has largely overcome challenges to providing equitable outcomes for all patients with acute coronary syndromes.

nvasive coronary angiography during hospital admission reduces the frequency of adverse outcomes (mortality, recurrent myocardial infarction, stroke) for patients with non-ST elevation acute coronary syndromes (NSTEACS). Guidelines recommend and immediate invasive strategy (within two hours of admission) for patients at very high risk, an early invasive strategy (within 24 hours) for patients at high risk, and an invasive strategy (within 72 hours) for patients at intermediate risk of cardiovascular events.<sup>2</sup> Similarly, time is critical when treating patients with an ST elevation myocardial infarction (STEMI).3 Reperfusion can be achieved by primary percutaneous coronary intervention (PCI) or fibrinolysis, but PCI is preferred if readily available, because of its better outcomes.<sup>4</sup> Further, emergency angiography is required for 18-32% of patients in whom fibrinolysis fails to achieve reperfusion, and angiography within 3-24 hours of fibrinolysis is recommended for all other patients to treat residual stenoses and to limit re-occlusion.<sup>2</sup>

Timely access to angiography is consequently advocated by Australian guidelines as an important component of care for patients with acute coronary syndrome (ACS). However, inequities in access have been described, driven in part by geographic challenges, and may be associated with poorer outcomes for patients presenting to hospitals that cannot provide angiography. 5-9

The aims of our study were to determine whether the availability of invasive coronary angiography at the hospital of presentation influences catheterisation rates for patients presenting with ACS; whether the clinical and risk characteristics of patients selected for angiography differ between hospitals of presentation according to whether they have on-site catheterisation laboratories; and whether presenting to a catheterisation-capable hospital is associated with better outcomes for patients with ACS.

### Methods

We analysed data from the Cooperative National Registry of Acute Coronary Events (CONCORDANCE). The registry

#### **Abstract**

**Objective:** To determine whether the availability of invasive coronary angiography at the hospital of presentation influences catheterisation rates for patients with acute coronary syndrome (ACS), and whether presenting to a catheterisation-capable hospital is associated with better outcomes for patients with ACS.

**Design, setting:** Retrospective cohort study; analysis of Cooperative National Registry of Acute Coronary Events (CONCORDANCE) data.

**Setting, participants:** Adults admitted with ACS to 43 Australian hospitals (including 31 catheterisation-capable hospitals), February 2009 – October 2018.

**Main outcome measures:** Major adverse cardiovascular events (myocardial infarction, stroke, congestive heart failure, cardiogenic shock, cardiovascular death) and all-cause deaths in hospital and by six and 12- or 24-month follow-up.

**Results:** The proportion of women among the 5637 patients who presented to catheterisation-capable hospitals was smaller than for the 2608 patients who presented to hospitals without catheterisation facilities (28% v 33%); the proportion of patients diagnosed with ST elevation myocardial infarction was larger (32% v20%). The proportions of patients who underwent catheterisation (81% v70%) or percutaneous coronary intervention (49% v35%) were larger for those who presented to catheterisation-capable hospitals. The baseline characteristics of patients who underwent catheterisation were similar for both presentation hospital categories, as were rates of major adverse cardiovascular events and all-cause death in hospital and by 6- and 12- or 24-month follow-up.

**Conclusions:** Although a larger proportion of patients who presented to catheterisation-capable hospitals underwent catheterisation, patients with similar characteristics were selected for the procedure, independent of the hospital of presentation. Major outcomes for patients were also similar, suggesting equitable management of patients with ACS across Australia.

collected information on patients who presented with ACS from the medical records of 43 metropolitan and regional hospitals from all Australian states and territories during 23 February 2009 – 16 October 2018 (Supporting Information, table 1). Catheterisation-capable hospitals were defined as those with, and catheterisation non-capable hospitals as those without cardiac catheterisation facilities. All patients were followed up by phone six months after discharge and again at 12 months (patients enrolled 2014–2018) or 24 months (patients enrolled 2009–2013).

### Patients: eligibility criteria and classification

Eligible for inclusion in the CONCORDANCE registry were all patients over 18 years of age admitted to hospital with presumed ACS who had elevated cardiac biomarker levels, an electrocardiographic abnormality, prior history of coronary heart disease, newly documented coronary heart disease, or two

high risk features predictive of recurrent in-hospital events: haemodynamic compromise (systolic blood pressure less than 90 mmHg and heart rate exceeding 100 per minute), left ventricular systolic dysfunction (left ventricular ejection fraction less than 0.40%), diabetes, or chronic kidney disease (estimated glomerular filtration rate less than 60 mL/min/1.73 m<sup>2</sup>).

We included patients with confirmed ACS for whom the index admission hospital contributed data to CONCORDANCE, so that complete medical history and in-hospital data were available for analysis. Patients were first classified according to whether they were admitted to hospitals with or without onsite cardiac catheterisation facilities; the catheterisation laboratory status did not change for any included hospital during the study period. Patients were then classified according to whether they underwent cardiac catheterisation during their index admission.

#### **Outcomes**

Outcomes were major adverse cardiovascular events (MACE) — myocardial infarction, stroke, congestive heart failure, shock, or cardiovascular death — and all-cause deaths in hospital and by follow-up at 12 or 24 months. The process measure was prescription on discharge of at least four of five indicated evidence-based medication types: aspirin, platelet ADP receptor-inhibiting agents (clopidogrel, prasugrel, or ticagrelor), a statin or other lipid-lowering medication, beta blockers, and angiotensin-converting enzyme inhibitors or angiotensin receptor blockers.

### Statistical analysis

Missing baseline categorical data were recoded as "no" or "normal"; that is, if something was not documented, it was deemed not to have been present. Analyses were not adjusted for other missing data. Demographic data, clinical characteristics, inhospital management, and events were compared for patients who presented to catheterisation-capable and non-capable hospitals. Categorical data are summarised as numbers and proportions; the statistical significance of differences was assessed in Rao–Scott  $\chi^2$  tests. Continuous data were summarised as means with standard deviation (SDs) and compared in unadjusted regression analyses in a generalised estimating equations (GEE) framework with an exchangeable correlation matrix. The normality of continuous data was assessed in Kolmogorov–Smirnov tests. Rao–Scott testing and the GEE analysis each accounted for clustering by hospital.

We assessed factors associated with outcomes by multivariable logistic regression in a GEE framework. For in-hospital outcomes, we included cardiac catheterisation, age (four categories), sex, Indigenous status, diabetes, peripheral artery disease, history of myocardial infarction, congestive heart failure, chronic renal failure, dementia, ACS diagnosis, catheterisation laboratory availability, urban/rural hospital, and hospital size (bed number; three groups) as independent variables. For follow-up outcomes, we included these variables as well as referral to a cardiac rehabilitation program and discharge prescription of four or more of the five evidence-based medication types. P < 0.05 (two-tailed) was deemed statistically significant. Statistical analyses were conducted in SAS 9.4.

### **Ethics approval**

Ethics approval for our study was granted by the human research ethics committee of the Concord Repatriation General

Hospital (reference, HREC/08/CRGH/180). Each participating site had appropriate approval for data collection.

### **Results**

CONCORDANCE data were available for 8245 patients: 5637 (68%) who had presented to 31 catheterisation-capable hospitals and 2608 (32%) who had presented to 12 catheterisation non-capable hospitals. The proportion of women among patients who presented to catheterisation-capable hospitals was smaller than for catheterisation non-capable hospitals (1575 [28%]) v 873 [33%]), but the mean ages of all patients were similar (65.3 years [SD, 13.4 years] v 66.7 years [SD, 13.5 years]). The proportions of patients with hypertension (3471 [62%] v1730 [66%]), chronic renal failure (438 [8%] v 310 [12%]), prior stroke (409 [7%] v 242 [9%]), and prior myocardial infarction (1705 [30%] v 922 [35%]) were significantly lower for patients who presented to catheterisation-capable hospitals, and that of patients diagnosed with STEMI significantly higher (1786 [32%] v 510 [20%]) (Box 1). At total of 1144 patients could not be contacted for the 6-month follow-up, and 2924 patients could not be contacted for the 12- or 24-month follow-up.

#### Cardiac catheterisation and revascularisation

The proportions of patients who underwent cardiac catheterisation (4557 [81%] v 1832 [70%]) or PCI (2739 [49%] v 915 [35%]) were each higher for those who presented to catheterisationcapable hospitals (Box 2). Catheterisation rates for people who presented to catheterisation-capable hospitals were higher than for those who presented to catheterisation non-capable hospitals for all ACS diagnoses: STEMI, 1698 (95%) v 454 (89%); non-STEMI (NSTEMI), 2170 (82%) v 1067 (74%); unstable angina, 689 (58%) v 311 (47%). The baseline characteristics of patients admitted to hospitals with or without catheterisation facilities were similar, both for those who underwent cardiac catheterisation (Supporting Information, table 2) and those who did not (Supporting Information, table 3). A larger proportion of patients who presented to catheterisation-capable hospitals were discharged with at least four of the five evidence-based medication types than of those who presented to hospitals without catheterisation facilities (4033 [72%] v 1728 [66%]) (Box 2).

## Patient outcomes in catheterisation-capable and catheterisation non-capable hospitals

In-hospital outcomes during the index admission were similar for hospitals with and without catheterisation facilities (MACE: 712 [13%] v 326 [13%]; all-cause death, 219 [4%] v 93 [4%]) (Box 3), as were outcomes at 6-month (MACE 283 [6%] v 154 [7%]; all-cause deaths, 146 [3%] v 90 [4%]; Box 4) and 12- or 24-month follow-up (MACE, 169 [32%] v 93 [36%]; all-cause deaths, 99 [3%] v 48 [3%]; Box 5).

### Factors associated with adverse outcomes: in-hospital outcomes

Presenting to a hospital with a catheterisation laboratory did not influence the frequency of in-hospital MACE (adjusted odds ratio [aOR], 0.90; 95% confidence interval [CI], 0.55–1.48) or death (aOR, 1.11; 95% CI, 0.76–1.62). Undergoing cardiac catheterisation was associated with lower rates of MACE (aOR, 0.48; 95% CI, 0.38–0.61) and death (aOR, 0.07; 95% CI, 0.04–0.13) (Box 6; online Supporting Information, figure 1).

The odds of in-hospital MACE was higher for older patients, for patients with diabetes, chronic renal failure, or dementia or

Characteristic	Catheterisation-capable hospital	Catheterisation non-capable hospital	Total	P
Number of patients	5637	2608	8245	
Age (years), mean (SD)	65.3 (13.4)	66.7 (13.5)	65.8 (13.4)	0.41
Age group (years)				0.17
< 55	1214 (22%)	512 (20%)	1726 (21%)	
55-64	1450 (26%)	561 (22%)	2011 (24%)	
65–74	1494 (27%)	722 (28%)	2216 (27%)	
≥ 75	1460 (26%)	809 (31%)	2269 (28%)	
Sex (women)	1575 (28%)	873 (33%)	2448 (30%)	< 0.001
Body mass index (kg/m²)				0.43
< 18 (underweight)	38 (1%)	21 (2%)	59 (1%)	
18 to < 25 (healthy)	988 (26%)	344 (26%)	1332 (26%)	
25 to < 30 (overweight)	1490 (39%)	488 (37%)	1978 (38%)	
≥ 30 (obese)	1314 (34%)	480 (36%)	1794 (35%)	
Indigenous Australians	199 (4%)	202 (8%)	401 (5%)	0.28
Smoking history	· •			0.66
Never smoked	2143 (38%)	1057 (41%)	3200 (39%)	
Ex-smoker	2008 (36%)	908 (35%)	2916 (35%)	
Current smoker	1474 (26%)	630 (24%)	2104 (26%)	
Comorbid conditions				
Hypertension	3471 (62%)	1730 (66%)	5201 (63%)	0.003
Diabetes	1550 (27%)	809 (31%)	2359 (29%)	0.14
Dyslipidaemia	3125 (56%)	1538 (59%)	4663 (57%)	0.11
Peripheral arterial disease	336 (6%)	188 (7%)	524 (6%)	0.20
Prior myocardial infarction	1705 (30%)	922 (35%)	2627 (32%)	0.008
Prior percutaneous coronary intervention	1235 (22%)	635 (24%)	1870 (23%)	0.10
Prior coronary artery bypass graft	662 (12%)	350 (13%)	1012 (12%)	0.21
Prior heart failure	452 (8%)	229 (9%)	681 (8%)	0.48
Prior stroke/transient ischemic attack	409 (7%)	242 (9%)	651 (8%)	0.041
Chronic renal failure	438 (8%)	310 (12%)	748 (9%)	< 0.001
Dementia/cognitive impairment	161 (3%)	103 (4%)	264 (3%)	0.08
Killip class				0.40
1	4955 (88%)	2302 (88%)	7257 (88%)	
2	538 (10%)	230 (9%)	768 (9%)	
3	100 (2%)	65 (2%)	165 (2%)	
4	44 (1%)	11 (< 1%)	55 (1%)	
GRACE risk score, mean (SD)	108.2 (31.9)	108.6 (32.3)	108.3 (32.0)	0.66
Cardiac arrest on admission	236 (4%)	73 (3%)	309 (4%)	0.040
Diagnosis			( ,	0.001
ST-elevation myocardial infarction	1786 (32%)	510 (20%)	2296 (28%)	
Non-ST-elevation myocardial infarction	2661 (47%)	1442 (55%)	4103 (50%)	
Unstable angina	1187 (21%)	655 (25%)	1842 (22%)	
Admitted to a rural hospital	1429 (25%)	1647 (63%)	3076 (37%)	0.06
Beds in hospitals	1123 (2370)	.5 17 (05.10)	3370 (3770)	< 0.001
Tertile 1 (< 330)	671 (12%)	1817 (70%)	2488 (30%)	0.001
Tertile 2 (330–600)	2246 (40%)	561 (22%)	2807 (34%)	
Tertile 3 (> 600)	2720 (48%)	230 (9%)	2950 (36%)	

GRACE = Global Registry of Acute Coronary Events;<sup>11</sup> SD = standard deviation.

### 2 Hospital investigations and treatment, and discharge medications for 8245 patients admitted with presumed acute coronary syndrome, Australia, Feb 2009 – Oct 2018

Characteristic	Catheterisation-capable hospital	Catheterisation non-capable hospital	Total	P
Number of patients	5637	2608	8245	
Cardiac catheterisation	4557 (81%)	1832 (70%)	6387 (78%)	< 0.001
Revascularisation (any type)	3153 (56%)	1131 (43%)	4284 (52%)	0.001
Coronary artery bypass graft	446 (8%)	224 (9%)	670 (8%)	0.47
Percutaneous coronary intervention	2739 (49%)	915 (35%)	3654 (44%)	< 0.001
Percutaneous coronary intervention: type				< 0.001
Primary	1606 [59%]	107 [12%]	1713 [47%]	
Rescue	170 [6%]	170 [19%]	340 [9%]	
Other	962 [35%]	637 [70%]	1599 [44%]	
Discharge medications				
Aspirin	4811 (85%)	2179 (84%)	6990 (85%)	0.31
Clopidogrel	2388 (42%)	1283 (49%)	3671 (45%)	0.06
Prasugrel	302 (5%)	39 (1%)	341 (4%)	< 0.001
Ticagrelor	1081 (19%)	318 (12%)	1399 (17%)	0.08
Coplavix (clopidogrel/aspirin)	212 (4%)	69 (3%)	281 (3%)	0.36
Lipid-lowering agent	5043 (89%)	2243 (86%)	7286 (88%)	0.022
Beta blocker	4190 (74%)	1885 (72%)	6075 (74%)	0.30
Angiotensin-converting enzyme inhibitors or angiotensin receptor blockers	3905 (69%)	1721 (66%)	5626 (68%)	0.14
Four or more evidence-based medicine types	4033 (72%)	1728 (66%)	5761 (70%)	0.018
Anticoagulant	487 (9%)	282 (11%)	769 (9%)	< 0.001
Referral to cardiac rehabilitation	3488 (62%)	1565 (61%)	5053 (62%)	0.85

### 3 In-hospital outcomes for 8245 patients admitted to hospital with presumed acute coronary syndrome, Australia, Feb 2009 - Oct 2018

Outcome	Catheterisation-capable hospital	Catheterisation non-capable hospital	Total	P
Number of patients	5637	2608	8245	
Major adverse cardiac events*	712 (13%)	326 (13%)	1038 (13%)	0.84
Myocardial infarction	133 (2%)	55 (2%)	188 (2%)	0.73
Stroke	31 (1%)	21 (1%)	52 (1%)	0.16
Congestive heart failure	478 (9%)	214 (8%)	692 (9%)	0.97
Cardiovascular death	192 (3%)	83 (3%)	275 (3%)	0.82
All-cause deaths	219 (4%)	93 (4%)	312 (4%)	0.74
Renal failure	329 (6%)	125 (5%)	454 (6%)	0.29
Major bleeding	428 (8%)	191 (7%)	619 (8%)	0.83
Cardiogenic shock	186 (3%)	51 (2%)	237 (3%)	< 0.001

cognitive impairment; it was higher for patients diagnosed with STEMI or NSTEMI than for those diagnosed with unstable angina (Box 6).

The odds of in-hospital death were higher for older patients, for patients with prior myocardial infarction, chronic renal failure, or dementia or cognitive impairment; they were higher for patients diagnosed with STEMI or NSTEMI than for those

diagnosed with unstable angina (Supporting Information, figure 1).

# Factors associated with adverse outcomes: six-month follow-up

Six-month follow-up was completed for 6759 patients (82%); the baseline characteristics of patients followed up and those

### 4 Outcomes by 6-month follow-up for 6759 patients admitted to hospital with presumed acute coronary syndrome, Australia, Feb 2009 – Oct 2018

Outcome	Catheterisation-capable hospital	Catheterisation non-capable hospital	Total	P
Number of patients	4642	2117	6759	
Major adverse cardiac events*	283 (6%)	154 (7%)	437 (7%)	0.18
Myocardial infarction	120 (3%)	65 (3%)	185 (3%)	0.51
Stroke	22 (0%)	14 (1%)	36 (1%)	0.41
Congestive heart failure	138 (3%)	79 (4%)	217 (3%)	0.11
Cardiovascular death	48 (1%)	24 (1%)	72 (1%)	0.81
All-cause deaths	146 (3%)	90 (4%)	236 (4%)	0.09
Major bleeding	34 (1%)	26 (1%)	60 (1%)	0.12
Re-hospitalised for heart disease or a bleeding event	941 (21%)	478 (23%)	1419 (22%)	0.25

<sup>\*</sup> Myocardial infarction, stroke, congestive heart failure, cardiogenic shock, or cardiovascular death.

### 5 Outcomes by 12- or 24-month follow-up for 4979 patients admitted to hospital with presumed acute coronary syndrome, Australia, Feb 2009 – Oct 2018

Outcome	Catheterisation-capable hospital	Catheterisation non-capable hospital	Total	P
Number of patients	3592	1387	4979	
Major adverse cardiac events*	169 (32%)	93 (36%)	262 (33%)	0.23
Myocardial infarction	86 (17%)	46 (18%)	132 (17%)	0.53
Stroke	13 (2%)	11 (4%)	24 (3%)	0.06
Congestive heart failure	80 (15%)	40 (16%)	120 (15%)	0.86
Cardiovascular death	28 (1%)	9 (1%)	37 (1%)	0.58
All-cause deaths	99 (3%)	48 (3%)	147 (3%)	0.32
Major bleeding	31 (6%)	16 (6%)	47 (6%)	0.85
Re-hospitalised for heart disease or a bleeding event	523 (15%)	253 (18%)	776 (16%)	0.10
* Myocardial infarction, stroke, congestive heart failure, cardiogenic shock, or cardiovascular death.				

not followed up were similar (Supporting Information, table 4). Presenting to a hospital with a catheterisation laboratory did not influence the odds of MACE (aOR, 1.04; 95% CI, 0.84–1.28) or death by six months (aOR, 1.01; 95% CI, 0.61–1.68). Undergoing cardiac catheterisation was associated with lower rates of MACE (aOR, 0.52; 95% CI, 0.39–0.69) and death (aOR, 0.60; 95% CI, 0.42–0.86) (Box 7; Supporting Information, figure 2).

The odds of MACE by 6-month follow-up were higher for older patients, for patients with diabetes mellitus, peripheral arterial disease, prior myocardial infarction or heart failure, or chronic renal failure, and for Indigenous Australian patients; they were higher for patients diagnosed with STEMI or NSTEMI than for those diagnosed with unstable angina (Box 7).

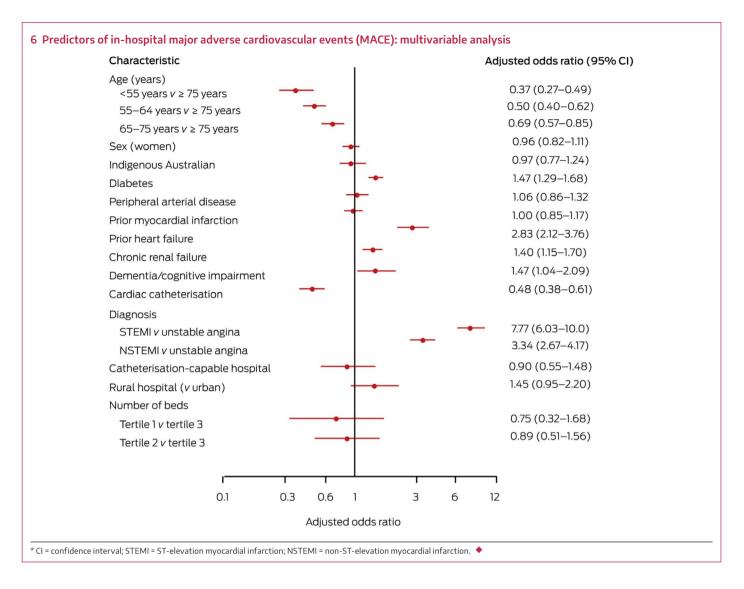
The odds of death by 6-month follow-up were higher for older patients, for patients with prior heart failure, chronic renal failure, or dementia or cognitive impairment, and for Indigenous Australian patients; they were higher for patients diagnosed with STEMI or NSTEMI than for those diagnosed with unstable angina. It was lower for patients referred to cardiac rehabilitation (Supporting Information, figure 2).

# Factors associated with adverse outcomes: 12- or 24-month follow-up

Twelve- or 24-month follow-up was completed for 4979 patients (60%); the baseline characteristics of patients followed up and those not followed up were similar (Supporting Information, table 5). Presenting to a hospital with a catheterisation laboratory did not influence the odds of MACE (aOR, 0.95; 95% CI, 0.65–1.39) or death (aOR, 1.11; 95% CI, 0.61–2.04). Undergoing cardiac catheterisation was associated with lower rates of MACE (aOR, 0.63; 95% CI, 0.44–0.91) and death (aOR, 0.50; 95% CI, 0.33–0.74) (Box 8, Supporting Information, figure 3).

The odds of MACE by 12- or 24-month follow-up were higher for patients with peripheral arterial disease, prior heart failure, chronic renal failure, or dementia or cognitive impairment; they were higher for patients diagnosed with STEMI or NSTEMI than for those diagnosed with unstable angina (Box 8).

The odds of death by 12- or 24-month follow-up were higher for older patients, and for patients with prior myocardial infarction or heart failure or chronic renal failure; they were higher for patients diagnosed with NSTEMI than for those diagnosed with



unstable angina, and lower for those referred to cardiac rehabilitation (Supporting Information, figure 3).

#### Discussion

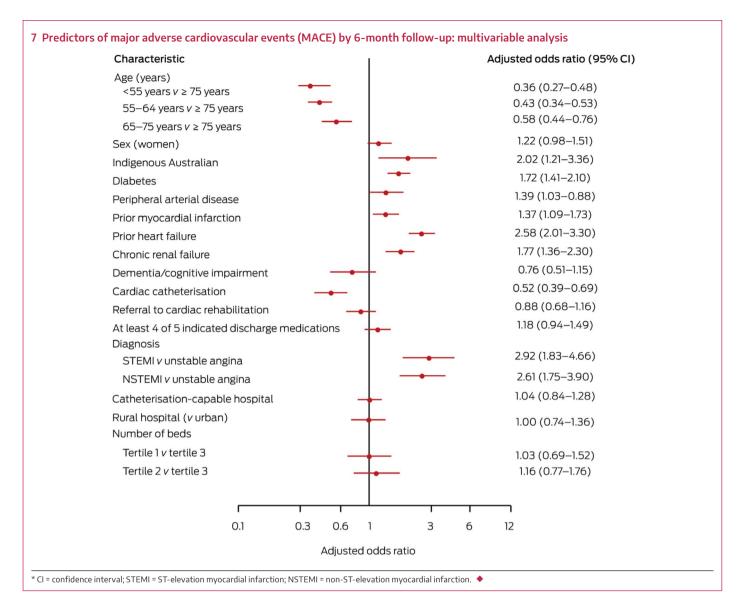
In our comprehensive analysis of 8245 patients who presented with ACS to 43 Australian hospitals during 2009–2018, the availability of coronary angiography at the hospital of presentation influenced catheterisation rates. However, the clinical characteristics of patients who underwent catheterisation were similar for both hospital categories, which suggests that practice patterns and decisions to offer invasive treatment are similar in both settings. Despite differences in the proportions of patients who ultimately underwent catheterisation, outcomes for patients with ACS who presented to catheterisation-capable and non-capable hospitals were similar, suggesting that they are not significantly influenced by the availability of on-site coronary angiography in the hospital of presentation.

The angiography rates we report — 81% in catheterisation-capable hospitals, 70% in hospitals without these facilities — are higher than those found by previous studies. The Heart Protection Project, conducted in 27 Australian hospitals during 2003–2005, reported invasive angiography rates of 66.4% in catheterisation-capable and 20.1% in catheterisation non-capable hospitals; the 2012 SNAPSHOT ACS audit (Australia and New

Zealand) reported rates of 61.5% and 50.8% respectively.<sup>6</sup> The differences presumably reflect growing acceptance of coronary angiography as the standard of care for patients with ACS at high risk of cardiovascular events, and the increased capacity to transfer patients to catheterisation-capable hospitals that contribute data to the CONCORDANCE registry.

Despite differences in the proportions who received angiography after presenting to catheterisation-capable and non-capable hospitals, the clinical characteristics of patients selected for angiography were similar. Diagnoses of STEMI were more frequent among people who presented to hospitals with catheterisation laboratories, probably reflecting pre-hospital triage of patients and direction of patients with STEMI to primary PCI hospitals. This could also explain the mildly higher comorbidity rates (prior myocardial infarction, prior stroke, hypertension, chronic renal failure) among patients presenting to hospitals without catheterisation facilities, reflecting the higher proportion of patients with NSTEACS. Similarly, catheterisation rates were higher among patients presenting to catheterisation-capable hospitals for all ACS diagnoses, probably reflecting greater comorbidity among patients who presented to catheterisation non-capable hospitals. 12

Outcomes in hospital and at 6 and 12 or 24 months after discharge were similar for patients admitted to catheterisation-capable and non-capable hospitals. A number of baseline characteristics — prior heart failure, chronic renal failure, peripheral arterial



disease, diabetes mellitus, dementia or cognitive impairment, and diagnoses of STEMI and NSTEMI — were associated with poor outcomes in both the short and long terms, consistent with previous reports. After adjusting for these factors, cardiac catheterisation was associated with lower incidence of MACE and death at all time points, indicating the importance of invasive coronary angiography and appropriate revascularisation. Importantly, patients with similar characteristics were offered cardiac catheterisation, regardless of whether the hospital to which they presented had a catheterisation laboratory or not, suggesting that triage and transfer systems in hospitals enrolled in CONCORDANCE are sufficiently mature to overcome previously documented inequities in access to invasive procedures. 5-7.9

### Limitations

Confounding in our retrospective cohort study and loss of patients to follow-up may have affected our findings. While the CONCORDANCE registry included both urban and rural tertiary hospitals from all Australian states and territories, smaller hospitals in rural and remote areas were not included; our findings may not reflect practice in these less well resourced hospitals. Analyses were not adjusted for multiple comparisons.

### **Conclusions**

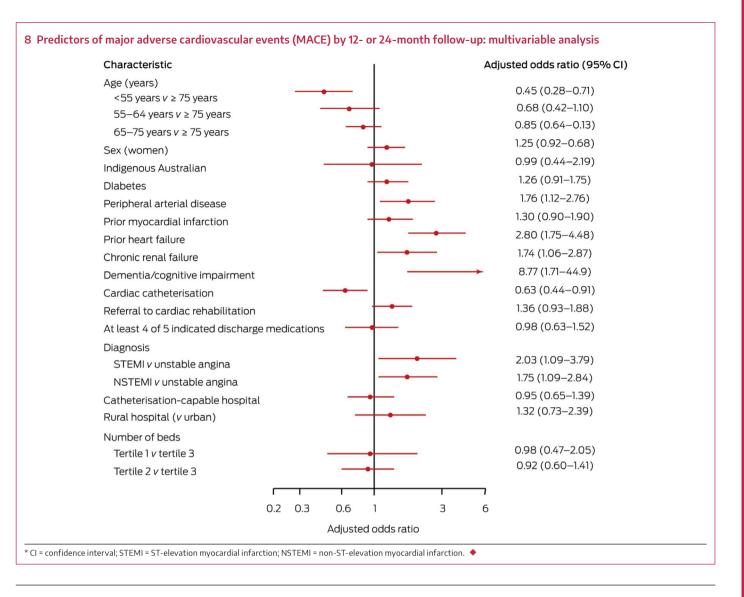
It has been reported that the management of patients presenting with ACS to Australian hospitals differed according to whether the hospital of first presentation had a catheterisation laboratory. We found, however, that catheterisation rates were higher than previously reported, regardless of whether the hospital of presentation was catheterisation-capable or not. Adverse outcomes were less frequent among patients who received coronary angiography, and patients with similar characteristics received this treatment, whether they presented to catheterisation-capable or non-capable hospitals, consistent with the similar outcomes for these patients.

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### Supporting Information