

openhart Prehospital identification of ST-segment elevation myocardial infarction and mortality (ANZACS-QI 61)

Becky Yi-Wen Liao ^{1,2}, Mildred Ai Wei Lee,¹ Bridget Dicker ^{3,4},
Verity F Todd,^{3,4} Ralph Stewart ², Katrina Poppe ⁵, Andrew Kerr^{5,6}

► Additional supplemental material is published online only. To view, please visit the journal online (<http://dx.doi.org/10.1136/openhrt-2021-001868>).

To cite: Liao BY-W, Lee MAW, Dicker B, *et al.* Prehospital identification of ST-segment elevation myocardial infarction and mortality (ANZACS-QI 61). *Open Heart* 2022;**9**:e001868. doi:10.1136/openhrt-2021-001868

Received 22 September 2021
Accepted 5 January 2022

ABSTRACT

Background Early recognition of ST-segment elevation myocardial infarction (STEMI) is needed for timely cardiac monitoring and reperfusion therapy.

Methods Three anonymously linked New Zealand national datasets (July 2016–November 2018) were used to assess the utilisation of ambulance transport in STEMI cases, the concordance between ambulance initial clinical impressions and hospital STEMI diagnoses, and the association between initial paramedic clinical impressions and 30-day mortality. The St John Ambulance electronic record captures community call-outs and paramedic initial clinical impressions. The national cardiac (ANZACS-QI) registry and national administrative datasets capture all New Zealand public hospital admission diagnoses and mortality data.

Results Of 5465 patients with STEMI, 73% were transported to hospital by ambulance. For these patients, the initial paramedic impression was STEMI in 50.7%, another acute coronary syndrome (ACS) diagnosis in 19.9% and non-ACS diagnosis in 29.7%. Only 37% of the 5465 patients with STEMI were both transported by ambulance and clinically suspected of STEMI by paramedics. Compared with patients with paramedic-‘suspected STEMI’, 30-day mortality was over threefold higher for patients thought to have a non-ACS condition (10.9% and 34.9%, respectively), but after adjustment for available covariates, this was substantially ameliorated (HR 1.48, 95% CI 1.22 to 1.80).

Conclusions In this national data linkage study, only 4 out of every 10 patients with STEMI were both transported by ambulance and had STEMI suspected by paramedics. Although patients with STEMI not suspected of an ACS diagnosis by paramedics had the highest mortality rate, this is largely explained by the different risk profile of these patients.

INTRODUCTION

Timely and accurate diagnosis of ST-segment elevation myocardial infarction (STEMI) is a prerequisite for implementation of optimal evidence-based treatments including antithrombotic therapy and coronary reperfusion.^{1–8} The St John Ambulance service is a charity with funding from government and community donations, attends to most

Key questions

What is already known about this subject?

► Timely and accurate diagnosis of ST-segment elevation myocardial infarction (STEMI) is important to facilitate cardiac monitoring and acute reperfusion therapy. Prior studies have reported variable accuracy in STEMI diagnoses by paramedics, but the overall proportion of all hospitalised patients with STEMI who are transported by ambulance and had STEMI suspected by paramedics is not known.

What does this study add?

► This national data linkage study anonymously linked the New Zealand national cardiac (ANZACS-QI) registry, national administrative datasets and the St John Ambulance electronic records to identify all patients hospitalised with STEMI. This allowed us to report the proportion presenting via ambulance and the accuracy of the paramedic clinical impressions. Only 4 out of every 10 hospitalised patients with STEMI were both transported by ambulance and had STEMI suspected by paramedics. Patients with STEMI who were not suspected of an acute coronary syndrome diagnosis by paramedics had a higher mortality rate than those suspected of STEMI or who self-transported to hospital, although this is largely explained by the different risk profile of these patients.

How might this impact on clinical practice?

► Increasing both the proportion of patients transferred by ambulance and improving identification of at-risk patients by paramedics are needed to optimise the known benefits of early and accurate STEMI identification.



© Author(s) (or their employer(s)) 2022. Re-use permitted under CC BY-NC. No commercial re-use. See rights and permissions. Published by BMJ.

For numbered affiliations see end of article.

Correspondence to

Dr Becky Yi-Wen Liao;
beckywliao@gmail.com

of the medical emergency call-outs in New Zealand (NZ). Virtually all emergency hospitalisations in NZ are admitted to public hospitals, including over 99% of acute coronary syndrome (ACS) admissions.⁹ In 2016, St John Ambulance implemented an electronic clinical record which contains information on patient clinical status and the paramedic initial clinical impression, including ‘suspected STEMI’ and ‘other suspected

cardiac'. Anonymised linkage of individual ambulance electronic records with public hospitalisation and mortality databases allows us to construct a comprehensive dataset of ambulance-to-hospital transfers, hospitalisations and outcomes for most of NZ. These linked datasets were used to investigate the utilisation of ambulance transport in STEMI cases, and to evaluate the concordance between the paramedic initial clinical impressions and the final hospital diagnoses. Our aims were to determine the proportion of patients with a final hospital diagnosis of STEMI who were transported by ambulance with a paramedic-'suspected STEMI', and to assess the impact of discordance between prehospital and final STEMI diagnoses on mortality outcomes.

METHOD

This national data linkage study included consecutive cases captured in each of the three health datasets between 01 July 2016 and 30 November 2018.

Terms used in this study

The definition of 'suspected STEMI' in St John records required both clinical impression and ECG suggestive of STEMI by paramedics. When their initial impression is ACS or other cardiac conditions but felt not to meet the criteria for STEMI, then it is classified as 'other suspected ACS'. The initial clinical impression is generally that of the paramedic prior to any electronic transmission of the ECG to an interventional centre. Remaining cases which do not meet either definition are classified as 'ACS not suspected'.

Datasets

The St John electronic record system collects information for all ambulance call-outs, including patient demographics, clinical indications, transport times, and prehospital assessment and management. The ambulance data were obtained for the whole of NZ except for patients transported to Wellington, Hutt and Wairarapa Hospitals because they use a different ambulance service (contributing only 10% of the NZ total).

The All NZ Acute Coronary Syndrome Quality Improvement (ANZACS-QI) registry has captured over 99% of NZ patients with ACS investigated with coronary angiography since 2015. The dataset and methodology of the ANZACS-QI registry have been previously reported, capturing ACS diagnosis types—STEMI, non-ST-elevated ACS (NSTEMI) and non-ACS diagnoses.¹⁰

The national administrative datasets include the National Minimum Dataset (NMDs) which provides International Classification of Diseases (ICD) 10AM-coded hospitalisation records, the Mortality Collection that provides mortality data, and the National Non-Admitted Patient Collection (NNAPC) which records patients presented to the emergency department who do not require hospitalisation and are discharged alive. NNAPC does not record ICD-coded diagnostic information. Online supplemental table 1 shows the ICD-10AM codes

used to define ACS diagnosis subtypes in the NMDs (STEMI, non-STEMI (NSTEMI), unstable angina (USA), myocardial infarction (MI) unspecified), and other coronary heart disease diagnoses. Both primary and secondary codes were used. Age, gender, ethnicity and an area-based index of socioeconomic deprivation (New Zealand Deprivation Index 2013 (NZDep13))¹¹ were obtained from the national datasets. Ethnicity was prioritised using a modified version of NZ Standard Ethnicity Protocol.^{12,11} Admission hospitals were classified as those participating in a routine primary percutaneous coronary intervention (PCI) service, those with a catheter laboratory but without routine primary PCI availability, or rural hospitals without a catheter laboratory. For STEMI cases, 'system time' was from ambulance dispatch to hospital arrival.

Dataset linkage and cohort construction

Every patient in contact with NZ health system is assigned a unique identifier, the National Health Index (NHI). The NHI can be used to anonymously link multiple electronic databases using a common encrypted NHI.¹⁰

Combined National Dataset: ACS hospitalisation episodes in the NMDs were defined using a previously validated 'bundling' method to account for interhospital transfers within a single ACS episode of care.⁹ For NMDs ACS bundles with more than one ACS diagnosis, ACS type was prioritised in the order STEMI>NSTEMI>USA>MI unspecified. When discrepancies arose between the final diagnoses in NMDs and ANZACS-QI, the latter was prioritised. The ANZACS-QI and NMDs admissions were linked and patients admitted to hospitals not in the St John catchment were excluded to form a Combined National Dataset.

St John dataset: from the raw St John dataset (789 466 records), duplicate entries and multiple call-outs of the same episode were excluded (n=25 865). A further 183 428 records were excluded as they were records of hospital transfers, prehospital deaths or for patients not transferred to hospital, where hospitalisation diagnostic data were not available. The final 580 173 St John acute hospital transports and 2 256 567 Combined National Dataset admissions were merged, resulting in a final cohort of 2 396 391 records for analysis (online supplemental figure 1).

Outcomes

The primary endpoint, 30-day post-admission all-cause mortality, was obtained for patients with a final STEMI diagnosis using the National Mortality dataset. These are reported for the first STEMI presentation per patient.

Statistics

Results were presented as mean with SD and/or median with IQR for continuous variables and frequency (percentage) for categorical variables. Comparison between groups (St John initial diagnosis or St John attendance) was done using Pearson's χ^2 test for

categorical data or non-parametric Mann-Whitney U test for continuous variables as the data were not normally distributed. The significance of the difference between the final diagnosis of STEMI according to the initial St John clinical impression and method of transport to hospital was tested using log-rank test. Unadjusted Kaplan-Meier curves showed the association between final diagnosis of STEMI according to initial St John clinical impression and method of transport to hospital, and mortality. The proportional hazard assumption was tested using SAS ASSESS statement in PROC PHREG and was met. Multivariable Cox regression models were used to estimate the adjusted hazard of ambulance paramedic 'other suspected ACS', 'ACS not suspected' and 'not attended by ambulance' compared with 'suspected STEMI' for 30-day mortality. Covariates included were age (continuous), gender, ethnicity, NZDep13, modified non-cardiac Charlson Comorbidity Index, coronary intervention status and hospital. All p values were two tailed and $p < 0.05$ was considered statistically significant. Data were analysed using SAS V.9.4 (SAS Institute), and survival plots were created using RStudio V.1.1.442.

Patient and public involvement

It was not appropriate to involve patients or the public in the design, conduct, reporting or dissemination plans of our research.

RESULTS

There were 5465 admissions with a final diagnosis of STEMI (table 1) and 3989 (73%) were transported to hospital by ambulance and 1486 (27%) self-presented. Of those transported by ambulance, 2016 (50.7%) had a prehospital clinical impression of suspected STEMI, 792 (19.9%) were suspected of another ACS diagnosis and 1181 (29.7%) were not suspected of an ACS.

For the 3557 cases where paramedics suspected a STEMI, the positive predictive value was 57%. Of the remaining cases, 16% were admitted with another coronary heart disease (CHD) diagnosis, predominantly NSTEMI. Another 21% were admitted with non-cardiac diagnosis and 7% were alive but did not require hospital

admission. Therefore, of all the cases suspected by paramedics of having STEMI, 72% had a final cardiac diagnosis.

Table 2 shows the characteristics of those cases with a final diagnosis of STEMI according to the initial paramedic impression and the mode of presentation. Cases identified as suspected STEMI by paramedics, as opposed to another diagnosis, were more likely to be younger and male. They also had less medical comorbidity when compared with those with a non-cardiac initial impression. Māori (45%) or Pacific people (42%) with STEMI were less likely to be initially identified as having a STEMI than Indian (57%), other Asian (58%) and European/other (52%) people. Only 36% of STEMI cases transported initially to rural hospitals without catheter laboratories were suspected STEMI prehospital compared with 53% transported to hospitals participating in a routine primary PCI service and 47% to hospitals with a catheter laboratory but without routine primary PCI. The mean time from first medical contact to hospital arrival was shorter when St John suspected a cardiac condition (54.6, 54.0 and 59.3 min for suspected STEMI, other ACS and non-ACS, respectively ($p < 0.001$)). Self-presentation to hospital, in contrast to ambulance transfer, was more common among older patients, men and non-European/other ethnic groups (Māori (36%), Pacific (31%), European/other (24%)).

Of the 3989 patients transported by ambulance with a final diagnosis STEMI, 2016 (51%) had an initial paramedic clinical impression of STEMI. As a consequence of this diagnostic sensitivity, combined with 73% of patients being transported by ambulance, only 37% of all STEMI cases were identified prehospital as a suspected STEMI (table 3). This proportion varied by age—younger patients, men and European/other patients with STEMI were most likely to be transported by ambulance with a clinically suspected STEMI (European/other 38.7% vs Māori and Pacific groups, 28.7% and 28.9%) as were those without multiple comorbidities.

In the 38 332 cases with an initial ambulance paramedic clinical impression of 'other suspected ACS', 792 (2.1%)

Table 1 Ambulance initial clinical impressions and the final diagnosis available in the Combined National Dataset for all cases

	Final diagnosis in the Combined National Dataset (column %)					Total
	STEMI	NSTEMI	Other CHD	Non-CHD	Not admitted	
Total	5465	28 304	14 862	2 184 334	163 426	2 396 391
Ambulance-suspected cardiac						
Suspected STEMI	2016 (36.9)	472 (1.7)	86 (0.6)	733 (0.1)	250 (0.2)	3557 (0.1)
Other suspected ACS	792 (14.5)	6876 (24.3)	2484 (16.7)	21 931 (1.0)	6249 (3.8)	38 332 (1.6)
Ambulance 'ACS not suspected'	1181 (21.6)	8785 (31.0)	3749 (25.2)	367 642 (16.8)	156 927 (96.0)	538 284 (22.5)
Not attended by ambulance	1476 (27.0)	12 171 (43.0)	8543 (57.5)	1 794 028 (82.1)	—	1 816 218 (75.8)

CHD, coronary heart disease; NSTEMI, non-ST-elevated acute coronary syndrome; STEMI, ST-segment elevation myocardial infarction.

Table 2 Characteristics of all cases with a STEMI diagnosis according to the ambulance service initial clinical impressions and method of transport to hospital

Ambulance initial clinical impression									
	Suspected STEMI		Other suspected ACS		ACS not suspected		All transported by ambulance		Self-presented
	(n=2016)	(n=792)	(n=792)	(n=792)	(n=1181)	P value (1 vs 2)	P value (1 vs 3)	P value (2 vs 3)	(n=1476)
Age (years), mean (SD)	66.4 (13.5)	68.4 (13.9)	68.4 (13.9)	68.4 (13.9)	72.7 (14.5)	0.001	<0.001	<0.001	64.3 (13.5)
Age group, n (%)						0.009	<0.001	<0.001	<0.001
<45	106 (5.3)	30 (3.8)	30 (3.8)	30 (3.8)	40 (3.4)				88 (6.0)
45–54	281 (13.9)	107 (13.5)	107 (13.5)	107 (13.5)	108 (9.1)				270 (18.3)
55–64	534 (26.5)	176 (22.2)	176 (22.2)	176 (22.2)	186 (15.7)				391 (26.5)
65–74	501 (24.9)	199 (25.1)	199 (25.1)	199 (25.1)	263 (22.3)				381 (25.8)
75–84	393 (19.5)	171 (21.6)	171 (21.6)	171 (21.6)	299 (25.3)				238 (16.1)
85+	201 (10.0)	109 (13.8)	109 (13.8)	109 (13.8)	285 (24.1)				108 (7.3)
Female, n (%)	634 (31.4)	271 (34.2)	271 (34.2)	271 (34.2)	514 (43.5)	0.158	<0.001	<0.001	440 (29.8)
Ethnicity, n (%)						0.971	<0.001	<0.001	<0.001
Māori	184 (9.1)	76 (9.6)	76 (9.6)	76 (9.6)	150 (12.7)				232 (15.7)
Pacific	80 (4.0)	29 (3.7)	29 (3.7)	29 (3.7)	83 (7.0)				85 (5.8)
Indian	86 (4.3)	35 (4.4)	35 (4.4)	35 (4.4)	30 (2.5)				84 (5.7)
Other Asian	79 (3.9)	28 (3.5)	28 (3.5)	28 (3.5)	29 (2.5)				74 (5.0)
European/other	1587 (78.7)	624 (78.8)	624 (78.8)	624 (78.8)	889 (75.3)				1001 (67.8)
NZDep13, n (%)						0.74	0.536	0.986	0.094
1–2 (least deprived)	327 (16.2)	124 (15.7)	124 (15.7)	124 (15.7)	173 (14.6)				224 (15.2)
3–4	375 (18.6)	135 (17.0)	135 (17.0)	135 (17.0)	194 (16.4)				251 (17.0)
5–6	385 (19.1)	140 (17.7)	140 (17.7)	140 (17.7)	210 (17.8)				232 (15.7)
7–8	452 (22.4)	164 (20.7)	164 (20.7)	164 (20.7)	248 (21.0)				291 (19.7)
9–10 (most deprived)	439 (21.8)	184 (23.2)	184 (23.2)	184 (23.2)	273 (23.1)				365 (24.7)
Missing	38 (1.9)	45 (5.7)	45 (5.7)	45 (5.7)	83 (7.0)				113 (7.7)
Non-cardiac Charlson score						<0.001	<0.001	<0.001	0.074
0	1551 (76.9)	541 (68.3)	541 (68.3)	541 (68.3)	619 (52.4)				1043 (70.7)
1–2	326 (16.2)	160 (20.2)	160 (20.2)	160 (20.2)	302 (25.6)				252 (17.1)
3+	139 (6.9)	91 (11.5)	91 (11.5)	91 (11.5)	260 (22.0)				181 (12.3)

Continued

Table 2 Continued

Ambulance initial clinical impression								
	Other suspected		ACS not suspected		All transported by			
	Suspected STEMI	ACS	ACS	P value	ambulance	Self-presented		
	(n=2016)	(n=792)	(n=1181)	(1 vs 2)	(1 vs 3)	(2 vs 3)	(n=1476)	P value
Hospital status, n (%)				<0.001	<0.001	0.404		0.892
Primary PCI centre	1408 (69.8)	506 (63.9)	735 (62.2)				2649 (66.4)	987 (66.9)
Non-primary PCI centre	551 (27.3)	241 (30.4)	389 (32.9)				1181 (29.6)	428 (29.0)
Rural hospital without cath lab	57 (2.8)	45 (5.7)	57 (4.8)				159 (4.0)	61 (4.1)
System time, median (IQR)	54.6 (42.1–74.0)	54.0 (40.8–71.5)	59.3 (44.3–80.7)	0.218	<0.001	<0.001	55.8 (42.4–75.9)	–
Coronary intervention status								
No cath				<0.001	<0.001	<0.001		0.512
Cath with intervention	270 (13.4)	177 (22.3)	627 (52.7)				1074 (26.9)	381 (24.7)
Cath without intervention	1574 (78.1)	514 (64.9)	417 (35.3)				2505 (62.8)	930 (64.3)
	172 (8.5)	101 (12.8)	137 (11.6)				410 (10.3)	165 (11.0)
ACS, acute coronary syndrome; cath, catheterisation; NZDep13, New Zealand Deprivation Index 2013; PCI, percutaneous coronary intervention; STEMI, ST-segment elevation myocardial infarction.								

Table 3 Proportion of patients with a final diagnosis of STEMI who were transported by ambulance with a paramedic-suspected STEMI

Final STEMIs transported by ambulance and suspected of STEMI prehospital/all final STEMI	
All	2016/5465 (36.9%)
Age group, years	
<45	106/264 (40.2%)
45–54	281/766 (36.7%)
55–64	534/1287 (41.5%)
65–74	501/1344 (37.3%)
75–84	393/1101 (35.7%)
85+	201/703 (28.6%)
Gender	
Male	1382/3606 (38.3%)
Female	634/1859 (34.1%)
Ethnicity	
Māori	184/642 (28.7%)
Pacific	80/277 (28.9%)
Indian	86/235 (36.6%)
Other Asian	79/210 (37.6%)
European/other	1587/4101 (38.7%)
NZDep13	
1–2 (least deprived)	327/848 (38.6%)
3–4	375/955 (39.3%)
5–6	385/967 (39.8%)
7–8	452/1155 (39.1%)
9–10 (most deprived)	439/1261 (34.8%)
Missing	38/279 (13.6%)
Hospital status	
Primary PCI centre	1408/3636 (38.7%)
Non-primary PCI centre	551/1609 (34.2%)
Rural hospital without cath lab	57/220 (25.9%)
Non-cardiac Charlson Index	
0	1551/3754 (41.3%)
1–2	326/1040 (31.3%)
3+	139/671 (20.7%)
Coronary intervention status	
No cath	270/1455 (18.6%)
Cath with intervention	1574/3435 (45.8%)
Cath without intervention	172/575 (29.9%)

cath, catheterisation; NZDep13, New Zealand Deprivation Index 2013; PCI, percutaneous coronary intervention; STEMI, ST-segment elevation myocardial infarction.

had a STEMI diagnosis and 18 145 (24.4%) had a diagnosis of NSTEMI or other CHD (table 1). There were 57.2% diagnosed with a non-ACS condition, and 16.3% who did not require admission and for whom no further diagnostic information is available.

Outcomes for STEMI cases according to mode of transport to hospital and the initial paramedic clinical impression are shown in table 4 and figure 1. Of the 5465 presentations with STEMI, 259 were repeat presentations which left 5206 (95.3%) first STEMI presentations for outcome analysis. The characteristics of this 'first presentation' cohort are similar to those shown in table 2 for all cases (online supplemental table 3).

The unadjusted 30-day mortality rates for patients with a final diagnosis of STEMI were similar for ambulance-suspected STEMI (9.0% (95% CI 7.8% to 10.3%)), suspected 'other ACS condition' (10.9% (95% CI 8.8% to 13.3%)) and for self-presenters (10.3% (95% CI 8.8% to 12.0%)). However, those transported by ambulance but not suspected of an ACS had the highest 30-day mortality rate (34.9% (95% CI 32.2% to 37.8%), $p<0.001$). After adjusting for all demographic and clinical covariates (table 4, Model C), 30-day mortality was lower for self-presenters with STEMI than for patients with STEMI identified by ambulance paramedics (HR 0.73, 95% CI 0.58 to 0.92). The risk for patients transported by ambulance but not initially suspected of an ACS was ameliorated but remained higher than for patients with STEMI identified by ambulance paramedics (HR 1.48, 1.22 to 1.80), and remained similar for ambulance-suspected 'other ACS conditions' (HR 0.85, 0.65 to 1.11).

DISCUSSION

St John Ambulance attends most of the call-outs in NZ and by combining its electronic dataset with the national datasets, we have a unique opportunity to assess both the accuracy of prehospital clinical impressions and their association with longer term outcomes. Only half of the patients with STEMI transported by ambulance were accurately identified and 20% were suspected of another ACS diagnosis, with the rest not initially thought to have an ACS. Over 60% of patients with STEMI self-presented or were not identified by paramedics and so missed out on opportunities for early reperfusion by activation of the cardiac catheterisation laboratory or receipt of prehospital fibrinolysis. The unadjusted all-cause mortality was similar for ambulance-transported patients suspected of either STEMI or another ACS condition, and in those who self-presented, but mortality was more than three times as high in those not suspected of an ACS. After adjustment for covariates, the excess mortality risk associated with not being suspected by paramedics of having a STEMI was ameliorated, but remained 1.5 times higher than for those initially identified as having a STEMI.

Table 4 Cox regression models of 30-day mortality for patients with a final STEMI diagnosis

Variables	Model A		Model B		Model C	
	HR (95% CI)	P value	HR (95% CI)	P value	HR (95% CI)	P value
Ambulance cardiac status						
Suspected STEMI (ref)	1	–	1	–	1	–
Other suspected ACS	1.22 (0.94 to 1.59)	0.14	1.11 (0.85 to 1.44)	0.45	0.85 (0.65 to 1.11)	0.22
ACS not suspected	4.41 (3.69 to 5.27)	<0.01	3.29 (2.74 to 3.95)	<0.01	1.48 (1.22 to 1.80)	<0.01
Not attended by ambulance	1.14 (0.92 to 1.42)	0.24	1.23 (0.99 to 1.54)	0.06	0.73 (0.58 to 0.92)	0.01
Demographics						
Age, per year	–	–	1.045 (1.039 to 1.051)	<0.01	1.015 (1.008 to 1.022)	<0.01
Sex	–	–				
Male			0.99 (0.86 to 1.15)	0.94	1.14 (0.98 to 1.32)	0.09
Female (ref)			1	–	1	–
Ethnicity	–	–	–	–		
Māori					1.09 (0.84 to 1.41)	0.53
Pacific					1.27 (0.92 to 1.76)	0.15
Indian					0.94 (0.58 to 1.53)	0.81
Asian					0.78 (0.48 to 1.27)	0.31
NZ Euro/other (ref)					1	–
NZDep13	–	–	–	–		
1–2 (ref)					1	–
3–4					1.09 (0.84 to 1.41)	0.51
5–6					1.24 (0.97 to 1.60)	0.09
7–8					1.27 (1.00 to 1.62)	0.05
9–10					1.33 (1.04 to 1.70)	0.02
Modified Charlson Comorbidity Index	–	–	–	–		
0 (ref)					1	–
1–2					1.01 (0.84 to 1.21)	0.91
≥3					1.24 (1.03 to 1.49)	0.02
Coronary intervention status	–	–	–	–		
No angiography (ref)					1	–
Angiography without intervention					0.25 (0.19 to 0.34)	<0.01
Angiography with intervention					0.12 (0.10 to 0.16)	<0.01
Hospital status	–	–	–	–		
Routine 24/7 PCI primary PCI					1.13 (0.96 to 1.33)	0.13
Base hospital w/o 24/7 PCI (ref)					1	–
Rural hospital					1.04 (0.76 to 1.44)	0.8

Model A: unadjusted.

Model B: adjusted by age and sex.

Model C: adjusted for age, sex, ethnicity, NZDep quintile, non-cardiac comorbidity (0, 1–2, >2), coronary intervention status (no angiography, angiography without intervention, angiography with intervention), hospital status (primary PCI, non-primary PCI with catheterisation lab, hospital without catheterisation lab).

ACS, acute coronary syndrome; NZDep13, New Zealand Deprivation Index 2013; PCI, percutaneous coronary intervention; STEMI, ST-segment elevation myocardial infarction.

Demographic differences

Māori (the indigenous population of NZ) and Pacific patients with a final hospital STEMI diagnosis were less likely to have been suspected of having a STEMI by

paramedics compared with other ethnic groups. They were also more likely to have self-presented to hospital rather than called an ambulance. As a result, Māori and Pacific groups are less likely to receive timely STEMI

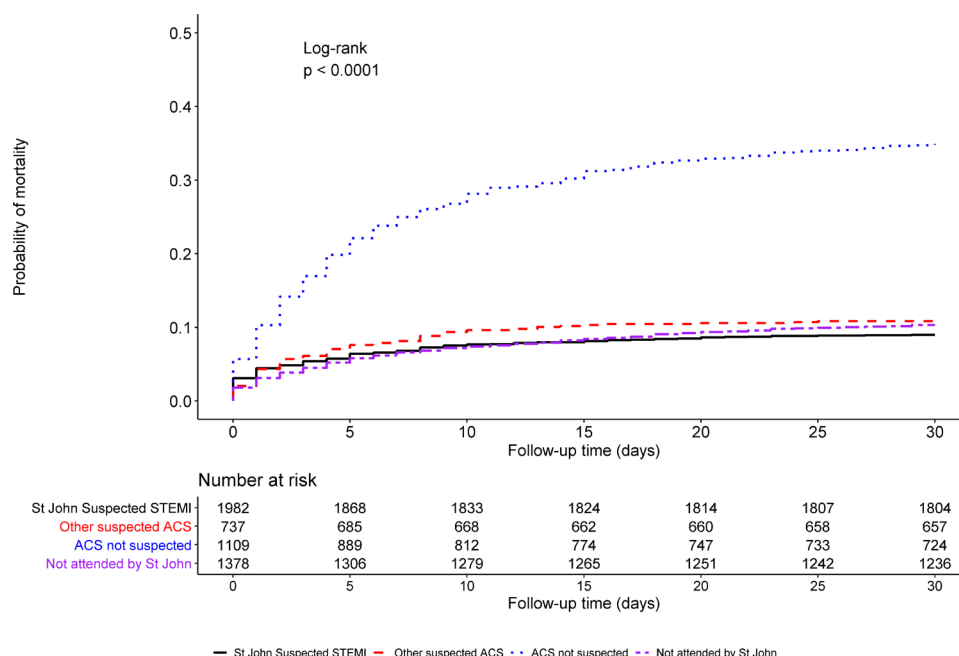


Figure 1 Unadjusted all-cause mortality for patients with STEMI* according to method of patient transport and initial ambulance paramedic clinical impression. *For patients with more than one STEMI presentation, only the first presentation was used in the outcomes analysis. ACS, acute coronary syndrome; STEMI, ST-segment elevation myocardial infarction.

therapy which is critical for improving clinical outcomes. This may contribute to their documented poorer outcomes post-MI.¹³

Patients with STEMI admitted to routine primary PCI-capable hospitals were more likely identified with STEMI by paramedics, compared with non-routine PCI-capable or rural hospitals (53%, 46% and 36%, respectively). While this may be due to factors such as paramedic training and appropriate utilisation of ECGs at the scene, the findings are also biased by patients with very clear STEMI in rural hospital catchments being transported directly to primary PCI-capable hospitals, leaving more equivocal cases going to rural hospitals. Prior analyses have not found differences in all-cause mortality based on the type of initial receiving hospital which may be due to appropriate identification and bypass of high-risk patients directly to primary PCI centres.¹⁴

Ambulance versus self-transport

Seventy-three per cent of the patients with STEMI were transported to hospital by ambulance. This compares favourably with international data. An observational analysis of 37 634 patients with STEMI treated at over 300 US hospitals found that emergency medical services were used for 59%–68% of patients.^{15–17} Nevertheless, in the current study, 27% of the STEMI cases were not attended by St John, a sizeable group without prehospital access to acute care and defibrillators. These patients will have longer time to reperfusion treatment as previously reported in the ANZACS-QI registry and from the REACT trial.^{18 19}

Paramedic diagnostic accuracy and implications

Prehospital clinical impression of STEMI was confirmed in 57% of cases. Prior studies have reported higher diagnostic accuracies for STEMI. One study of 354 patients had suspected ACS assessed by their mobile emergency care unit (ambulance with two emergency medical staff) at 87.5% of preliminary STEMI diagnoses confirmed.²⁰ Another Danish study similarly reported accuracy of 89%.²¹ A third study assessed 103 paramedics' responses to five different settings of chest pain and ECGs, finding 93% sensitivity and 85% specificity for paramedic diagnosis of STEMI.²² In contrast, another cohort study using paramedics' surveys reported a lower sensitivity 75% and specificity 53% for STEMI detection, including 96% correct for inferior STEMI but only 51% correct for lateral STEMI, and no correlation between paramedic training, experience and accuracy.²²

Outcomes

Patients with STEMI who were transferred by ambulance but not initially suspected of an ACS had over three times the mortality risk of those initially identified by paramedics as STEMI. The adverse outcomes are in part due to covariates including age and comorbidities; and after adjustment, the excess hazard was markedly reduced to just 1.5 times excess risk. Some of this residual excess risk may be related to unmeasured and therefore unadjusted comorbidity. In particular, the national datasets do not have information about clinical acuity at the time of presentation, and it is possible that some patients with STEMI without a clear STEMI diagnosis may be more unwell. We are unable to determine, in this observational

study, whether some of the remaining excess risk is due to the delay in appropriate monitoring and treatment for these patients. It is also possible that patients who call an ambulance may be sicker than those who self-present. This is supported by the finding that after adjustment, self-presenters had better outcomes than ambulance-transferred patients.

Clinical implications

Increasing both the proportion transferred by ambulance and improving identification of at-risk patients by paramedics may improve prehospital STEMI identification. This may facilitate earlier reperfusion therapy whether primary PCI or fibrinolysis. Consistent and increased utilisation of prehospital ECGs may help identify at-risk patients earlier. Ongoing training of paramedics on ECG interpretation and recognition of ACS-related symptoms is needed. The relatively low positive predictive value of 57% of the ambulance clinical impression of STEMI is of concern, and needs to be addressed as prehospital fibrinolysis programmes are rolled out, to avoid patient harm related to unnecessary fibrinolysis. Strategies to increase public awareness of ACS could include media campaigns, advanced cardiac life support community courses and school programmes, with goals of shortening the time to calling an ambulance and enhancing ambulance utilisation. The data linkage methodology used for this study will allow routine audit and ongoing quality improvement initiatives.

Limitations

Patients who died prehospital were not included in this study as final diagnosis data depended on hospital admission. Some St John records in the initial dataset did not have NHIs so could not be included in this study. Records with missing NHIs were predominantly for patients not transported to hospital (personal communication, Bridget Dicker). Some patients may have developed STEMI in hospital therefore could not have been identified by St John. However, using additional data available for nearly 80% of the patients with STEMI, we could estimate how many would have had a delayed diagnosis due to the ECG findings developing after admission. Exclusion of those patients would have increased the sensitivity of the St John clinical impression from 51% to 53%. Although the available data did not record the exact proportion of patients who did not receive a prehospital ECG but prior audit from St John Ambulance has shown 99% of suspected cardiac cases had a 12-lead ECG (personal communication, Bridget Dicker). Our study results were based on real-world national experience of suspected STEMI management in NZ, with our own unique patient population and healthcare systems, therefore findings are not necessarily generalisable to other countries and healthcare systems.

CONCLUSION

This large national data linkage study found over 6 out of 10 patients with STEMI potentially missed out on early reperfusion due to self-presentation to hospital or not being identified by paramedics as having a STEMI. Patients with STEMI transported by ambulance but not initially suspected of having an ACS had worse prognosis, in part explained by differences in baseline demographic and clinical factors.

Author affiliations

¹Department of Cardiology, Middlemore Hospital, Auckland, New Zealand

²Greenlane Cardiovascular Services, Auckland City Hospital, Auckland, New Zealand

³Paramedicine Research Unit, Paramedicine Department, School of Clinical Sciences, Auckland University of Technology, Auckland, New Zealand

⁴Clinical Audit and Research, St John New Zealand, Auckland, New Zealand

⁵Section of Epidemiology and Biostatistics, School of Population Health, University of Auckland, Auckland, New Zealand

⁶School of Medicine, University of Auckland, Auckland, New Zealand

Acknowledgements We would like to thank the Clinical Audit and Research team at St John and the Middlemore Cardiac Trust for their support of this study. ANZACS-QI programme implementation, coordination and analysis: the ANZACS-QI software was developed and supported by Enigma Solutions. Programme implementation is coordinated by the National Institute for Health Innovation (NIHI) at the University of Auckland. The ANZACS-QI programme is funded by the New Zealand Ministry of Health. We thank the National Health Board Analytic Services enabling use of the national datasets. We also thank the VIEW team at the School of Population Health, University of Auckland, for the curation and linkage of the national data. ANZACS-QI Governance Group: Andrew Kerr (chair), Dean Boddington, Gary Sutcliffe, Gerry Devlin, Harvey White, John Edmond, Jonathon Tisch, Kim Marshall, Mayanna Lund, Michael Williams (deputy chair), Nick Fisher, Seif El Jack and Sue Riddle. ANZACS-QI Project management: Kristin Sutherland (project manager), Charmaine Flynn (northern coordinator) and Maxine Rhodes (southern coordinator). Data analysis: Mildred Lee. Data management: Billy Wu (SOPH), Michelle Jenkins (NIHI) and John Faatui (NIHI). We acknowledge all the New Zealand cardiologists, physicians, nursing staff, radiographers and patients who have supported and contributed to ANZACS-QI.

Contributors Study design—BY-WL and AK. Data collection—all co-authors. Data analysis—BY-WL, MAWL and AK. Data interpretation—all co-authors. Manuscript writing—BY-WL and AK. Critical review of the manuscript and final approval of the manuscript—all co-authors. Guarantor—AK.

Funding This work was supported by the New Zealand Health Research Council, Wellington (grant number 11/800 to AK). KP is supported by a New Zealand Heart Foundation Hynds Senior Fellowship. BY-WL and MAWL are supported by the Middlemore Hospital Cardiology Trust.

Disclaimer Researchers are independent from funders, and the funders had no role in the study design, collection, analysis or interpretation of data.

Competing interests None declared.

Patient consent for publication Not required.

Ethics approval ANZACS-QI is part of the Vascular Informatics Using Epidemiology and the Web (VIEW) programme. The VIEW programme was approved by the Northern Region Ethics Committee Y in 2003 (AKY/03/12/314), with subsequent amendments to include the ANZACS-QI registries, and with annual approval by the National Multi-region Ethics Committee since 2007 (MEC07/19/EXP). The St John Ambulance service locality review approved the use of St John data in this study. All data were anonymised in the analyses so individual patient consent was not required. The corresponding author affirms that the manuscript is an honest, accurate and transparent account of the study being reported, that no important aspects of the study have been omitted.

Provenance and peer review Not commissioned; externally peer reviewed.

Data availability statement Data are available upon reasonable request. The data underlying this article cannot be shared publicly because they are made available to the investigators by the Ministry of Health under a Multi-region Ethics Committee

approval. The data will be shared on reasonable request to the corresponding author and approval by the VIEW Governance Group.

Open access This is an open access article distributed in accordance with the Creative Commons Attribution Non Commercial (CC BY-NC 4.0) license, which permits others to distribute, remix, adapt, build upon this work non-commercially, and license their derivative works on different terms, provided the original work is properly cited, appropriate credit is given, any changes made indicated, and the use is non-commercial. See: <http://creativecommons.org/licenses/by-nc/4.0/>.

ORCID iDs

Becky Yi-Wen Liao <http://orcid.org/0000-0001-9910-013X>

Bridget Dicker <http://orcid.org/0000-0002-8151-7356>

Ralph Stewart <http://orcid.org/0000-0002-6167-1225>

Katrina Poppe <http://orcid.org/0000-0002-4418-4476>

REFERENCES

- 1 Roffi M, Patrono C, Collet J-P, *et al.* 2015 ESC guidelines for the management of acute coronary syndromes in patients presenting without persistent ST-segment elevation: Task force for the management of acute coronary syndromes in patients presenting without persistent ST-segment elevation of the European Society of cardiology (ESC). *Eur Heart J* 2016;37:267–315.
- 2 Ibanez B, James S, Agewall S, *et al.* 2017 ESC guidelines for the management of acute myocardial infarction in patients presenting with ST-segment elevation: the task force for the management of acute myocardial infarction in patients presenting with ST-segment elevation of the European Society of cardiology (ESC). *Eur Heart J* 2018;39:119–77.
- 3 Levine GN, Bates ER, Blankenship JC. ACCF/AHA/SCAI guideline for percutaneous coronary intervention: a report of the American College of cardiology Foundation/American heart association Task force on practice guidelines and the Society for cardiovascular angiography and interventions. *Circulation* 2011;124:e574–651.
- 4 Amsterdam EA, Wenger NK, Brindis RG. ACC/AHA Task force members; Society for cardiovascular angiography and interventions and the Society of thoracic surgeons. 2014 AHA/ACC guideline for the management of patients with non-ST-elevation acute coronary syndromes: Executive summary: a report of the American College of Cardiology/American heart association Task force on practice guidelines. *Circulation* 2014;130:2354–94.
- 5 Johnston S, Brightwell R, Ziman M. Paramedics and pre-hospital management of acute myocardial infarction: diagnosis and reperfusion. *Emerg Med J* 2006;23:331–4.
- 6 Keeley EC, Hillis LD. Primary PCI for myocardial infarction with ST-segment elevation. *N Engl J Med* 2007;356:47–54.
- 7 Keeley EC, Boura JA, Grines CL. Primary angioplasty versus intravenous thrombolytic therapy for acute myocardial infarction: a quantitative review of 23 randomised trials. *Lancet* 2003;361:13–20.
- 8 McNamara RL, Wang Y, Herrin J, *et al.* Effect of door-to-balloon time on mortality in patients with ST-segment elevation myocardial infarction. *J Am Coll Cardiol* 2006;47:2180–6.
- 9 Kerr AJ, Lee M, Jiang Y, *et al.* High level of capture of coronary intervention and associated acute coronary syndromes in the all New Zealand acute coronary syndrome quality improvement cardiac registry and excellent agreement with national administrative datasets (ANZACS-QI 25). *N Z Med J* 2019;132:19–29.
- 10 Kerr A, Williams MJ, White H, *et al.* The all New Zealand acute coronary syndrome quality improvement programme: implementation, methodology and cohorts (ANZACS-QI 9). *N Z Med J* 2016;129:23–36.
- 11 Atkinson J, Salmond C, Crampton P. *NZDep2013 index of deprivation*. Wellington: Department of Public Health, University of Otago, 2014.
- 12 HISO 10001:2017 ethnicity data protocols, 2017. Available: <https://www.health.govt.nz/publication/hiso-100012017-ethnicity-data-protocols>
- 13 Mazengarb J, Grey C, Lee M, *et al.* Inequity in one-year mortality after first myocardial infarction in Māori and Pacific patients: how much is associated with differences in modifiable clinical risk factors? (ANZACS-QI 49). *N Z Med J* 2020;133:40–54.
- 14 Lee S, Miller R, Lee M, *et al.* Outcomes after ST-elevation myocardial infarction presentation to hospitals with or without a routine primary percutaneous coronary intervention service (ANZACS-QI 46). *N Z Med J* 2020;133:64–81.
- 15 Kaul P, Welsh RC, Liu W, *et al.* Temporal and provincial variation in ambulance use among patients who present to acute care hospitals with ST-elevation myocardial infarction. *Can J Cardiol* 2016;32:949–55.
- 16 So DYF, Ha ACT, Turek MA, *et al.* Comparison of mortality patterns in patients with ST-elevation myocardial infarction arriving by emergency medical services versus self-transport (from the prospective Ottawa Hospital STEMI registry). *Am J Cardiol* 2006;97:458–61.
- 17 Kerr A, Lee M, Grey C, *et al.* Acute reperfusion for ST-elevation myocardial infarction in New Zealand (2015–2017): patient and system delay (ANZACS-QI 29). *N Z Med J* 2019;132:41–59.
- 18 Shiomi H, Morimoto T, Kitaguchi S, *et al.* The ReACT Trial: Randomized Evaluation of Routine Follow-up Coronary Angiography After Percutaneous Coronary Intervention Trial. *JACC Cardiovasc Interv* 2017;10:109–17.
- 19 Hansen LH, Mikkelsen S. Ischaemic heart disease: accuracy of the prehospital diagnosis—a retrospective study. *Emerg Med Int* 2013;2013:754269.
- 20 Andersen MS, Nielsen TT, Christensen EF. A study of police operated dispatch to acute coronary syndrome cases arising from 112 emergency calls in Aarhus County, Denmark. *Emerg Med J* 2006;23:705–6.
- 21 Trivedi K, Schuur JD, Cone DC. Can paramedics read ST-segment elevation myocardial infarction on prehospital 12-lead electrocardiograms? *Prehosp Emerg Care* 2009;13:207–14.
- 22 Mencl F, Wilber S, Frey J, *et al.* Paramedic ability to recognize ST-segment elevation myocardial infarction on prehospital electrocardiograms. *Prehosp Emerg Care* 2013;17:203–10.

SUPPLEMENTARY APPENDIX

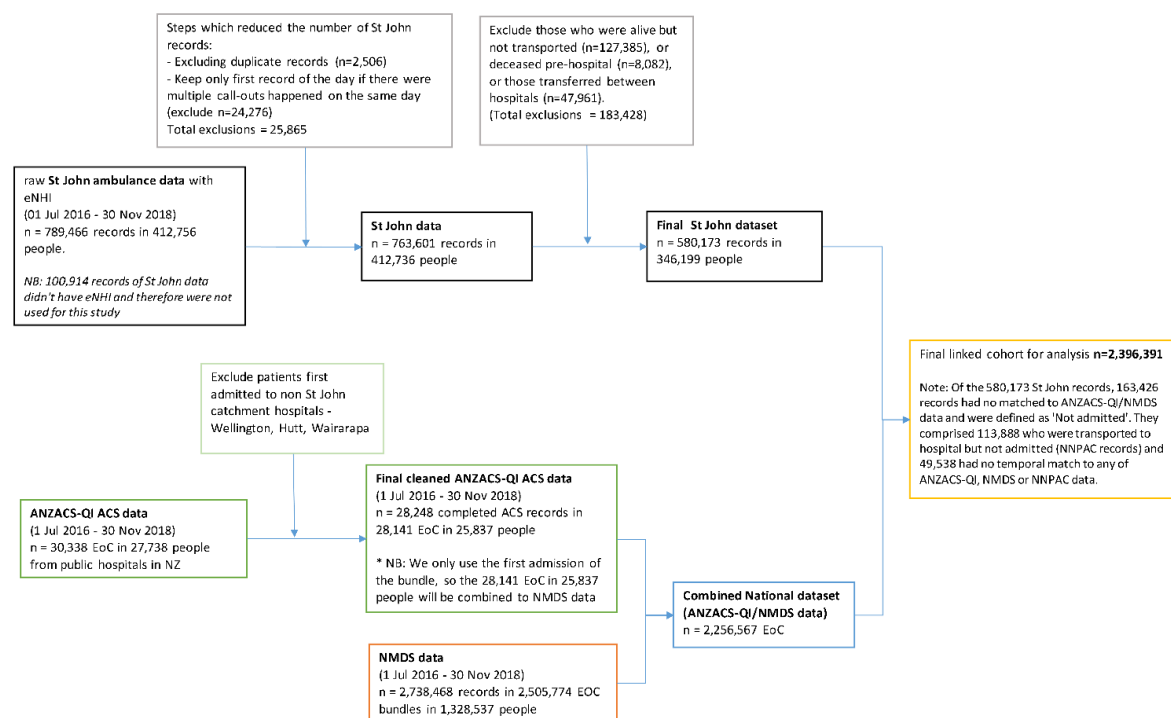
Table S1: List of ICD codes for ACS and other CHD

ACS Type	ICD codes	Description
STEMI	I210	Acute transmural myocardial infarction of anterior wall
	I211	Acute transmural myocardial infarction of inferior wall
	I212	Acute transmural myocardial infarction of other sites
	I213	Acute transmural myocardial infarction of unspecified site
	I220	Subsequent myocardial infarction of anterior wall
	I221	Subsequent myocardial infarction of inferior wall
	I228	Subsequent myocardial infarction of other sites
	I229	Subsequent myocardial infarction of unspecified site
NSTEMI	I214	Acute subendocardial myocardial infarction
	I222	Subsequent non-ST elevation (NSTEMI) myocardial infarction
USA	I200	Unstable angina
MI Unspecified	I219	Acute myocardial infarction, unspecified
Other CHD	I201	Angina pectoris with documented spasm
	I208	Other forms of angina pectoris
	I209	Angina pectoris, unspecified
	I230	Haemopericardium as current complication following acute myocardial infarction
		Atrial septal defect as current complication following acute

	I231	myocardial infarction
		Ventricular septal defect as current complication following
	I232	acute myocardial infarction
		Rupture of cardiac wall without haemopericardium as current
	I233	complication following acute MI
		Rupture of chordae tendineae as current complication following
	I234	acute myocardial infarction
		Rupture of papillary muscle as current complication following
	I235	acute myocardial infarction
		Thrombosis of atrium, auricular appendage, and ventricle as
	I236	current complications following acute MI
		Other current complications following acute myocardial
	I238	infarction
		Coronary thrombosis not resulting in myocardial infarction
	I240	Other forms of acute ischaemic heart disease
	I248	Acute ischaemic heart disease, unspecified
	I249	Aneurysm of heart
	I253	Coronary artery aneurysm
	I254	Ischaemic cardiomyopathy
	I255	Silent myocardial ischaemia
	I256	Cardiac arrest with successful resuscitation
	I460	Cardiac arrest, unspecified
	I469	

CHD: coronary heart disease

Appendix Figure 1: Flow chart summarising cohort construction



EOC, episode of care; NMDS, National minimum dataset; ACS, acute coronary syndrome

Appendix Table 2: Characteristics of patients with a first presentation with a STEMI diagnosis according to the initial St John clinical impressions and method of transport to hospital.

	St John suspected STEMI (n=1,982)	St John other suspected cardiac (n=737)	St John suspected non cardiac condition (n=1,109)	P value (1 vs 2)	P value (1 vs 3)	P value (2 vs 3)	Transported by St John (n=3,828)	Self presented (n=1,378)	P value
Age (years), Mean (SD)	66.3 (13.5)	68.4 (13.8)	72.6 (14.5)	<.001	<.001	<.001	68.5 (14.1)	64.3 (13.6)	<.001
Age group, n (%)									
<45	105 (5.3)	27 (3.7)	37 (3.3)				169 (4.4)	84 (6.1)	
45 -54	280 (14.1)	101 (13.7)	105 (9.5)				486 (12.7)	256 (18.6)	
55-64	525 (26.5)	160 (21.7)	176 (15.9)	0.004	<.001	<.001	861 (22.5)	365 (26.5)	<.001
65-74	493 (24.9)	187 (25.4)	245 (22.1)				925 (24.2)	347 (25.2)	
75-84	386 (19.5)	160 (21.7)	280 (25.3)				826 (21.6)	225 (16.3)	
85+	193 (9.7)	102 (13.8)	266 (24.0)				561 (14.7)	101 (7.3)	
Female, n (%)	620 (31.3)	254 (34.5)	479 (43.2)	0.114	<.001	<.001	1353 (35.3)	405 (29.4)	<.001
Ethnicity, n (%)									
Maori	178 (9.0)	69 (9.4)	137 (12.4)				384 (10.0)	217 (15.8)	
Pacific	76 (3.8)	25 (3.4)	79 (7.1)				180 (4.7)	81 (5.9)	
Indian	85 (4.3)	32 (4.3)	29 (2.6)	0.983	<.001	<.001	146 (3.8)	81 (5.9)	<.001
Other Asian	78 (3.9)	28 (3.8)	25 (2.3)				131 (3.4)	71 (5.2)	
European/Other	1565 (79.0)	583 (79.1)	839 (75.7)				2987 (78.0)	928 (67.3)	
NZ Dep13, n (%)									
1-2	324 (16.3)	124 (16.8)	172 (15.5)				620 (16.2)	221 (16.0)	
3-4	372 (18.8)	131 (17.8)	193 (17.4)				696 (18.2)	247 (17.9)	
5-6	381 (19.2)	137 (18.6)	209 (18.8)	0.817	0.570	0.969	727 (19.0)	231 (16.8)	0.085
7-8	448 (22.6)	162 (22.0)	247 (22.3)				857 (22.4)	287 (20.8)	
9-10	433 (21.8)	176 (23.9)	269 (24.3)				878 (22.9)	360 (26.1)	
Missing	24 (1.2)	7 (0.9)	19 (1.7)				50 (1.3)	33 (2.2)	

Hospital status, n (%)									
Primary PCI centre	1386 (69.9)	478 (64.9)	686 (61.9)				2550 (66.6)	928 (67.3)	
Non primary PCI centre	540 (27.3)	218 (29.6)	367 (33.1)	0.001	<.001	0.275	1125 (29.4)	393 (28.5)	0.822
Rural hospital without cathlab	56 (2.8)	41 (5.6)	56 (5.0)				153 (4.0)	57 (4.1)	
System time (from the first medical contact to arrival at hospital, mins)									
n	1972	734	1098	0.198	<.001	<.001	3804	-	-
Mean (SD)	62.6 (33.8)	60.1 (29.8)	67.6 (35.8)				63.6 (33.7)		
Median (IQR)	54.7 (41.9-74.0)	53.9 (40.7-71.6)	59.1 (44.3-80.9)				55.8 (42.4-76.0)		
Non-cardiac Charlson									
0	1546 (78.0)	520 (70.6)	595 (53.7)	<.001	<.001	<.001	2661 (69.5)	996 (72.3)	0.064
1-2	317 (16.0)	141 (19.1)	281 (25.3)				739 (19.3)	227 (16.5)	
3+	119 (6.0)	76 (10.3)	233 (21.0)				428(11.2)	155 (11.2)	
Coronary intervention status									
No Cath	208 (10.5)	137 (18.6)	556 (50.1)	<.001	<.001	<.001	901 (23.5)	309 (22.4)	0.170
Cath with intervention	1604 (80.9)	507 (68.8)	425 (38.3)				2536 (63.3)	904 (65.6)	
Cath without intervention	170 (8.6)	93 (12.6)	128 (11.5)				391 (10.2)	165 (12.0)	

NZDep13, New Zealand deprivation index 2013;PCI, percutaneous coronary intervention; Cath, catheterisation laboratory

Appendix Table 3: Proportion of patients with a final diagnosis of STEMI who were transported by ambulance with a paramedic suspected STEMI.

	Final STEMIs transported by ambulance / All final diagnoses of STEMI	Ambulance final STEMIs with suspected STEMI pre-hospital/ All ambulance final STEMI	Final STEMIs transported by ambulance and suspected of STEMI pre-hospital / all final STEMI
All	3989/5465 (73.0%)	2016/3989 (50.5%)	2016/5465 (36.9%)
Age group, years			
<45	176/264 (66.7%)	106/176 (60.2%)	106/264 (40.2%)
45-54	496/766 (64.8%)	281/496 (56.7%)	281/766 (36.7%)
55-64	896/1287 (69.6%)	534/896 (59.6%)	534/1287 (41.5%)
65-74	963/1344 (71.7%)	501/963 (52.0%)	501/1344 (37.3%)
75-84	863/1101 (78.4%)	393/863 (45.5%)	393/1101 (35.7%)
85+	595/703 (84.6%)	201/595 (33.8%)	201/703 (28.6%)
Gender			
Male	2570/3606 (71.3%)	1382/2570 (53.8%)	1382/3606 (38.3%)
Female	1419/1859 (76.3%)	634/1419 (44.7%)	634/1859 (34.1%)
Ethnicity			
Māori	410/642 (63.9%)	184/410 (44.9%)	184/642 (28.7%)
Pacific	192/277 (69.3%)	80/192 (41.7%)	80/277 (28.9%)
Indian	151/235 (64.3%)	86/151 (57.0%)	86/235 (36.6%)
Other Asian	136/210 (64.8%)	79/136 (58.1%)	79/210 (37.6%)
European/Other	3100/4101 (75.6%)	1587/3100 (51.2%)	1587/4101 (38.7%)
NZ Dep13			
1-2 (least deprived)	624/848 (73.6%)	327/624 (52.4%)	327/848 (38.6%)
3-4	704/955 (73.7%)	375/704 (53.3%)	375/955 (39.3%)
5-6	735/967 (76.0%)	385/735 (52.4%)	385/967 (39.8%)
7-8	864/1155 (74.8%)	452/864 (52.3%)	452/1155 (39.1%)
9-10 (most deprived)	896/1261 (71.1%)	439/896 (49.0%)	439/1261 (34.8%)
Missing	166/279 (59.5%)	38/166 (22.9%)	38/279 (13.6%)
Hospital status			
Primary PCI centre	2649/3636 (72.9%)	1408/2649 (53.2%)	1408/3636 (38.7%)
Non primary PCI centre	1181/1609 (73.4%)	551/1181 (46.7%)	551/1609 (34.2%)
Rural hospital without	159/220 (72.3%)	57/159 (35.8%)	57/220 (25.9%)

cathlab			
Non-cardiac Charlson index			
0	2711/3754 (72.2%)	1551/2711 (57.2%)	1551/3754 (41.3%)
1-2	788/1040 (75.8%)	326/788 (41.4%)	326/1040 (31.3%)
3+	490/671 (73.0%)	139/490 (28.4%)	139/671 (20.7%)
Coronary intervention status			
No Cath	1074/1455 (73.8%)	270/1074 (25.1%)	270/1455 (18.6%)
Cath with intervention	2505/3435 (72.9%)	1574/2505 (62.8%)	1574/3435 (45.8%)
Cath without intervention	410/575 (71.3%)	172/410 (42.0%)	172/575 (29.9%)