

National Cardiovascular Disease Database (NCVD)

Inaugural

**Report of the
Acute Coronary Syndrome (ACS) Registry**

2006

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FOREWORD

I extend my heartiest congratulations to the National Cardiovascular Disease Database (NCVD) team for recognising the critical status of Acute Coronary Syndrome in Malaysia through the production and publication of this inaugural NCVD ACS Registry Report Year 2006.

This report provides an estimation of acute coronary syndrome cases and describes the provision of acute coronary care services in our country. The data presented here details the trend of acute coronary syndrome and current practices related to acute coronary care services. As the report comprehensively covers such vital information, it can easily be utilised by cardiologists and physicians to improve patient outcomes and be used as reference material for policymakers to plan better cardiac care services in the future.

I would like to commend the Clinical Research Centre, the National Institute of Health, Malaysia and the National Heart Association of Malaysia for coordinating and supporting this registry, an important endeavour in our strive to create awareness of acute coronary syndrome and the importance of its treatment and management.

I wish you all the best and thank you for this effort.

Y. Bhg. Tan Sri Dato' Seri Dr. Hj Mohd Ismail Merican

Director-General of Health Malaysia

Ministry of Health Malaysia



AMERICAN COLLEGE OF CARDIOLOGY
Heart House, 2400 N Street NW, Washington DC, 20037

On behalf of the American College of Cardiology (ACC), I extend my sincerest congratulations for the inaugural report of the Malaysian Acute Coronary Syndrome Registry.

There have been tremendous advances in ACS treatment over the last 25 years. Despite these advances, however, it is clear that evidence is unevenly translated into clinical practice. In virtually all countries and practice settings where it has been measured, there are significant gaps between guideline recommendations and actual clinical care. In the United States (US), the Institute of Medicine has said there is a “quality chasm” in healthcare, and has strongly advocated the need to make healthcare delivery more effective, safe, equitable, timely, efficient, and patient-centered.

To achieve these aims of higher quality of care, a critical first step is to measure how care is currently being delivered. Only in this way, can gaps in care delivery be identified and targeted quality improvement interventions are implemented. In this way, quality improvement itself becomes “evidence-based”, and is more likely to lead to substantive improvements in patient outcomes.

With this context, the Malaysian Acute Coronary Syndrome Registry and National Cardiovascular Disease Database (NCVD) are a tremendous accomplishment. This is a world-leading effort that can demonstrate the importance of clinical registries as a facilitator of improving quality of care and patient outcomes. The data reported here can and should lead to targeted local and national quality improvement efforts for ACS care. The ongoing existence of the registry will allow tracking of the impact of these quality improvement efforts on processes of care and patient outcomes. The inclusion of longitudinal care and patient outcomes in the NCVD is a laudable goal and places this effort ahead of almost all others in the world.

The American College of Cardiology National Cardiovascular Data Registry (NCDR) currently includes an ACS registry as well as four other quality improvement registry programs in the United States. The mission of the NCDR is to improve the quality of cardiovascular patient care by providing information, knowledge and tools, implementing quality initiatives; and supporting research that improves patient care and outcomes.

This is clearly a shared mission with the Malaysian Acute Coronary Syndrome Registry/NCVD. As such, I hope that we can work collaboratively to show the world how to optimally measure and improve the quality of care and outcomes of patients with cardiovascular disease.

Sincerely,

John S. Rumsfeld, MD PhD FACC FAHA

Chief Science Officer and Chairman of the Management Board,
American College of Cardiology National Cardiovascular
Data Registry (NCDR)
Email: jrumfel@acc.org



To

Professor Dr. SIM Kui-Hian
President 2008-2010
National Heart Association of Malaysia (NHAM)

Ludwigshafen / Germany, July 18, 2008

Dear Professor Sim,

Thank you very much for providing to me the Report of the Malaysian Acute Coronary Syndrome (ACS) Registry which was started in 2006 with the support of the Ministry of Health and the National Heart Association of Malaysia (NHAM).

I congratulate you on this important contribution providing new insights into clinical practice of current ACS management in Malaysia. In contrast to randomized controlled trials (RCT), which consider highly selected patients treated in specialized centers, prospective registries document the current status of treatment of consecutive patients in clinical practice. They therefore provide information about patients, who usually are excluded from RCT, e.g. women, older patients, high risk patients or patients with multiple concomitant diseases. Your comprehensive report for the first time provides important data on the clinical presentation of ACS in Malaysia, on acute treatment in daily practice as well as on hospital and 1-year outcome.

Overall, there are a lot of similarities to data presented in European ACS registries, but also very specific differences especially with respect to the patient population, which in Malaysia is significantly younger and does have a different risk profile. These differences in baseline characteristics already might influence treatment decisions as well as outcome. This is only one of many arguments which underline the importance of specific Malaysian data.

The mission of the ESC: to improve the quality of life of the European population by reducing the impact of cardiovascular disease

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Most importantly, prospective registries play a key role in quality assurance in cardiovascular medicine. They help identifying gaps between treatment in “real life” and recommendations of guidelines and even are a feasible and effective tool for quality assessment and for improving adherence to existing treatment guidelines.

I very much would like to encourage you to continue this important project in the future.

Yours sincerely,

Anselm K. Gitt, MD, FESC
Chairman ESC Euro Heart Survey Programme 2004-2008

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ACKNOWLEDGEMENTS

The National Cardiovascular Disease Database (NCVD) would like to give its appreciation to everyone who has helped make this report possible.

We would especially like to thank the following:

- Our source data providers, cardiologists and physicians working in the participating sites
- Clinical Research Centre, Ministry of Health
- Ministry of Health Malaysia
- The members of various expert panels

PREFACE

In this day and age, the practice of medicine in Malaysia and internationally demand evidence based data and consideration of health economics. Although cardiovascular (CV) disease is one of the top mortalities in Malaysia, we only have minimal data on many aspects of the CV disease in our country.

Under the 9th Malaysia Plan, CV disease is listed as one of the top 8 diseases for priority research in Malaysia. The NCVD - ACS Registry (2006) and NCVD - PCI Registry (2007) were started with grants from the Ministry of Health Malaysia (MOH) Malaysia as well as the National Heart Association of Malaysia (NHAM).

We would like to express our heart-felt thanks to MOH, NHAM, members of the governance board, chairs for NCVD - ACS Registries and the committee, chair of NCVD publication and the committee, CRC HKL project management team and last but not least, the many investigators and the research nurses across the participating hospitals in the country who sacrificed hours of labour without financial remuneration to turn a dream into reality.

We would like to dedicate our own first NCVD - ACS Registry report to everyone who has contributed to this registry.

Lastly, we look forward to 100% participation across the country in order to build a truly “national” cardiovascular database.

Prof. Dr. Sim Kui-Hian
Co-Chairman
NCVD Governance Board

Dato' Seri Dr. Robaayah Zambahari
Co-Chairman
NCVD Governance Board

FOREWORD

As Co-Chairs of NCVD Acute Coronary Syndrome (ACS), Dato' Dr. Jeyaindran Sinnadurai (HKL) and I would like to present the first report of the NCVD ACS database.

This report is an analysis of ACS admissions in 11 hospitals involving 3,422 patients in year 2006 and is the cooperative effort of hospitals under the Ministry of Health Malaysia (MOH), National Heart Institute (IJN) and University Malaya Medical Centre (UMMC).

The registry will be expanded later to involve all general hospitals nationwide. It is designed to analyze the characteristics of patients presenting with acute coronary syndromes and their progress in hospital until discharge as well as outcome until the first year of treatment. This ACS database is to compliment the percutaneous coronary intervention (PCI) database. This report has been the collaborative effort of many individuals, hospitals and their staff. It is made possible by funds from the MOH and the National Heart Association of Malaysia (NHAM) as co-sponsors.

On behalf of the NCVD ACS, we would like to express our sincere appreciation and thanks for the untiring efforts of all those who were involved in this registry including the hospital staff concerned. We also would like to thank the hospitals that responded to the CCU survey. We hope that this will be the start of a more comprehensive and ongoing database that will enable us to improve the quality of cardiovascular care in our country.

Dato' Dr. Jeyaindran Sinnadurai
Co-Chairman
NCVD-ACS Registry

Dato' Dr. Azhari Rosman
Co-Chairman
NCVD-ACS Registry

ABBREVIATIONS

ACE	Angiotensin Converting Enzyme
ACS	Acute Coronary Syndrome
ADP	Adenosine Diphosphate
CABG	Coronary Artery Bypass Graft
CAD	Coronary Artery Disease
CCU	Coronary Care Unit
CK	Creatinine Kinase
CK-MB	Creatinine Kinase, Muscle and Brain
CPG	Clinical Practice Guidelines
CRC	Clinical Research Centre
CRF	Case Report Form
CRW	Cardiac Rehabilitation Ward
CV	Cardiovascular
CVD	Cardiovascular disease
DBMS	Database Management System
EDC	Electronic Data Capture
GP	Glycoprotein
HDL	High Density Lipoprotein
ICCU	Intensive Coronary Care Unit
ICT	Information and Communication Technology
ICU	Intensive Care Unit
IT/IS	Information Technology and Information System
LDL	Low Density Lipoprotein
LMWH	Low Molecular Weight Heparin
LVEF	Left Ventricular Ejection Fraction
JPN	Jabatan Pendaftaran Negara
MOH	Ministry of Health
NCVD	National Cardiovascular Disease Database
NSTEMI	Non ST-segment elevation Myocardial Infarction
PCI	Percutaneous Coronary Intervention
PMP	Per Million Population
SAP	Statistical Analysis Plan
SC	Site Coordinator
SD	Standard Deviation
SDP	Source Data providers
SME	Subject Matter Expert
STEMI	ST-segment elevation Myocardial Infarction
TIMI	Thrombolysis In Myocardial Infarction
T n I	Troponin I
T n T	Troponin T
UA	Unstable angina

ABOUT NCVD

Introduction

The National Cardiovascular Database (NCVD) is a service supported by the Ministry of Health Malaysia (MOH) to collect information about cardiovascular disease in Malaysia, which will enable us to know the incidence of cardiovascular disease, and to evaluate its risk factors and treatment in the country. This information is useful in assisting the MOH, Non-Governmental Organizations, private healthcare providers and industry in programme planning and evaluation, leading to cardiovascular disease prevention and control.

The NCVD is established to integrate various existing databases in individual hospitals either in MOH hospitals or private and other data sources to achieve a nation-wide cardiovascular database.

Rationale for Acute Coronary Syndrome (ACS) registry

Several important issues arise when applying the rigorous standards and protocols from clinical trials into real-life practice which include questions like:

- Are the population and the patient groups in Malaysia similar to those being investigated in the clinical trials?
- Are our hospitals following the guidelines set out by the expert committees?
- Are we seeing the same results and benefits of implementing evidence-based strategies?
- Which strategy will be the best value in terms of cost-effectiveness for the Ministry?

Furthermore, at present much of what we understand about risk and likelihood of CV disease and indeed its incidence and prevalence are derived from 'Western' data. There is now an increasing awareness of ethnic variations and risk, socio-cultural and socio-economic influences as well as geographical variations. The risk prediction of ACS is also unclear and may be different from CV disease patients with chronic stable angina.

These are the reasons why a registry is needed, as it provides the real-life data that would represent the population.

Acute Coronary Syndrome (ACS) Registry

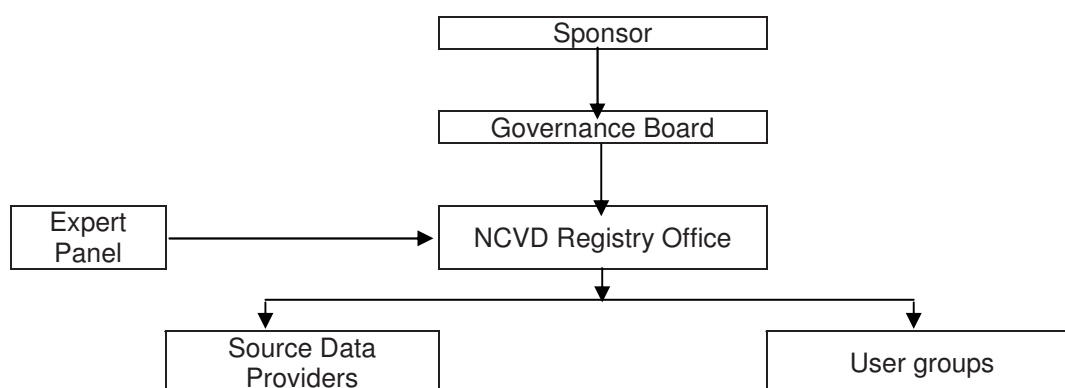
The NCVD-ACS registry is the first stage in realizing the rationale of a nation wide cardiovascular database. The ACS registry was officially launched on 31st March 2006.

The objectives are to:

- (i) Determine the number and the time trend of acute coronary syndromes in Malaysia.
- (ii) Determine the socio-demographic profiles of these patients to better identify high-risk groups in our Malaysian population.
- (iii) Determine the efficiency of and adherence to current guidelines of treatment guidelines
- (iv) Determine the cost to the nation of cardiovascular disease and the cost effectiveness of treatment and prevention programmes
- (v) Stimulate and facilitate research of cardiovascular disease using this database.

This report has met the first and second objectives. The rest of the objectives will be covered in future publications.

Organization of NCVD ACS registry



Sponsor

The NCVD is sponsored by the Ministry of Health Malaysia (MOH) and co-sponsored by the National Heart Association of Malaysia (NHAM). The registry has also been supported by several MOH organizations:

- Cardiology and Medicine Departments
- Clinical Research Centre

Governance Board

The Governance Board was established in year 2006 by sponsors to oversee operations of the National Cardiovascular Disease Database. The MOH, universities, professional bodies, NGOs and private healthcare providers are represented in this committee to ensure that the NCVD stays focused on its objectives, its continuing relevance and justification.

Current membership of the board is as follows:

Name	Organization
Prof Dr Sim Kui-Hian (Co-Chairman)	Head, Department of Cardiology & Clinical Research Centre, Sarawak General Hospital
Dato' Seri Dr Robaayah Zambahari (Co-Chairman)	Medical Director, National Heart Institute
Dato' Dr Omar Ismail	Head, Department of Cardiology, Penang Hospital
Dato' Dr Jeyaindran Sinnadurai	Consultant Pulmonary & Critical Care Physician, Department of Medicine, Kuala Lumpur Hospital
Prof Dr Wan Azman Wan Ahmad	Head, Department of Medicine, University Malaya Medical Centre
Dato' Dr Zaki Morad B Mohd Zaher	Professor, School of Medicine, International Medical University (IMU)
Dr Zainal Ariffin Omar	Deputy Director, Non-communicable Disease Section, MOH
Dato' Dr K Chandran	Head, Department of Medicine, Ipoh Hospital
Dato' Dr Haji Sapari Satwi	Head, Department of Medicine, Tengku Ampuan Afzan Hospital
Dr Lim Teck Onn	Director, Clinical Research Centre
Dr Hendrick M. Y. Chia	President, National Heart Association of Malaysia (2006-2008)
Dato' Dr Azhari Rosman	Hon. Secretary, National Heart Association of Malaysia (2006-2008)
Dato' Dr Khoo Kah Lin	Director, National Heart Foundation of Malaysia
Prof Dr Abdul Rashid Abdul Rahman	Professor, Faculty of Pharmacy, Cyberjaya University College of Medical Sciences

Steering Committee

The steering committee comprises individuals who are subject matter experts drawn from the various centres that are involved in the MOH, universities and private hospitals. They are convened to decide on the initial data collection process, develop the pro forma and data content as well as guide future development. They ensure that the database has a sound technical as well as scientific basis.

The role of the steering committee is to:

- Establish policy and procedures for the registry's conduct
- Motivate source data providers (SDP) to continue participation in the registry
- Disseminate information about the registry
- Communicate results locally and internationally.
- Approve, and if necessary validate, the statistical analysis plan,
- Undertake Quality Control of the reported data
- Determine policy and procedures for the operations of the database.
- Establish the Registry Coordinating Centre and appoint its project team members
- Direct the activities of the Registry Coordinating Centre

The current membership of steering committee is as follows:

Name	Organization
Dato' Dr Jeyaindran Sinnadurai (Co-Chairman)	Kuala Lumpur Hospital
Dato' Dr Azhari Rosman (Co-Chairman)	National Heart Institute
Assoc. Prof Dr Chin Sze Piau (Secretary)	International Medical University (IMU)
Dr Liew Chee Tat	Penang Hospital
Dr Ang Choon Kiat	Sarawak General Hospital
Dr Chong Wei Peng	University Malaya Medical Centre
Dr Lu Hou Tee	Sultanah Aminah Hospital

NCVD: Registry Coordinating Centre

Clinical Registry Manager	Ms S Gunavathy Selvaraj
Clinical Registry Associate	Ms Noor Amirah Muhamad

Supporting Staff from the Clinical Research Centre

The Clinical Research Centre (CRC) of the Ministry of Health provide technical support for the NCVD-ACS Registry. The clinical epidemiologists provide methodological and epidemiological inputs while the database is supported on CRC's IT infrastructure.

Clinical Epidemiologist	Dr Jamaiah Haniff Dr Anita Das Ms S Gunavathy Selvaraj
Statistician	Mr Muhammad Adam
ICT Manager	Madam Celine Tsai Pao Chien
Database Administrator	Ms Lim Jie Ying
Application Developer	Ms Amy R. Porle
Network Administrator	Mr Kevin Ng Hong Heng Mr Adlan Ab. Rahman
Webmaster & Desktop Publisher	Ms Azizah Alimat
Clinical Data Manager	Ms Teo Jau Shya

Biostatistics Consultants

Dr Sharon Chen Won Sun
Dr Hoo Ling Ping
Mr Tan Wei Hao
Ms Jasmine Chew
Ms Norhafizah Bt. Ab. Manan

Medical Writing Committee

A Committee has been constituted to prepare the registry regular or interim report, and to prepare the manuscript for journal submission for a particular study based on registry data.

The current members of medical writing committee of ACS registry are as follows:

Name	Organization
Prof Dr Wan Azman Wan Ahmad (Chairman)	University Malaya Medical Centre (UMMC)
Dato' Dr Azhari Rosman	National Heart Institute
Assoc. Prof Dr Chin Sze Piaw	International Medical University (IMU)
Dr Chong Wei Peng	University Malaya Medical Centre (UMMC)
Dr Alan Fong Yean Yip	Sarawak General Hospital
Dr Saravanan Krishnan	Sultanah Aminah Hospital
Dr Hazlyna Kamaruddin	National Heart Institute

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INTRODUCTION

Acute coronary syndrome (ACS) remains an important cause of death and hospitalization in Malaysia. It is a clinical spectrum of ischaemic heart disease ranging from unstable angina (UA), non ST-elevation myocardial infarction (NSTEMI) to ST-elevation myocardial infarction (STEMI) depending upon the degree and acuteness of coronary occlusion. The most common cause is the reduced myocardial perfusion due to atherosclerotic plaque rupture, fissuring or ulceration with superimposed thrombosis and coronary vasospasm.

In STEMI there is myocardial necrosis following an acute total coronary occlusion. UA and NSTEMI are considered to be closely related conditions where the coronary artery narrowing is non-occlusive. The clinical presentations are similar, but they differ in severity and whether the ischemia is severe enough to cause sufficient myocardial damage that result in significant elevation of cardiac biomarkers.

Much progress has been made in the management of ACS especially in the last two decades. Cardiovascular medicine is the most evidence-based medicine, thanks to the many good large randomized multi-centre clinical trials that involved new drugs, new treatment modalities and different treatment strategies. To further improve the management of ACS many expert bodies in the world have come up with guidelines in the management of this condition. In Malaysia we have 2 Clinical Practice Guidelines (CPG) for ACS i.e. CPG on Management of Acute ST-Segment Elevation Myocardial Infarction (STEMI) 2007- (2nd Edition) and CPG on UA/NSTEMI 2002.

The update of this second guideline is in the pipeline. To record how we manage this condition in the real world, we have created this registry with the help of many expert bodies. We are very grateful for every assistance and take great pride in publishing our own registry. This is our registry and it belongs to all of us and for the first time we have our own data to refer to at national and international meetings.

This is the fruit of your hard work and it was done in the spirit of 'Malaysia Boleh' and we would like to acknowledge all those who have made this report possible. This event will be another milestone in the history of cardiovascular medicine in Malaysia.

The report is divided into 5 chapters and after each chapter important summary points will be highlighted.

Chapter 1: Provision of Acute Coronary Care Services in Malaysia

Chapter 2: Patient Characteristics

Chapter 3: Clinical Presentations and Investigation

Chapter 4: Treatment

Chapter 5: Outcome

CHAPTER 1

PROVISION OF ACUTE CORONARY CARE SERVICES IN MALAYSIA

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In 2006, there were a total of 31186 admissions to the 73 coronary care units (CCU) in Malaysia, of which 12534 admissions were due to Acute Coronary Syndrome (ACS) (Table 1). The incidence of ACS admission was therefore 47.1 per 100,000 population in 2006. Assuming half of all coronary heart disease (CHD) first presented with ACS and only half were admitted to CCU with a third who died before being admitted into hospital, a rough estimate of the incidence of CHD in Malaysia is 141 per 100,000 population.

The 37 CCUs in MOH hospitals took care of the majority (60%) of ACS admissions, the 3 university hospitals' CCUs cared for another 10% while the private sector accounted for 27% of admissions. As expected, the economically developed states like Penang, Perak, Selangor/Wilayah Persekutuan have disproportionately large numbers of ACS admissions, while the less developed states (Kedah/Perlis, Terengganu, Sabah and Sarawak) were under-resourced, the surprising exception being Kelantan. Thus, the pattern parallels the availability of acute coronary care services in these states.

The 73 CCUs in the country provided 414 CCU beds. Table 1.2 shows the utilization of these resources. The MOH sector is clearly under-resourced relative to the demands it faces, resulting in over 4000 ACS being denied admission into its CCU in 2006. It has a 30% shortfall in CCU beds, and even if non-acute cardiac admissions are excluded, it is still short of 10% of its required bed strength.

Table 1.3 shows the cardiac care ACS patients received after being admitted into CCU. A remarkable 59% of patients in IJN had thrombolytic. Patients in private hospitals, including IJN, are more likely to receive invasive coronary interventions (emergency angiogram, PCI and coronary artery bypass graft [CABG]).

Figure 1.1 and 1.2 shows that the likelihood of ACS patients receiving emergency angiogram, PCI and CABG are driven by availability of cardiologist and on-site invasive cardiac catheterization laboratory (cath lab) facility.

Similarly, the likelihood of receiving emergency coronary artery bypass graft (CABG) correlates with availability of cardiac surgical services (Figure 1.3). What is the optimum level for the provision of these services however remains to be determined, and whether availability of such emergency services translated into better health outcomes are addressed in another chapter in this report.

Summary Points:

- The incidence of ACS admission to CCU was 47 per 100,000 populations in 2006.
- MOH Hospitals received 60% of ACS while Private Hospitals account for only 27%. However the MOH sector is clearly under-resourced in terms of CCU beds, on-site Cardiologists, Catheterization Laboratory and Cardiac Surgical Facilities.
- The likelihood of ACS patients receiving intervention (PCI or CABG) is driven by the availability of these resources.

Table 1.1 Acute Coronary/ Cardiac Care Services and Admissions in Malaysia 2006

	Populati on	CCU		Total beds		CCU beds		CCU nurses		Cardiolo gist		Cath lab		Cardiac surgical service		CCU admission		All acute admission		ACS admission		
		No	(%)	No	(%)	No	(%)	No	(%)	No	(%)	No	(%)	No	(%)	No	(%)	No	(%)	No	(%)	
Malaysia	26.6	73	30191	414	559	163	6	36	1	31186	20144	756	12534	471								
By sector																						
Private		32	6357	214	176	81		28		13820	6286		3398									
MOH		44	31	52	31	50		78		44	31		27									
		37	21118	169	305	37		4		14257	11809		7580									
		51	41	55	23	23		11		46	59		60									
IJUN		1	211	12	13	27		1		1307	465		335									
		1	3	2	17	4		3		4	2		3									
University		3	2505	19	65	17		3		1802	1583		1221									
		4	5	12	10	10		8		6	8		10									
States																						
P.Pinang	1.5	9	3206	54	63	24	16	6	4	3	6124	2242	1502	1140	764							
		12	11	13	11	15		17		20	11		9									
Melaka	0.7	4	1485	31	15	11	15	3	4	4	1961	1243	1714	745	1027							
		5	5	7	3	7		8		6	6		6									
Johor	3.2	5	2293	30	34	9	3	2	1	1	2846	1938	611	1243	392							
		7	8	7	6	6		6		9	10		10									
Perak	2.3	7	2829	25	58	4	2	1	0	1	2635	2253	987	1772	776							
		10	9	6	10	2		3		8	11		14									
Selangor & Kuala Lumpur	6.4	22	10,118	148	220	86	13	15	2		9211	5634	876	3683	573							
		30	34	36	39	53		42		30	28		29									
Negeri Sembilan	1.0	4	1260	17	20	1	1	1	1	0	780	621	645	374	389							
		5	4	4	4	1		3		3	3		3									
Kedah & Perlis	2.1	5	2093	21	33	1	0	1	0	0	2190	1878	890	623	295							
		7	7	5	6	1		3		7	9		5									
Terengganu	1.0	1	821	4	2	0	0	0	0	0	362	276	265	181	173							
		1	3	1	0	0		0		1	1		1									
Pahang	1.5	4	1317	18	28	4	3	1	1	0	841	753	518	526	362							
		5	4	4	5	2		3		3	4		4									
Kelantan	1.5	3	1778	17	36	6	4	2	1	1	2380	1956	1278	1438	939							
		4	6	4	6	4		6		8	10		11									
Sarawak	2.4	5	1785	15	22	13	6	3	1	1	690	585	248	314	133							
		7	6	4	4	8		8		2	3		3									
Sabah	3.0	4	1206	34	28	3	1	1	0	1	1166	766	255	496	166							
		5	4	8	5	2		3		4	4		4									

*PMP = Per Million Population
Note: Percentage is to the nearest decimal point

Table 1.2 Utilization of Acute Coronary/Cardiac Services in Malaysia 2006

	Pop in 000,000	Current bed	Bed occup. rate	Use for all acute	Use for ACS	Use for non cardiac	ACS denied	ACS denied % of all ACS	Required bed strengths (RBS)		RBS if non cardiac excluded	
									No	(% shortfall)	No	(% shortfall)
Malaysia	26.6	414	89	65	40	35	6703	53	515	80	375	110
Sector												
Private		214	83	51	27	49	1635	48	238	90	162	132
MOH		169	93	77	49	23	4722	62	242	70	190	89
IJN		12	83	36	26	64	177	53	13	92	7	171
University		19	104	76	58	24	168	14	21	90	16	119
States												
P.Pinang	1.5	54	148	42	21	58	917	80	71	76	28	193
Melaka	0.7	31	59	55	33	45	460	62	35	89	28	111
Johor	3.2	30	136	75	48	25	700	56	40	75	29	103
Perak	2.3	25	145	74	58	26	1116	63	61	41	41	61
Selangor & Kuala Lumpur	6.4	148	75	65	43	35	1463	40	164	90	128	116
Negeri Sembilan	1.0	17	47	69	42	31	472	126	21	81	18	94
Kedah & Perlis	2.1	21	93	83	28	17	182	29	22	95	18	117
Terengganu	1.0	4	59	66	43	34	96	53	5	86	4	100
Pahang	1.5	18	41	78	54	22	187	36	19	95	18	100
Kelantan	1.5	17	114	72	53	28	723	50	24	71	17	100
Sarawak	2.4	15	24	76	41	24	46	15	15	100	14	107
Sabah	3.0	34	64	57	37	43	340	69	38	89	32	106

Note: Percentage is to the nearest decimal point.

Table 1.3 Cardiac Care provided for ACS in Malaysia 2006

	Population no in 000,000	ACS admits		PMP*		Trombolytic		PMP*		Angiogram		PMP*		PCI		PMP*		CABG		PMP*	
		No	%	No	%	No	%	No	%	No	%	No	%	No	%	No	%	No	%	No	%
Malaysia	26.64	12534	100	471	38	179	33	4156	33	156	20	2552	96	858	7	32					
Sector																					
Private		3398	27					2168	64			1478		534	16						
MOH		7580	60					1403	19			801		264	3						
IJN		335	3					257	77			134		39	12						
University		1221	10					328	27			139		21	2						
States																					
Pulau Pinang	1.49	1140	9	764	33	250	47	534	47	358	32	367	246	119	10	80					
Melaka	0.73	745	6	1027	58	594	35	261	35	360	16	122	168	59	8	81					
Johor	3.17	1243	10	392	27	106	36	450	36	142	25	307	97	89	7	28					
Perak	2.28	1772	14	776	28	218	52	924	52	405	26	457	200	141	8	62					
Selangor & Kuala Lumpur	6.43	3683	29	573	40	228	34	1238	34	193	25	908	141	277	8	43					
Negeri Sembilan	0.96	374	3	389	53	208	18	66	18	69	11	40	42	14	4	15					
Kedah & Perlis	2.11	623	5	295	55	163	21	128	21	61	13	78	37	26	4	12					
Terengganu	1.04	181	1	173	50	48	18	32	18	31	11	19	18	6	3	6					
Pahang	1.45	526	4	362	59	215	18	93	18	64	10	55	38	19	4	13					
Kelantan	1.53	1438	11	939	36	336	14	197	14	129	8	119	78	44	3	29					
Sarawak	2.36	314	3	133	45	60	23	71	23	30	9	29	12	11	3	5					
Sabah	3.00	496	4	166	21	34	33	162	33	54	10	51	17	53	11	18					

*PMP = Per Million Population
Note: Percentage is to the nearest decimal point.

Figure 1.1 Relationship between availability of cardiologist and provision of emergency Coronary angiogram and Percutaneous Coronary Intervention (PCI) for patients admitted with ACS in 2006

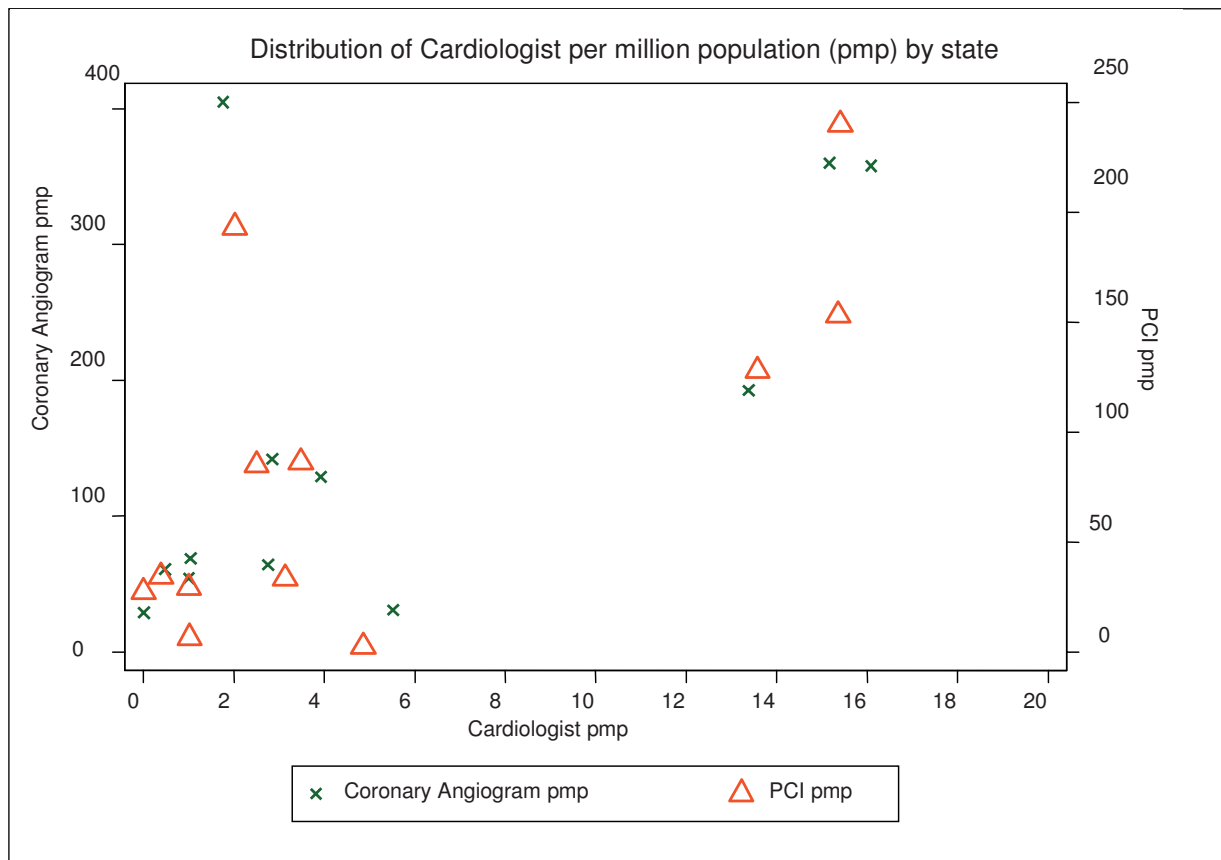


Figure 1.2 Relationship between availability of Cath Lab and provision of emergency Coronary angiogram and Percutaneous Coronary Intervention (PCI) for patients admitted with ACS in 2006

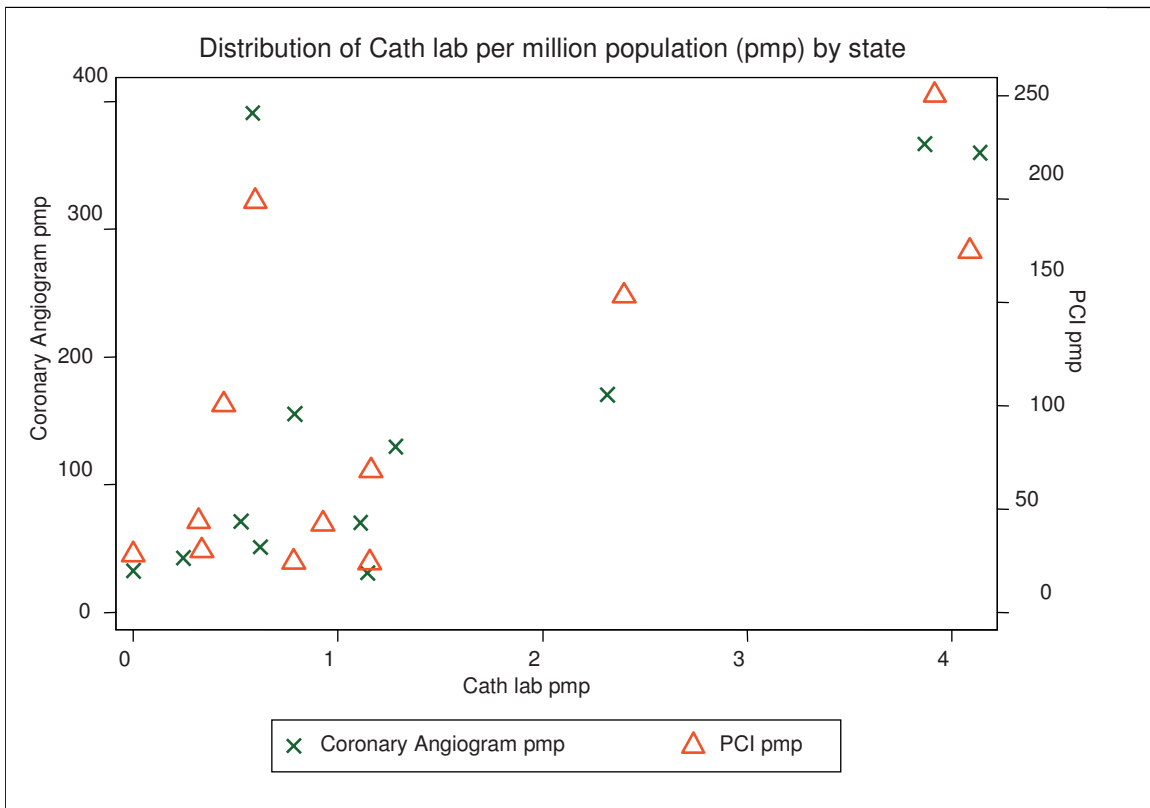
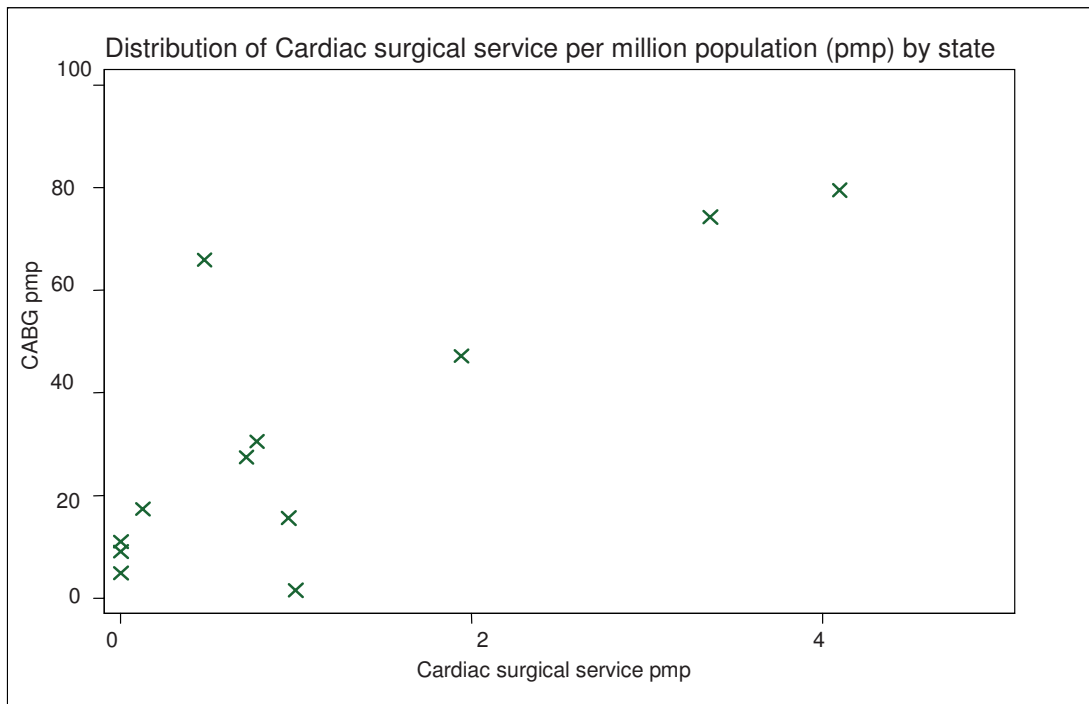


Figure 1.3 Relationship between availability of cardiac surgical services and provision of emergency CABG for patients admitted with ACS in 2006



CHAPTER 2

PATIENT CHARACTERISTICS

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Introduction

In 2006, a total of 3422 patients had baseline characteristics recorded in the Acute Coronary Syndrome section of the National Cardiovascular Database (ACS; NCVD). These were divided into patient demographics, significant past medical history and anthropometric measurements (Table 2.1)

Demographics

Of the ethnic distribution, 49% of patients were Malay, 23% Chinese, 23% Indian, and approximately 4% representing other indigenous groups as well as non-Malaysian nationals. The ethnic groups were subdivided into 12 categories to include the most prevalent groups: namely Malay, Chinese, Indian, Orang Asli, Kadazan, Melanau, Murut, Bajau, Bidayuh, Iban, Malaysians of other ethnicities and non-Malaysian nationals. The mean age of the patients was 59 years (range 21-100); 23% of the patients were below 50 years; 31% of patients were aged between 50 and 60 years old; 26% were aged between 60 and 70 years old and the remaining 21% aged 70 years and older. In term of gender, 75% of the patients were male.

Significant past medical history

Smoking history was subdivided to patients having never smoked, 'former smokers' and current smokers. Our findings revealed 40% of patients had never smoked prior to admission, 24% were former smokers and 33% were current smokers. A significant number of patients had a family history of premature cardiovascular disease. Of the 61% of this data field completed, it was noted that 19.7% of patients did have a 'positive' family history. In a recent Public Health Survey, dyslipidaemia, hypertension and diabetes were noted to be prevalent amongst Malaysian adults³. Not surprisingly, of the 59% of the dyslipidaemia data field completed, 55.9% of patients had a diagnosis of the condition prior to presentation with ACS. Of the 84% of the hypertension data field completed, 72.6% of patients had a diagnosis of the condition prior to presentation of ACS. Of the 80% of the diabetes data field completed, 55.0% of patients had a diagnosis of the condition prior to admission.

Having a prior history of myocardial infarction increases the risk of subsequent ACS compared to those who had not. Of the 70% of this data field completed, 22.9% of patients had a prior history of a myocardial infarction prior to the index admission with ACS. Similarly, a history of documented coronary artery disease could increase this risk. Of the 66% of this data field completed, 22.7% of the patients had a positive finding of documented coronary artery disease prior to the index admission with ACS.

In terms of symptoms of angina, of the 74% of the data field completed for chronic angina prior to admission, 20.2% of patients were found to have this condition. Of the 79% of the data field

completed for new onset angina prior to admission, 57.0% of patients had this condition prior to the index admission for ACS. Heart failure, particularly of ischaemic origin, is associated with poorer long term clinical outcomes. Of the 75% of this data field completed, 10.7% of patients had a prior history of heart failure prior to admission.

Other non-cardiac co-morbid conditions were also investigated. The result shows that 4% of patients had a history of chronic lung disease prior to admission, with 75% of this data field completed. 7% of patients had renal disease prior to admission, with 75% of this data field completed. 4% of patients had a prior history of cerebrovascular disease, with 75% of this data field completed. 1% of patients had a prior history of peripheral vascular disease, with 74% of this data field completed. Combining all the variables above, 91% of the data fields were completed; 97.8% of patients had at least one of the above-mentioned cardiovascular risk factors at the index admission with ACS.

Analysis of patients with coronary artery disease, aggregating subjects with a prior history of myocardial infarction, with angiographically-proven coronary stenosis of greater than 50%, with chronic angina and new onset angina, 80% of the data fields were completed; 80.0% of patients had symptoms or established documented evidence of coronary artery disease prior to the index admission with ACS.

Anthropometrics

Patient anthropometric data was subdivided into Body Mass Index (BMI), Waist-Hip Ratio (WHR) and waist circumference.

The mean BMI was 25.8 ± 4.51 ; the median BMI was 25.2 (13.2-62.4). 75% of subjects had a BMI>23. 40.7% of patients had a calculated WHR obtained. The mean WHR was 0.97 ± 0.08 ; the median WHR was 0.96 (0.54-1.85). 28% of male subjects had a WHR>1.0; 88% of female subjects had a WHR>0.85. 34.0% had a waist circumference measurement performed. The mean value was 89.8 ± 14.6 cm; the median value was 90cm (36-160); 50% of male subjects had a value over 90cm; 80% of female patients had a value over 80cm.

Patient characteristics and different types of ACS presentations (Table 2.6)

Subdividing ACS presentations to ST-elevation myocardial infarction (STEMI; n=1445), non-STEMI (n=1132) and unstable angina (UA; n=845), we found that 42% of patients were admitted with STEMI, 33% with non-STEMI, and 25% unstable angina.

Mean ages for patients presenting with STEMI, NSTEMI and UA were 56, 62 and 60 years respectively; the patient group aged between 50 and 60 years old accounting for 32%, 29% and 31%

of each type of ACS presentation respectively. Comparing gender of ACS presentation in STEMI, NSTEMI and UA group males comprised 85%, 69% and 66% in each group, respectively. On ethnicity, Malays accounted for 54% of patients admitted with STEMI, 45% for NSTEMI and 46% with UA.

Fifty percent of patients admitted with STEMI were current smokers, compared to 23% in the NSTEMI group, and 18% in the UA group. "Never been smokers" accounted for 29% of the STEMI group, 49% for the NSTEMI group and 48% of the UA group. Twelve percent of patients admitted with STEMI had a family history of premature cardiovascular disease, compared with 11% in the NSTEMI group and 13% in the UA group. Nineteen percent in the STEMI group, 41% in the NSTEMI group and 46% in the UA group recorded history of dyslipidaemia. Forty-seven percent in the STEMI group, 70% in the NSTEMI group and 73% in the UA group had a history of hypertension. Thirty-six percent in the STEMI group, 51% in the NSTEMI group and 47% in the UA group had a history of diabetes.

Ten percent in the STEMI group, 19% in the NSTEMI group and 24% in the UA group had a prior history of myocardial infarction. Five percent of patients in the STEMI group, 20% of patients in the NSTEMI group and 24% of patients in the UA group had a previously documented significant coronary artery disease. Accordingly, 7% in the STEMI group, 17% in the NSTEMI group and 25% in the UA group had a prior history of chronic stable angina; however, 43% in the STEMI group, 48% in the NSTEMI group and 43% in the UA group had new onset angina. Three percent in the STEMI group, 14% in the NSTEMI group and 10% in the UA group had a recorded history of heart failure prior to the index admission with ACS.

Two percent in the STEMI group, 5% in the NSTEMI group and 5% in the UA group had a prior history of chronic lung disease. Four percent in the STEMI group, 13% in the NSTEMI group and 6% in the UA group had a prior history of renal disease. Three percent in the STEMI group, 6% in the NSTEMI group and 4% in the UA group had a prior history of cerebrovascular disease; <1% in the STEMI group; 2% in the NSTEMI group and 1% in the UA group had a prior history of peripheral vascular disease.

Ninety-one percent of patients in the STEMI group had at least one of the above mentioned cardiovascular risk factors at the index admission for ACS, compared to 97% in the NSTEMI group and 98% in the UA group.

Mean BMI for patients admitted with STEMI, NSTEMI and UA were 26, 25, and 26 respectively; patients with a BMI>23 accounted for 76%, 72% and 79% of the respective groups. The mean waist-hip ratio (WHR) of patients admitted with STEMI, NSTEMI and UA were 0.97, 0.97 and 0.96 respectively. Whilst the measurements in the male patients were similar, in women, 10% of patients in

the STEMI group, 14% in the NSTEMI group and 9% in the UA group had a WHR of ≤ 0.85 . Mean waist circumference (WC) for patients admitted with STEMI was 89cm; in NSTEMI 90cm and UA 92cm. Forty-eight percent of male patients admitted with STEMI had a WC ≥ 90 cm; compared to 47% in the NSTEMI group and 57% in the UA group. 80% of female patients admitted with STEMI had a WC ≥ 80 , compared to 79% in the NSTEMI group, and 82% in the UA group.

Commentary

Demographics

In 2006, Malays made up an estimated 50.4% of the total population of 26.64 million, Chinese 23.7%, Indian 7.1%, and Non-Malay Bumiputera 11%³. The distribution of Malay and Chinese patients admitted with ACS recorded in this registry for the same year was similar with the proportion of ethnic distribution in the country. While, there were disproportionately more Indian patients and disproportionately less non-Malay Bumiputera patients.

With the country's gender distribution of nearly 1:1, it was surprising to note that 75% of the subjects were male³; in comparison to the 66% of male found in the Global Registry of Acute Coronary Events (GRACE)². We used the GRACE Registry as the comparative in this Patient Characteristics as it is the largest ongoing, multicentre Registry, for ACS worldwide.

National statistics reported the life expectancy at birth in Malaysia in 2006 to be 74.1 years, with males living to 71.8 years and females 76.3 years³. The mean age for subjects in our registry was relatively young at 59 years. Eighty percent of subjects were aged less than 70 years, and significantly, 23% were aged less than 50 years. The median age of our subjects was 59, which was significantly younger than the 66 years of those found in the GRACE Registry.

Significant past medical history

In term of smoking habits amongst subjects in the Registry, 33% were current smokers, compared to 56.7% from the GRACE Registry. The National prevalence of current smokers in adults aged 25-64 years old in 2005/2006 was 25.5%¹. Despite the comparatively smaller proportion of current smokers in our Registry as compared with GRACE, our subjects present at a younger age. It is possible that our patients are more susceptible to chemicals in cigarette smoke. This is compounded by the observation that nearly a fifth of our subjects have a documented family history of cardiovascular disease.

Hypercholesterolaemia, hypertension and diabetes are prevalent in our country, with reported figures of 53.5%, 25.7% and 5.0% amongst adults aged 25-64 years in 2005/2006¹. Our findings

demonstrated that for subjects enrolled into our registry in 2006, 55.9% had dyslipidaemia, 72.6% had hypertension, and 55.0% had diabetes. GRACE Registry figures are 43.6%, 57.8% and 23.3%. Our observations suggest that hypertension and diabetes confers a disproportionately higher risk for developing ACS, when compared to our National population as a whole, and compared to the subjects recruited from the GRACE Registry.

For those with symptoms and known significant coronary artery disease, 80% of our subjects have at least one of the following: a history of angina prior to the index admission with ACS, known angiographically proven coronary artery disease with at least one vessel over 50% stenosis, or a previous documented myocardial infarction. In fact, only 20.2% of patients had chronic stable angina over two weeks prior to admission in contrast with 68.1% demonstrated by the GRACE Registry.

Other cardiovascular risk factors, with the exception of renal dysfunction (9.3% versus 7.2%) featured less commonly in our subjects compared to those in the GRACE registry: history of heart failure (10.7% versus 11.0%), cerebrovascular disease (5.3% versus 8.3%) and peripheral vascular disease (1.4% versus 10.3%).

In terms of patient characteristics, improved completion of data fields over the coming years may yet shed more light into these patterns.

Anthropometrics

Anthropometric findings suggest that the majority of our subjects were overweight and had an abnormally elevated abdominal circumference.

Patient characteristics and different types of ACS presentations

Our findings reveal that the majority of ACS admissions to our hospitals were STEMI. Furthermore, there are early indications that a large majority of them were male and of Malay ethnicity. The proportion of current smokers was higher in the STEMI group, when compared to NSTEMI and UA. This could indicate that smoking plays a larger role in massive plaque rupture, a hallmark of STEMI, amongst patients admitted with ACS in Malaysia. However, other established cardiovascular risk factors appeared more prevalent in patients presenting with NSTEMI and UA when compared to STEMI. Anthropological measurements did not seem to account for significant differences among the patients presenting with the different ACS presentations except for WHR in the female gender, where a lower ratio seemed to confer a larger protective effect in NSTEMI compared to STEMI and UA.

Summary Points:

- Of the 3,422 patients admitted with ACS to the 11 participating sites in 2006, 49% were Malay, 23% Chinese, 23% Indian and about 4% were others.
- 75% of the subjects were male and the female patients may be underrepresented. The mean age for subjects in our registry was also relatively young.
- Subdividing ACS presentations revealed that 42% had STEMI, 33% NSTEMI and 25% UA.
- Patients with STEMI had a younger mean age and comprised more males, Malays and active smokers compared with NSTEMI and UA groups.
- In this registry there was higher prevalence of established cardiovascular risk factors. Upon admission with ACS, majority of them has either history of MI or are known to have significant CAD.

References:

1. "Data and Statistics"; Non-Communicable Diseases Surveillance, Malaysia. www.dph.gov.my
2. Granger CB, Goldberg RJ, Dabbous O, et al. Predictors of Hospital Mortality in the Global Registry of Acute Coronary Events. *Arch Intern Med.* 2003;163:2345-2353.
3. "Key Statistics" and "Key Data"; Department of Statistics, Malaysia. www.statistics.gov.my

Table 2.1 Summary of patients characteristics for patient with ACS, Malaysia 2006

	Total=3422
1. DEMOGRAPHICS	
1.1 Age, years	
• Mean, SD	59 (12)
• Median (min, max)	59 (21,100)
1.2 Age group, no. %	
• 20 - <30	23 (1)
• 30 - <40	143 (4)
• 40 - <50	621 (18)
• 50 - <60	1054 (31)
• 60 - <70	881 (26)
• 70 - <80	571 (17)
• ≥80	129 (4)
1.3 Gender, no. %	
• Male	2569 (75)
• Female	853 (25)
1.4 Ethnic group, no. %	
• Malay	1684 (49)
• Chinese	786 (23)
• Indian	799 (23)
• Orang Asli	0 (0)
• Kadazan	2 (0)
• Melanau	0 (0)
• Murut	0 (0)
• Bajau	1 (0)
• Bidayuh	28 (1)
• Iban	48 (1)
• Other Malaysian	37 (1)
• Foreigner	37 (1)
2. OTHER CORONARY RISK FACTORS	
2.1 Smoking, no. %	
• Never	1370 (40)
• Former (quit >30 days)	805 (24)
• Current (any tobacco use within last 30 days)	1138 (33)
• Unknown	109 (3)
2.2 Family history of premature cardiovascular disease, no. %	
• Yes	404 (12)
• No	1684 (49)
• Not known	1334 (39)

	Total=3422
2.3 Antropometric	
BMI	
• N	1926
• Mean, SD	25.8 (4.4)
• Median, (min, max)	25.2 (13.2,60.4)
BMI, kg/m², no. %	
• <18.5	58 (3)
• 18.5-23	426 (22)
• > 23	1442 (75)
WHR	
• N	1394
• Mean, SD	0.97 (0.09)
• Median, (min, max)	0.96 (0.46,1.85)
WHR, no. %	
• Men	1091
• ≤ 1.0	786 (72)
• >1.0	305 (28)
• Women	303
• ≤ 0.85	35 (12)
• >0.85	268 (88)
Waist circumference, cm	
• N	1502
• Mean, SD	89.7 (14.4)
• Median, (min, max)	90 (36,160)
Waist circumference, cm, no. %	
• Men	1162
• ≤ 90	586 (50)
• > 90	576 (50)
• Women	340
• ≤ 80	68 (20)
• > 80	272 (80)
2.4 Co-morbidity	
Dyslipidaemia, no. %	
• Yes	1131 (33)
• No	902 (26)
• Not known	1389 (41)
Hypertension, no. %	
• Yes	2084 (61)
• No	786 (23)
• Not known	552 (16)

	Total=3422
Diabetes, no. %	
• Yes	1497 (44)
• No	1226 (36)
• Not known	699 (20)
Fasting blood glucose, mmol/L	
• N	2561
• Mean (SD)	8.2 (4)
• Median (min, max)	6.8 (3,29.9)
Myocardial infarction history, no. %	
• Yes	562 (16)
• No	1847 (54)
• Not known	1013 (30)
Documented CAD > 50% stenosis, no. %	
• Yes	508 (15)
• No	1734 (51)
• Not known	1180 (34)
Chronic angina (onset more than 2 weeks ago), no. %	
• Yes	502 (15)
• No	2012 (59)
• Not known	908 (27)
New onset angina (less than 2 weeks), no. %	
• Yes	1532 (45)
• No	1160 (34)
• Not known	730 (21)
Heart failure, no. %	
• Yes	284 (8)
• No	2289 (67)
• Not known	849 (25)
Chronic lung disease, no. %	
• Yes	130 (4)
• No	2431 (71)
• Not known	861 (25)
Renal disease, no. %	
• Yes	253 (7)
• No	2305 (68)
• Not known	864 (25)
Cerebrovascular disease, no. %	
• Yes	149 (4)
• No	2420 (71)
• Not known	853 (25)

	Total=3422
Peripheral vascular disease, no. %	
• Yes	37 (1)
• No	2492 (73)
• Not known	893 (26)
None of the above, no. %	
• Yes	67 (2)
• No	3050 (89)
• Not known	305 (9)
Coronary artery disease**, no. %	
• Yes	2199 (64)
• No	532 (16)
• Not known	691 (20)

* Not known includes patients who do not know their co-morbidities and missing data

**Coronary artery disease is defined as "Yes" on any of the following co-morbidities: 1) History of myocardial infarction, 2) Documented CAD >50% stenosis, 3) Chronic angina (onset more than 2 weeks ago), 4) New onset angina (less than 2 weeks).

Note: Percentage is to the nearest decimal point.

Figure 2.1.1 Age group (years) distribution for patients with ACS, Malaysia 2006

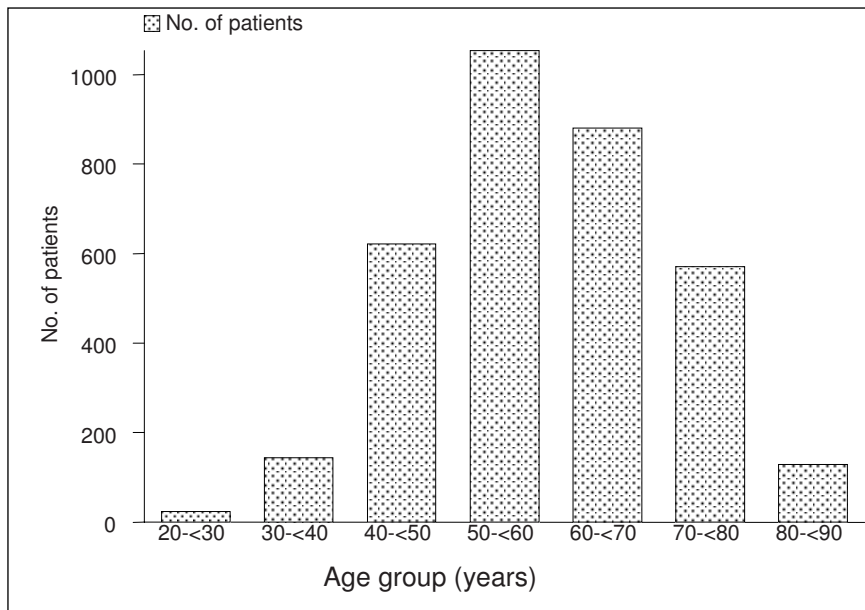


Figure 2.1.2 Gender distribution for patients with ACS, Malaysia 2006

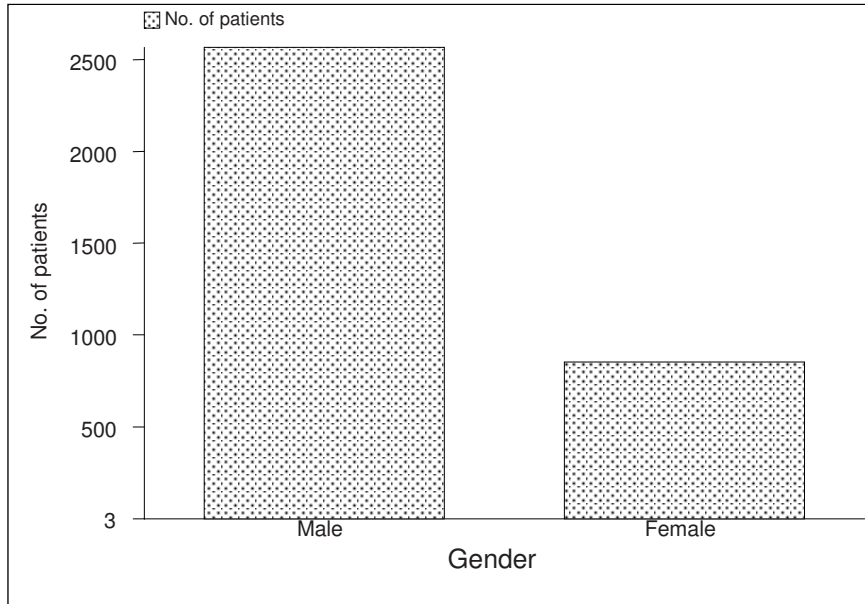


Figure 2.1.3 Ethnic group distribution for patients with ACS, Malaysia 2006

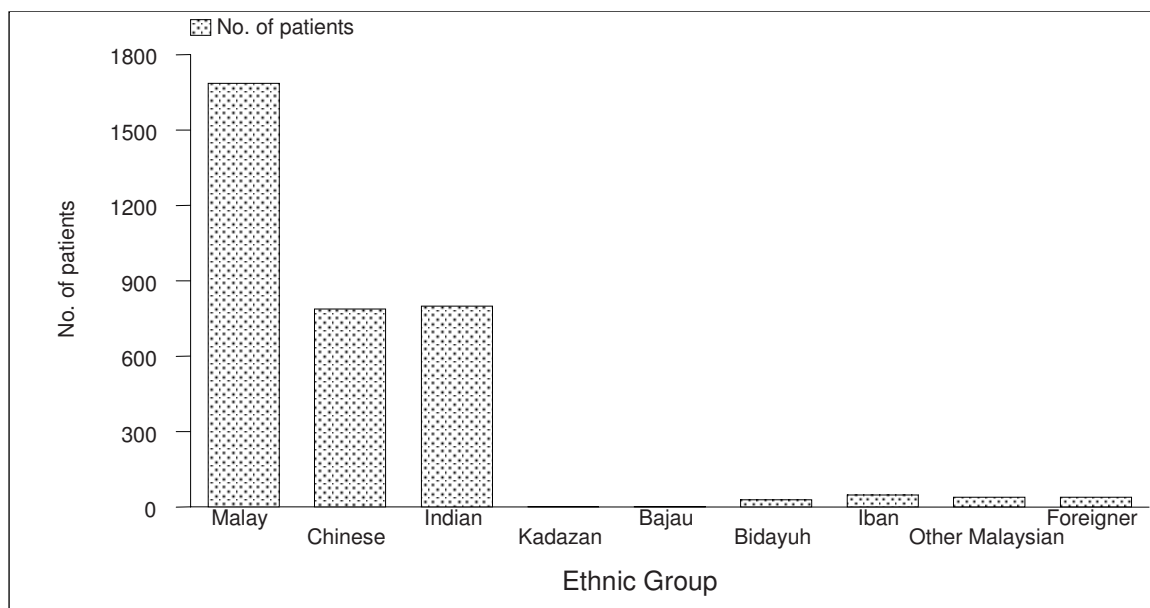
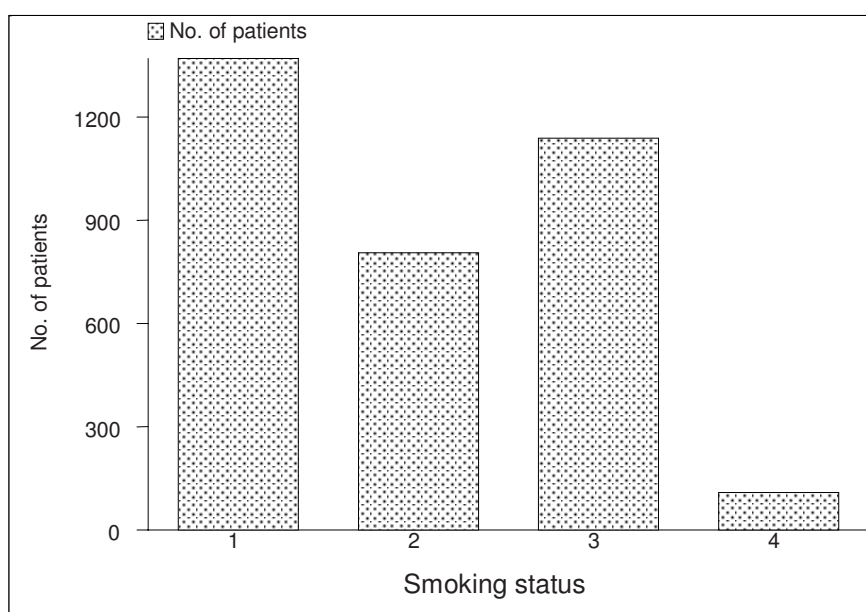


Figure 2.1.4 Smoking status for patients with ACS, Malaysia 2006



1. Never, 2. Former (quit >30 days), 3. Current (any tobacco use within last 30 days), 4. Unknown

Figure 2.1.5 Family history of premature cardiovascular disease for patients with ACS, Malaysia 2006

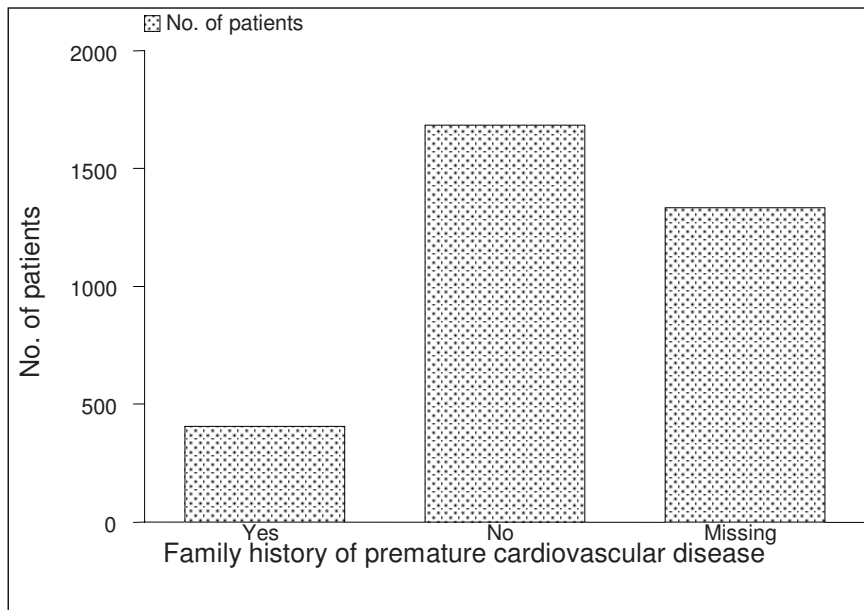


Figure 2.1.6 BMI for patients with ACS, Malaysia 2006

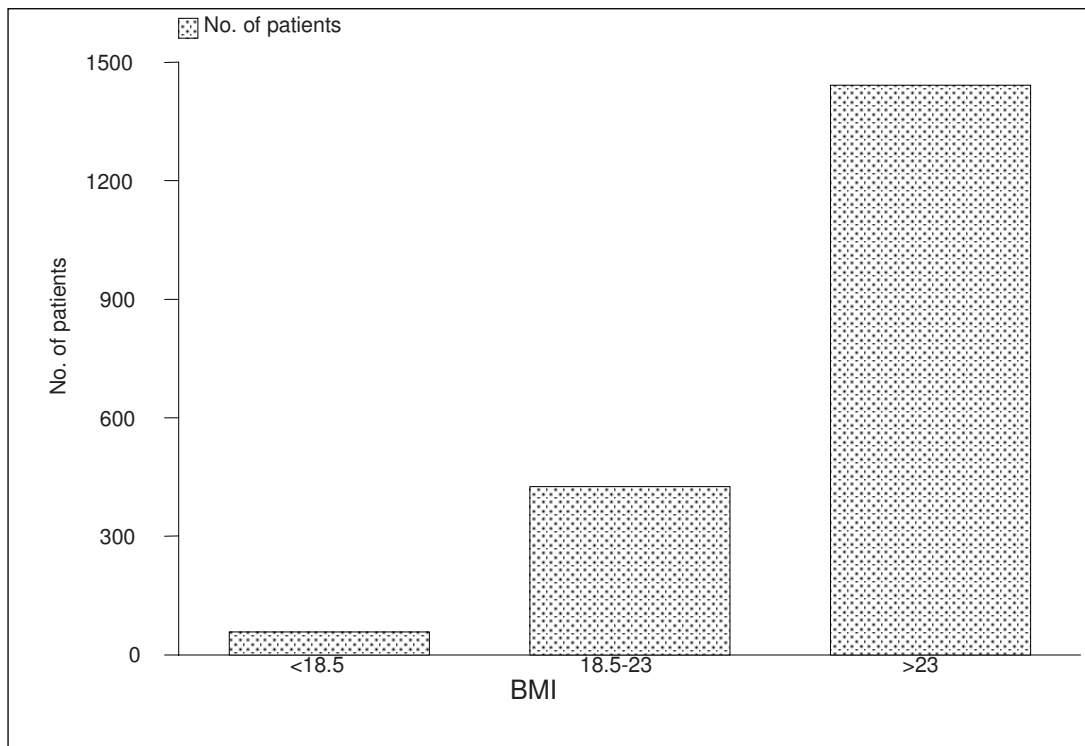


Figure 2.1.7 WHR for patients with ACS, Malaysia 2006

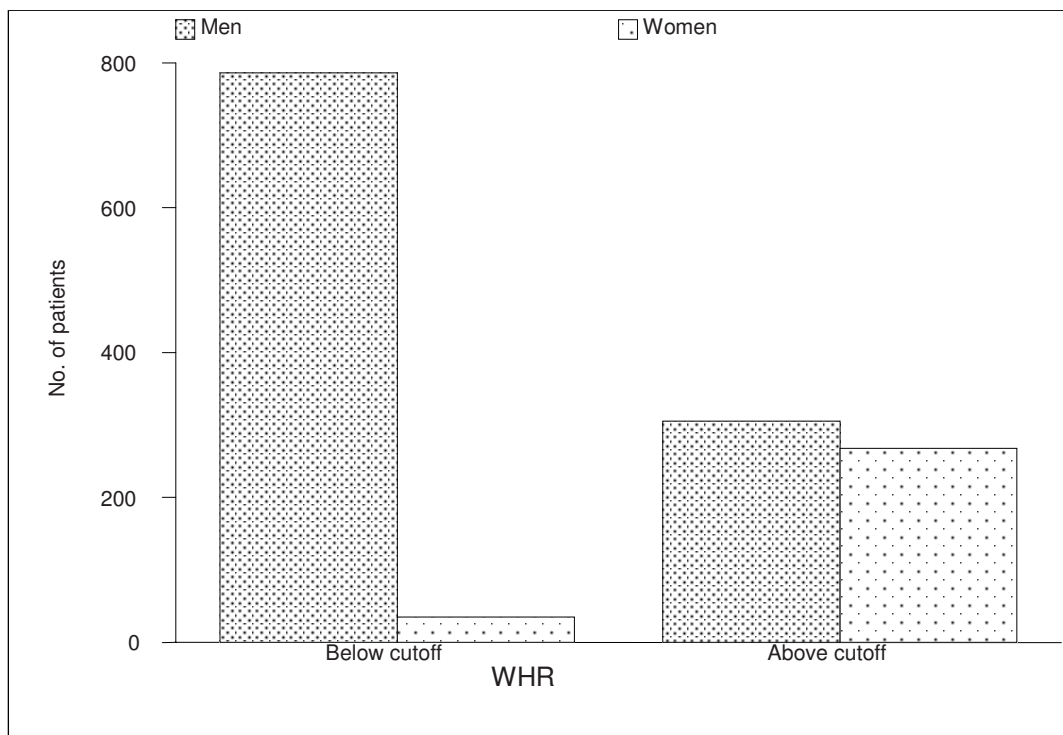


Figure 2.1.8 Waist circumference (cm) for patients with ACS, Malaysia 2006

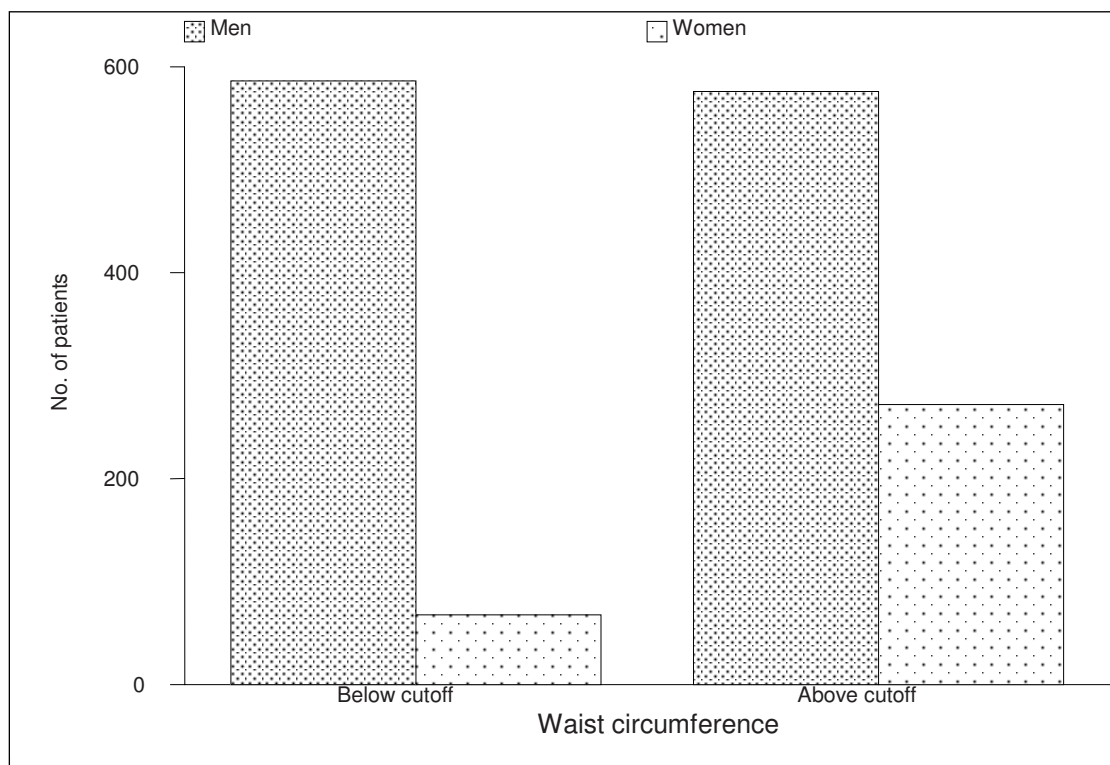
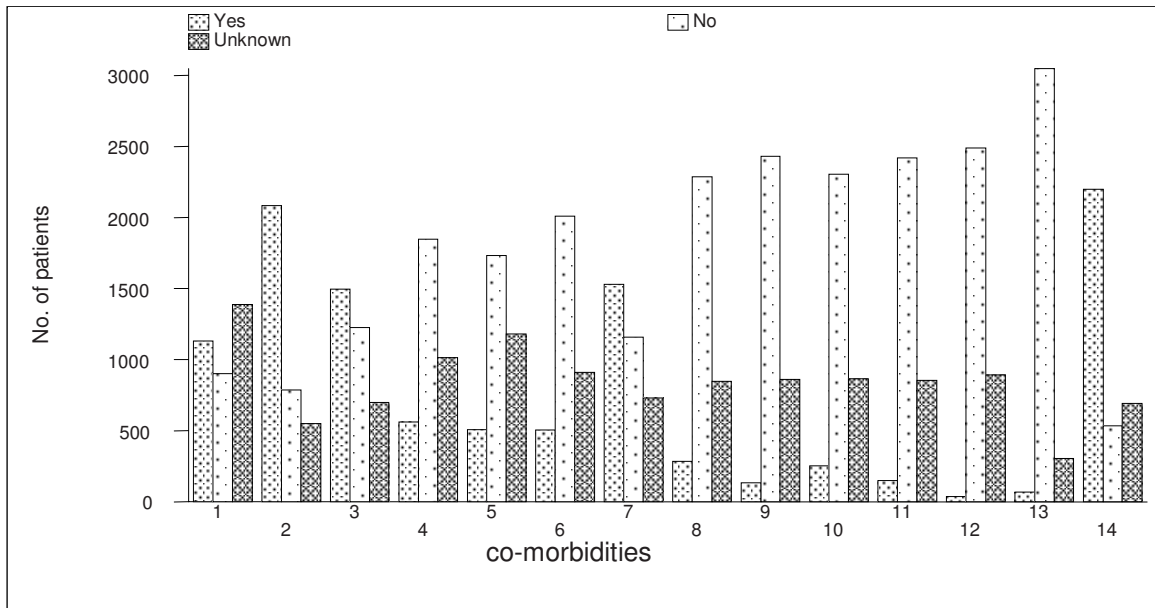


Figure 2.1.9 Co-morbidities for patients with ACS, Malaysia 2006



1. Dyslipidaemia, 2. Hypertension, 3. Diabetes, 4. History of myocardial infarction, 5. Documented CAD > 50% stenosis, 6. Chronic angina (onset more than 2 weeks ago), 7. New onset angina (less than 2 weeks), 8. Heart failure, 9. Chronic lung disease, 10. Renal disease, 11. Cerebrovascular disease, 12. Peripheral vascular disease, 13. None of the above, 14. Coronary artery disease*

Table 2.2.1 Distribution of patients with ACS by SDP, Malaysia 2006

	SDP	No.	%
1	University Malaya Medical Centre, Kuala Lumpur	802	23
2	National Heart Institute, Kuala Lumpur	456	13
3	Kuala Lumpur Hospital, Kuala Lumpur	413	12
4	Penang Hospital, Penang	482	14
5	Sarawak General Hospital, Sarawak	375	11
6	Sultanah Aminah Hospital, Johor	242	7
7	Sultanah Bahiyah Hospital, Kedah	160	5
8	Tuanku Ja'afar Hospital, Negeri Sembilan	146	4
9	Tuanku Fauziah Hospital, Perlis	53	2
10	Raja Perempuan Zainab II Hospital, Kelantan	141	4
11	Tengku Ampuan Afzan Hospital, Pahang	152	4
	Total	3422	100

* Each SDP started to contribute data at different time

Note: Percentage is to the nearest decimal point.

Figure 2.2.1 Distribution of patients with ACS by SDP, Malaysia 2006

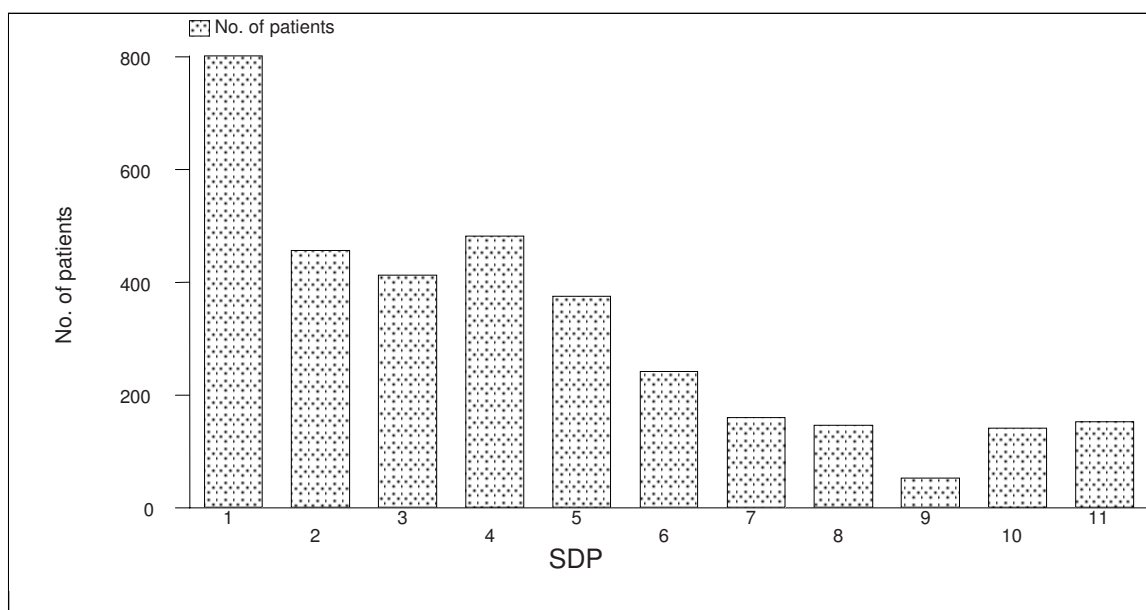


Table 2.2.2 SDP-ethnicity distribution of patients with ACS, Malaysia 2006 (row percent)

	SDP	Ethnic group									
		Malay		Chinese		Indian		Others*		Total	
		No.	%	No.	%	No.	%	No.	%	No.	%
1	University Malaya Medical Centre	264	33	180	22	335	42	23	3	802	100
2	National Heart Institute	216	47	85	19	141	31	14	3	456	100
3	Kuala Lumpur Hospital	223	54	74	18	91	22	25	6	413	100
4	Penang Hospital	190	39	172	36	113	23	7	1	482	100
5	Sarawak General Hospital	141	38	151	40	4	1	79	21	375	100
6	Sultanah Aminah Hospital	128	53	66	27	46	19	2	1	242	100
7	Sultanah Bahiyah Hospital	145	91	7	4	7	4	1	1	160	100
8	Tuanku Ja'afar Hospital	71	49	23	16	52	36	0	0	146	100
9	Tuanku Fauziah Hospital	49	92	1	2	2	4	1	2	53	100
10	Raja Perempuan Zainab II Hospital	134	95	7	5	0	0	0	0	141	100
11	Tengku Ampuan Afzan Hospital	123	81	20	13	8	5	1	1	152	100

*Others includes Orang asli, Kadazan, Melanau, Murut, Bajau, Bidayuh, Iban, other Malaysian and foreigner

Note: Percentage is to the nearest decimal point.

Figure 2.2.2 SDP-ethnicity distribution of patients with ACS, Malaysia 2006

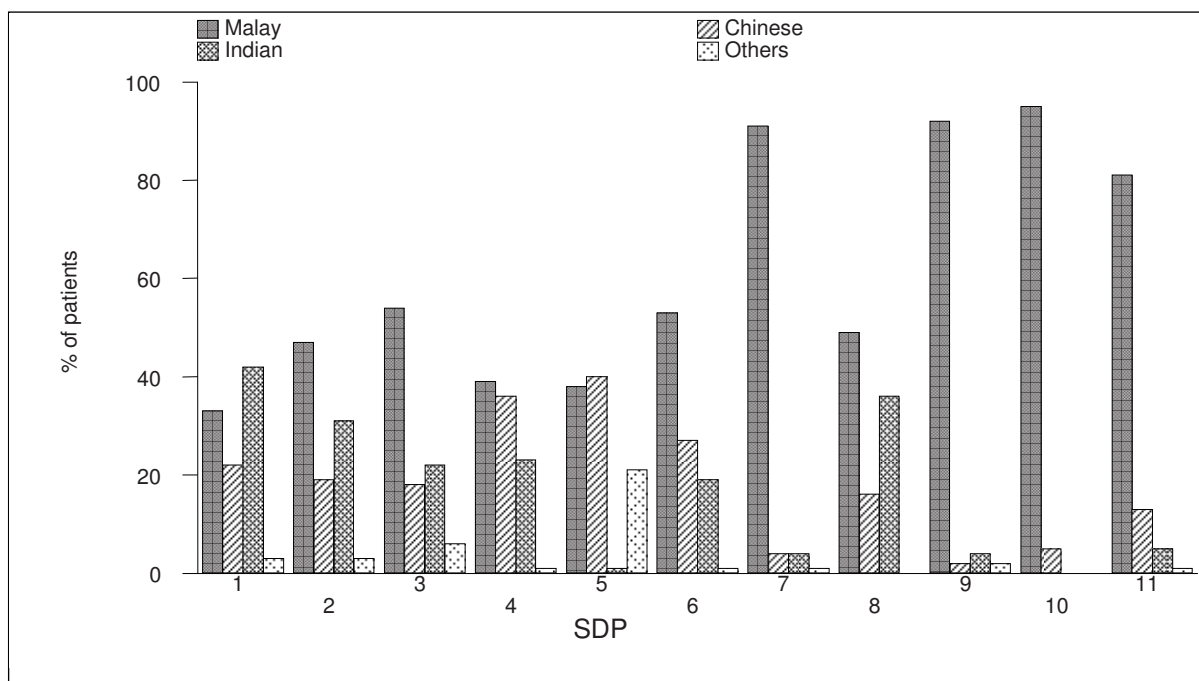


Table 2.2.3: SDP-ethnicity distribution of patients' admitted to participating sites, Malaysia 2006 (row percent)

	SDP	Ethnic group									
		Malay		Chinese		Indian		Others*		Total	
		No.	%	No.	%	No.	%	No.	%	No.	%
1	University Malaya Medical Centre	16657	41	11576	29	9826	24	2336	6	40395	100
2	National Heart Institute	5357	51	2096	20	2447	23	534	5	10434	100
3	Kuala Lumpur Hospital	7739	67	691	6	1565	13	1605	14	11600	100
4	Penang Hospital	18162	42	16615	39	6218	14	1916	4	42911	100
5	Sarawak General Hospital	12218	37	8156	25	92	0	12641	38	33109	100
6	Sultanah Aminah Hospital	37248	58	14125	22	7735	12	5485	8	64593	100
7	Sultanah Bahiyah Hospital	14381	82	1857	11	628	4	626	4	17492	100
8	Tuanku Ja'afar Hospital	23083	53	7571	17	10690	24	2306	5	43650	100
9	Tuanku Fauziah Hospital	23221	89	1505	6	353	1	919	4	25998	100
10	Raja Perempuan Zainab II Hospital	37716	94	1220	3	91	0	1095	3	40122	100
11	Tengku Ampuan Afzan Hospital	25112	78	3986	12	1363	4	1680	5	32141	100

* Patients age > 20 years old only

*Others includes Orang asli, Kadazan, Melanau, Murut, Bajau, Bidayuh, Iban, other Malaysian and foreigner

Note: Percentage is to the nearest decimal point.

Figure 2.2.3: SDP-ethnicity distribution of patients' admitted to participating sites, Malaysia 2006

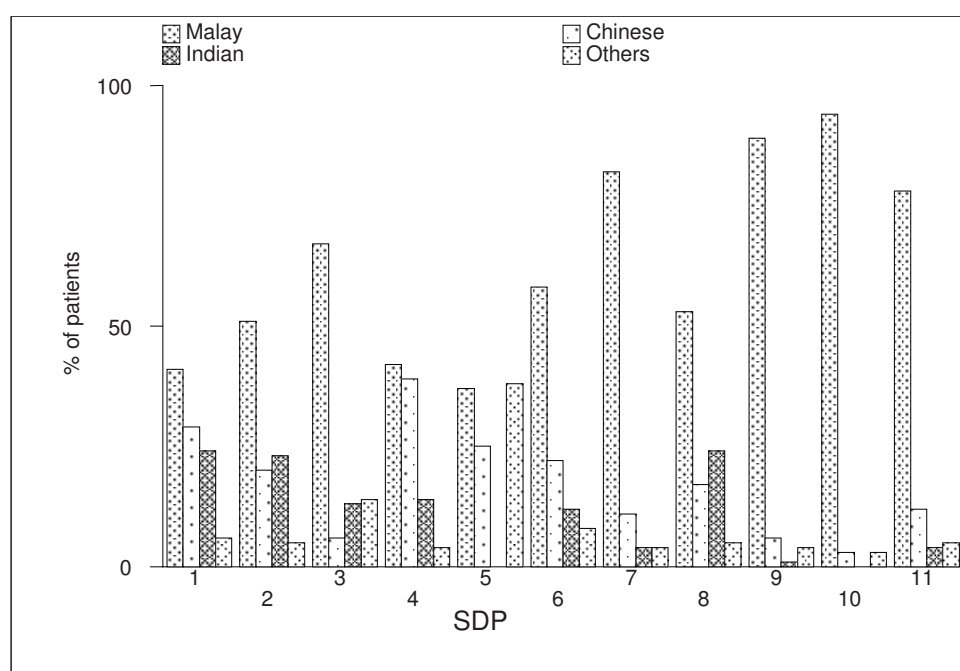


Table 2.2.4 SDP-gender distribution of patients with ACS, Malaysia 2006 (row percent)

	SDP	Gender					
		Male		Female		Total	
		No.	%	No.	%	No.	%
1	University Malaya Medical Centre	588	73	214	27	802	100
2	National Heart Institute	357	78	99	22	456	100
3	Kuala Lumpur Hospital	329	80	84	20	413	100
4	Penang Hospital	366	76	116	24	482	100
5	Sarawak General Hospital	263	70	112	30	375	100
6	Sultanah Aminah Hospital	212	88	30	12	242	100
7	Sultanah Bahiyah Hospital	103	64	57	36	160	100
8	Tuanku Ja'afar Hospital	77	53	69	47	146	100
9	Tuanku Fauziah Hospital	45	85	8	15	53	100
10	Raja Perempuan Zainab II Hospital	109	77	32	23	141	100
11	Tengku Ampuan Afzan Hospital	120	79	32	21	152	100

Note: Percentage is to the nearest decimal point.

Figure 2.2.4 SDP-gender distribution of patients with ACS, Malaysia 2006

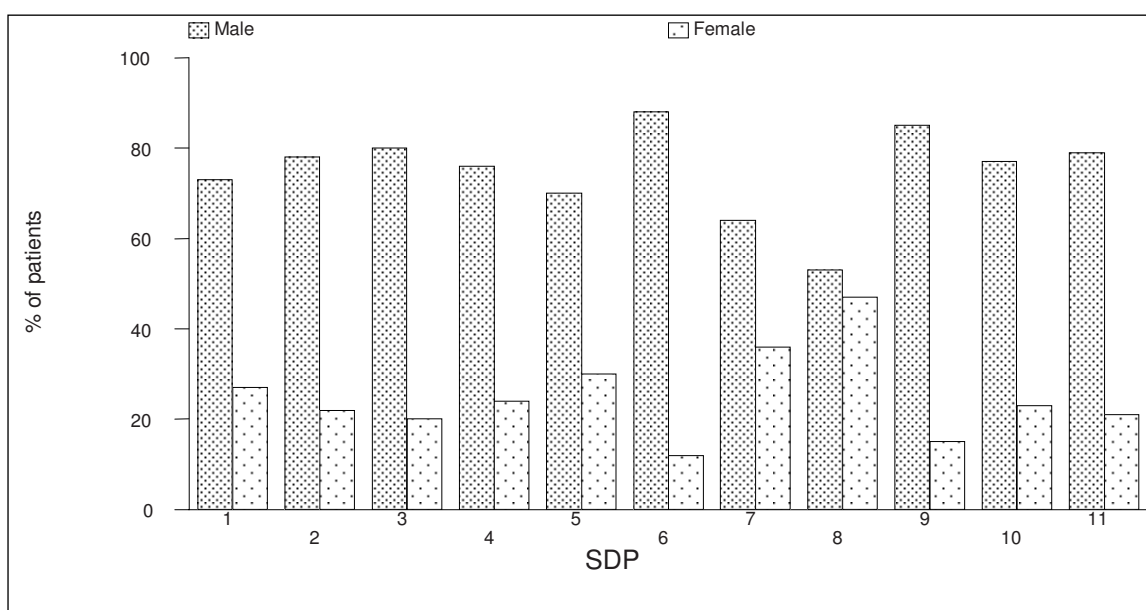


Table 2.2.5: SDP-gender distribution of patients admitted to participating sites, Malaysia 2006 (row percent)

	SDP	Gender					
		Male		Female		Total	
		No.	%	No.	%	No.	%
1	University Malaya Medical Centre	16233	40	24350	60	40395	100
2	National Heart Institute	3134	30	7300	70	10434	100
3	Kuala Lumpur Hospital	3970	34	7630	66	11600	100
4	Penang Hospital	19382	45	23529	55	42911	100
5	Sarawak General Hospital	9704	29	23405	71	33109	100
6	Sultanah Aminah Hospital	22690	35	41903	65	64593	100
7	Sultanah Bahiyah Hospital	6368	36	11124	64	17492	100
8	Tuanku Ja'afar Hospital	15202	35	28448	65	43650	100
9	Tuanku Fauziah Hospital	11036	42	14962	58	25998	100
10	Raja Perempuan Zainab II Hospital	14312	36	25810	64	40122	100
11	Tengku Ampuan Afzan Hospital	11532	36	20609	64	32141	100

⁺ Patients age > 20 years old only

Note: Percentage is to the nearest decimal point.

Figure 2.2.5: SDP-gender distribution of patients admitted to participating sites, Malaysia 2006

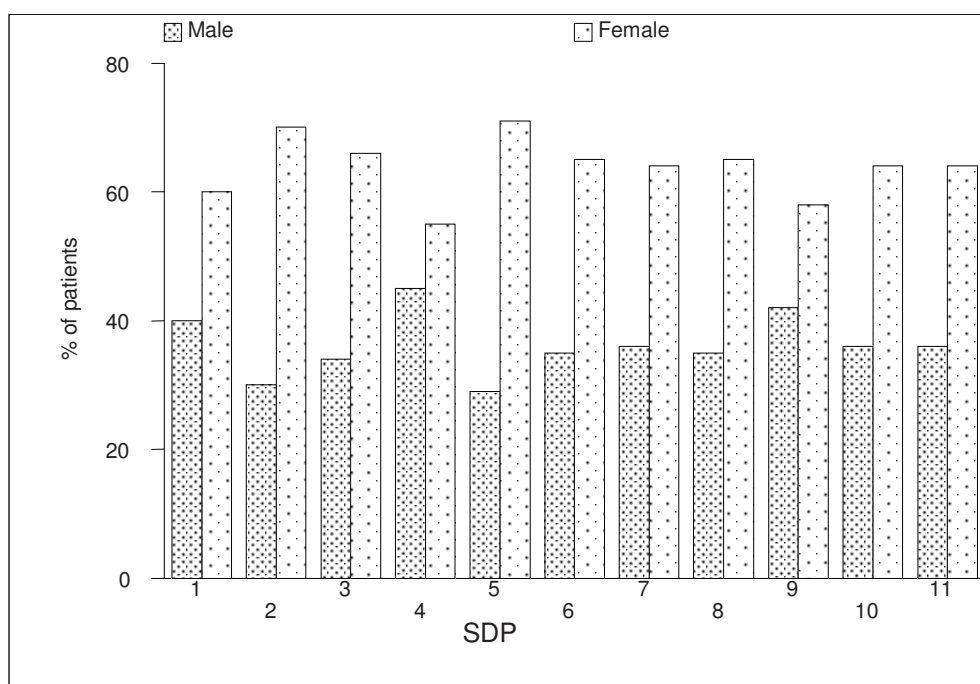


Table 2.3 Age-gender distribution for patients with ACS, Malaysia 2006

Age group	Gender			
	Male		Female	
	No.	%	No.	%
20 - <30	22	1	1	0
30 - <40	131	5	12	1
40 - <50	541	21	80	9
50 - <60	888	35	166	19
60 - <70	616	24	265	31
70 - <80	306	12	265	31
≥80	65	3	64	8
Total	2569	100	853	100

Note: Percentage is to the nearest decimal point.

Figure 2.3 Age-gender distribution for patients with ACS, Malaysia 2006

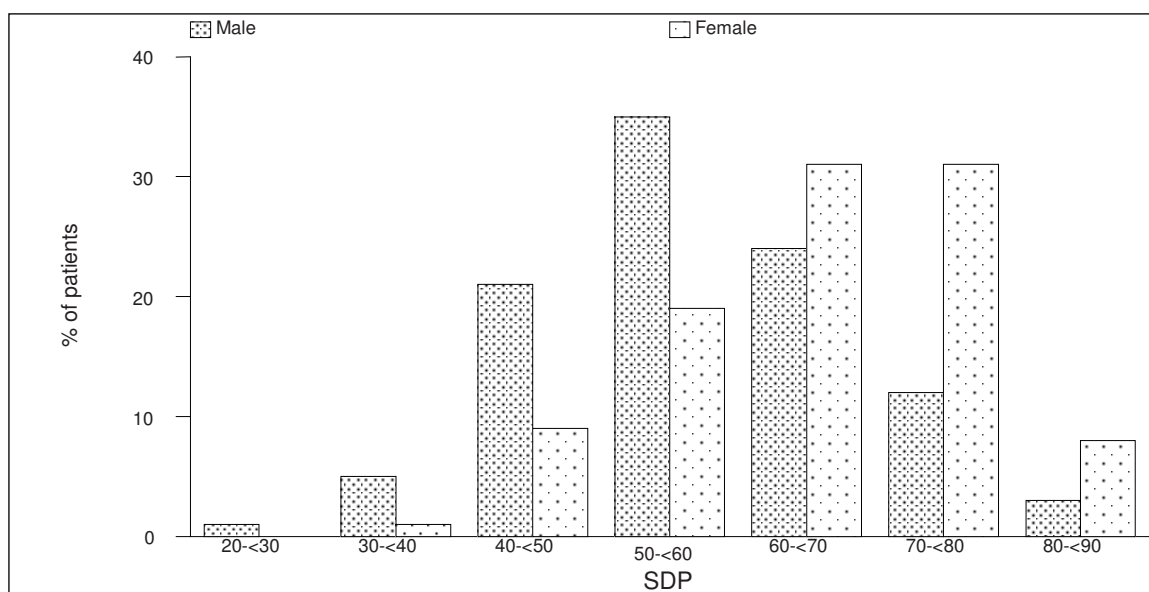


Table 2.3.1 Age-gender distribution for patients with ACS by ethnic group, Malaysia 2006

Gender	Age group	Ethnic group							
		Malay		Chinese		Indian		Others*	
		No.	%	No.	%	No.	%	No.	%
Men	20 - <30	10	1	5	1	5	1	2	2
	30 - <40	67	5	25	4	28	5	11	9
	40 - <50	286	22	91	16	133	23	31	26
	50 - <60	445	34	183	33	227	39	33	28
	60 - <70	324	25	137	24	130	22	25	21
	70 - <80	148	11	95	17	48	8	15	13
	≥80	24	2	25	4	15	3	1	1
	Total	1304	100	561	100	586	100	118	100
Women	20 - <30	1	0	0	0	0	0	0	0
	30 - <40	5	1	3	1	4	2	0	0
	40 - <50	41	11	7	3	32	15	0	0
	50 - <60	88	23	28	12	45	21	5	14
	60 - <70	117	31	68	30	62	29	18	51
	70 - <80	111	29	91	40	53	25	10	29
	≥80	17	4	28	12	17	8	2	6
	Total	380	100	225	100	213	100	35	100

*Others includes Orang asli, Kadazan, Melanau, Murut, Bajau, Bidayuh, Iban, other Malaysian and foreigner

Note: Percentage is to the nearest decimal point.

Figure 2.3.1a Age-gender distribution male patients with ACS by ethnic group, Malaysia 2006

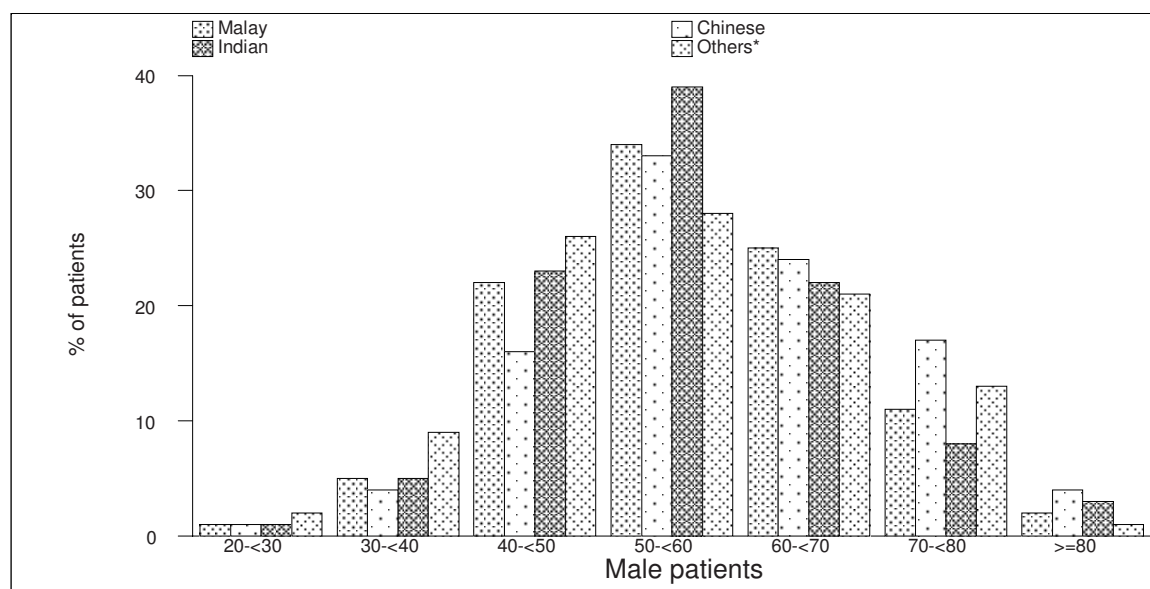


Figure 2.3.1b Age-gender distribution for female patients with ACS by ethnic group, Malaysia 2006

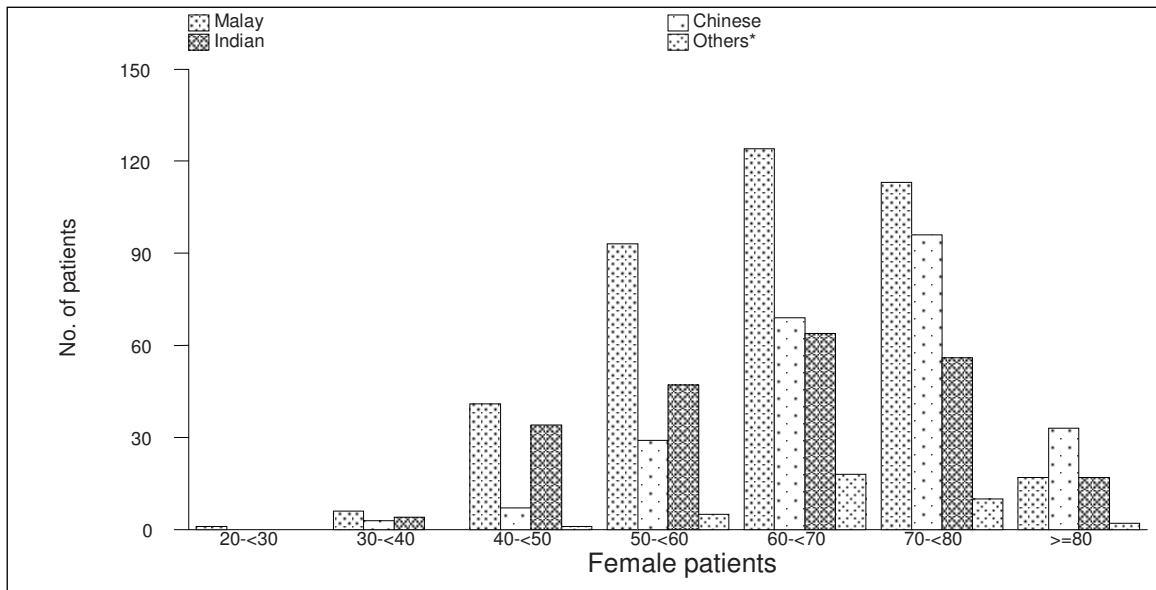


Table 2.3.2 Age-gender distribution for patients with ACS by pre-morbid diabetes, Malaysia 2006

Gender	Age group	Pre-morbid diabetes					
		Diabetic		Non-diabetic		Not known	
		No.	%	No.	%	No.	%
Male	20 - <30	1	0	15	2	6	1
	30 - <40	31	3	58	6	42	7
	40 - <50	194	19	212	22	135	23
	50 - <60	402	39	291	30	195	33
	60 - <70	262	26	224	23	130	22
	70 - <80	114	11	129	13	63	11
	≥80	18	2	32	3	15	3
	Total	1022	100	961	100	586	100
Female	20 - <30	0	0	0	0	1	1
	30 - <40	5	1	3	1	4	4
	40 - <50	44	9	23	9	13	12
	50 - <60	97	20	45	17	24	21
	60 - <70	149	31	83	31	33	29
	70 - <80	154	32	86	32	25	22
	≥80	26	5	25	9	13	12
	Total	475	100	265	100	113	100

Note: Percentage is to the nearest decimal point.

Figure 2.3.2a Age-gender distribution for male patients with ACS by pre-morbid diabetes, Malaysia 2006

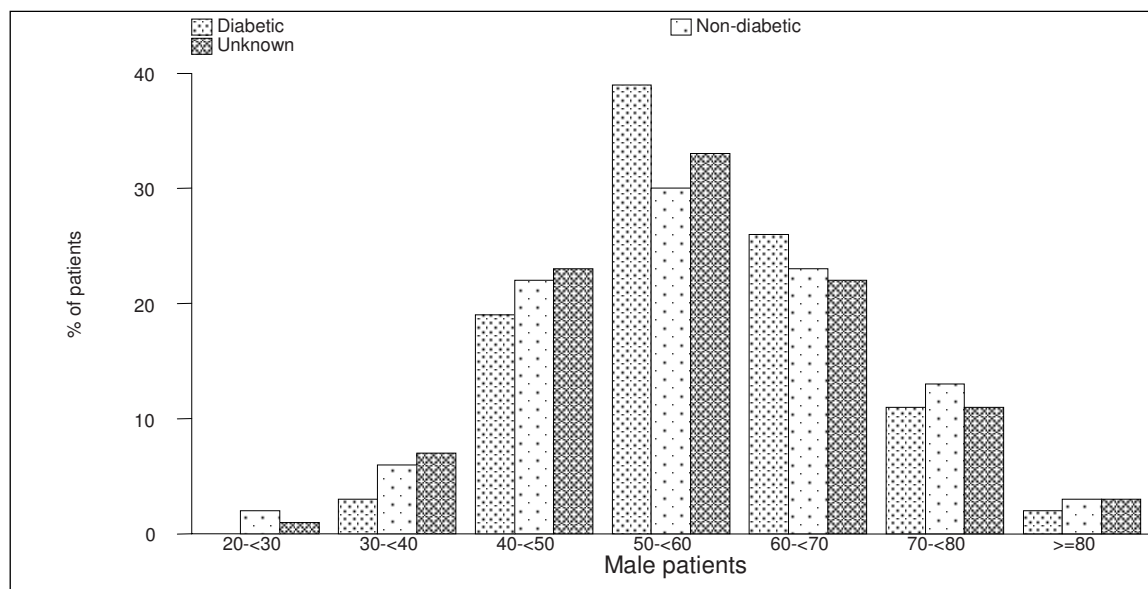


Figure 2.3.2b Age-gender distribution for female patients with ACS by pre-morbid diabetes, Malaysia 2006

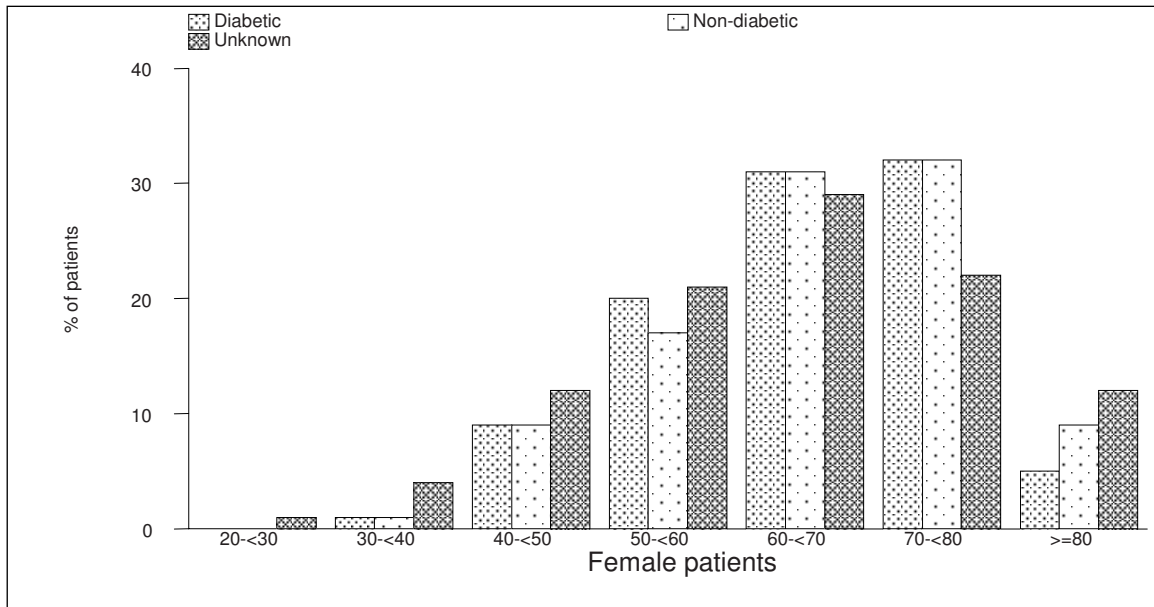


Table 2.3.3 Age-gender distribution for patients with ACS by pre-morbid hypertension, Malaysia 2006

Gender	Age group	Pre-morbid hypertension					
		Hypertensive		Non-hypertensive		Not known	
		No.	%	No.	%	No.	%
Male	20 - <30	2	0	15	2	5	1
	30 - <40	37	3	55	8	39	8
	40 - <50	235	17	178	26	128	27
	50 - <60	514	36	229	34	145	31
	60 - <70	394	28	125	18	97	21
	70 - <80	197	14	63	9	46	10
	≥80	39	3	16	2	10	2
	Total	1418	100	681	100	470	100
Female	20 - <30	0	0	0	0	1	1
	30 - <40	5	1	3	3	4	5
	40 - <50	52	8	14	13	14	17
	50 - <60	129	19	23	22	14	17
	60 - <70	214	32	32	30	19	23
	70 - <80	212	32	27	26	26	32
	≥80	54	8	6	6	4	5
	Total	666	100	105	100	82	100

Note: Percentage is to the nearest decimal point.

Figure 2.3.3a Age-gender distribution for male patients with ACS by pre-morbid hypertension, Malaysia 2006

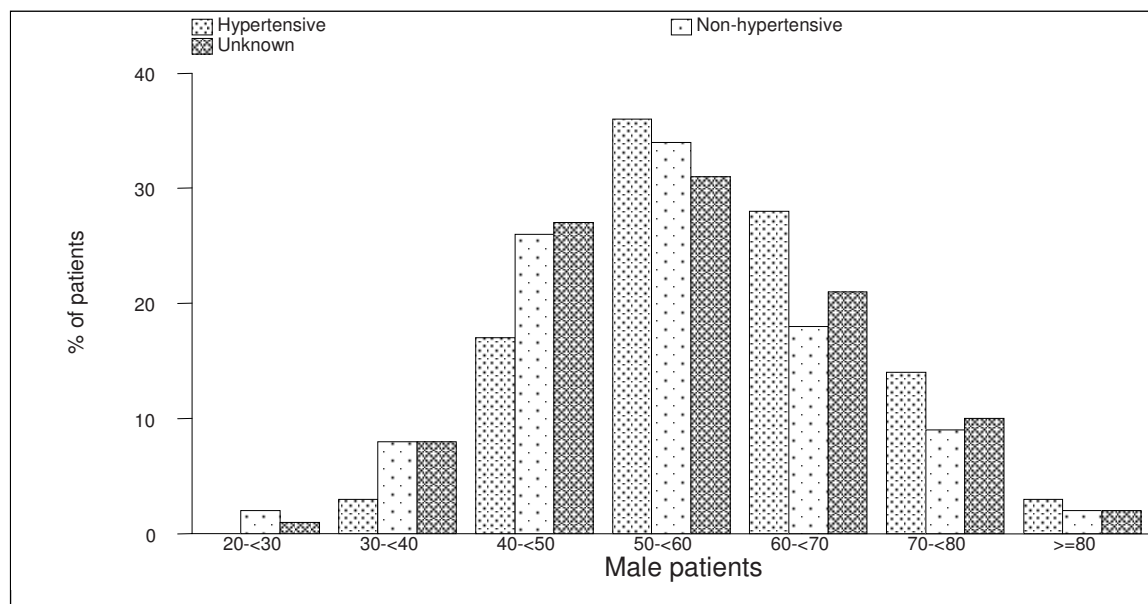


Figure 2.3.3b Age-gender distribution for female patients with ACS by pre-morbid hypertension, Malaysia 2006

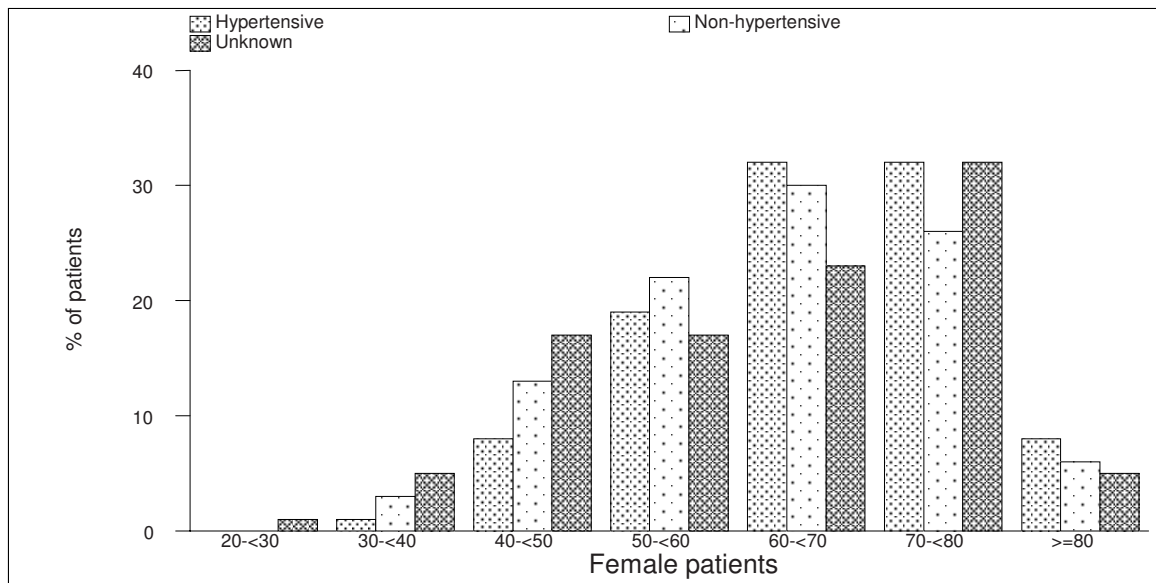


Table 2.3.4 Age-gender distribution for patients with ACS by pre-morbid dyslipidaemia, Malaysia 2006

Gender	Age group	Pre-morbid dyslipidaemia					
		Yes		No		Not known	
		No.	%	No.	%	No.	%
Male	20 - <30	2	0	11	2	9	1
	30 - <40	28	4	44	6	59	5
	40 - <50	145	18	152	22	244	23
	50 - <60	270	34	229	33	389	36
	60 - <70	239	30	140	20	237	22
	70 - <80	98	12	91	13	117	11
	≥80	15	2	22	3	28	3
	Total	797	100	689	100	1083	100
Female	20 - <30	0	0	0	0	1	0
	30 - <40	2	1	1	0	9	3
	40 - <50	27	8	24	11	29	9
	50 - <60	66	20	48	23	52	17
	60 - <70	108	32	64	30	93	30
	70 - <80	116	35	54	25	95	31
	≥80	15	4	22	10	27	9
	Total	334	100	213	100	306	100

Note: Percentage is to the nearest decimal point.

Figure 2.3.4a Age-gender distribution for male patients with ACS by pre-morbid dyslipidaemia, Malaysia 2006

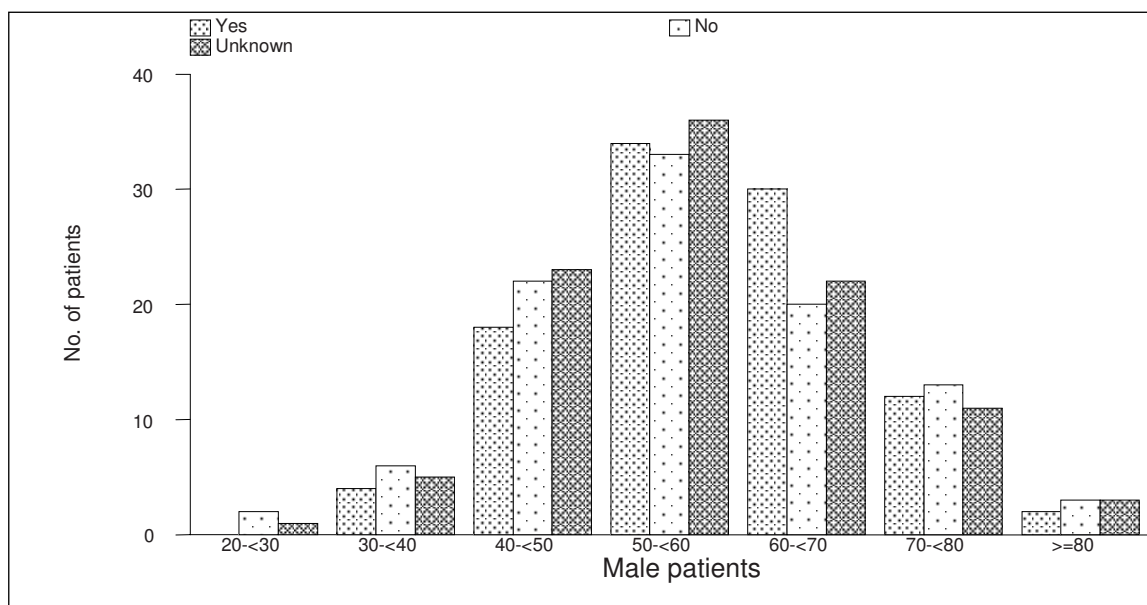


Figure 2.3.4b Age-gender distribution for female patients with ACS by pre-morbid dyslipidaemia, Malaysia 2006

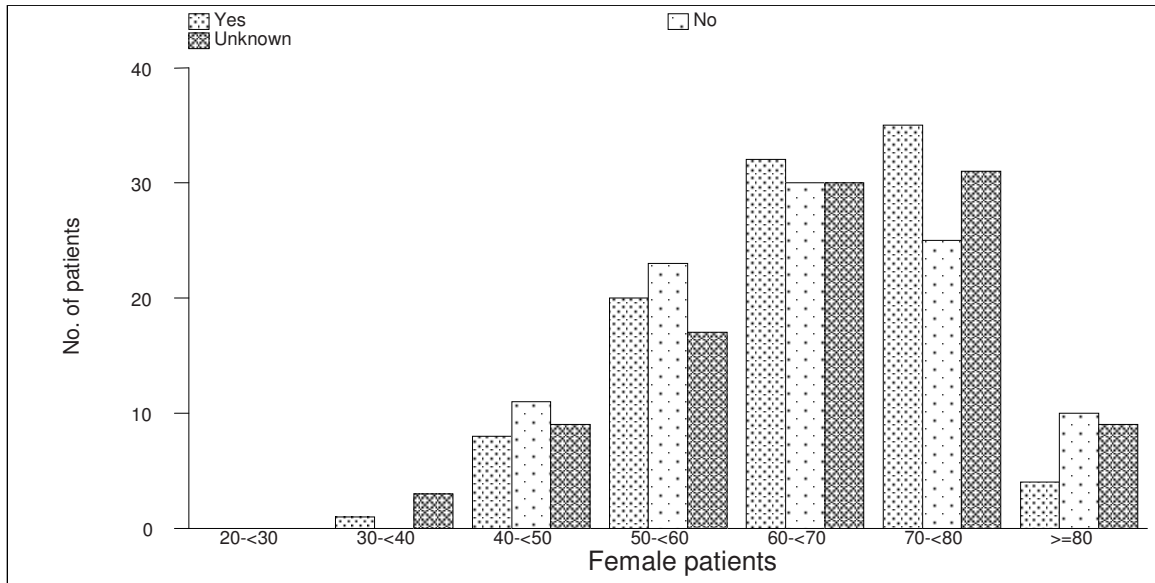


Table 2.3.5 Age-gender distribution for patients with ACS by family history, Malaysia 2006

Gender	Age group	Family history of premature cardiovascular disease					
		Yes		No		Not known	
		No.	%	No.	%	No.	%
Male	20 - <30	3	1	13	1	6	1
	30 - <40	36	11	55	4	40	4
	40 - <50	93	28	250	20	198	20
	50 - <60	130	39	394	32	364	36
	60 - <70	58	17	310	25	248	25
	70 - <80	14	4	176	14	116	12
	≥80	2	1	37	3	26	3
	Total	336	100	1235	100	998	100
Female	20 - <30	0	0	0	0	1	0
	30 - <40	2	3	2	0	8	2
	40 - <50	22	32	33	7	25	7
	50 - <60	14	21	86	19	66	20
	60 - <70	17	25	155	35	93	28
	70 - <80	12	18	138	31	115	34
	≥80	1	1	35	8	28	8
	Total	68	100	449	100	336	100

Note: Percentage is to the nearest decimal point.

Figure 2.3.5a Age-gender distribution for male patients with ACS by family history, Malaysia 2006

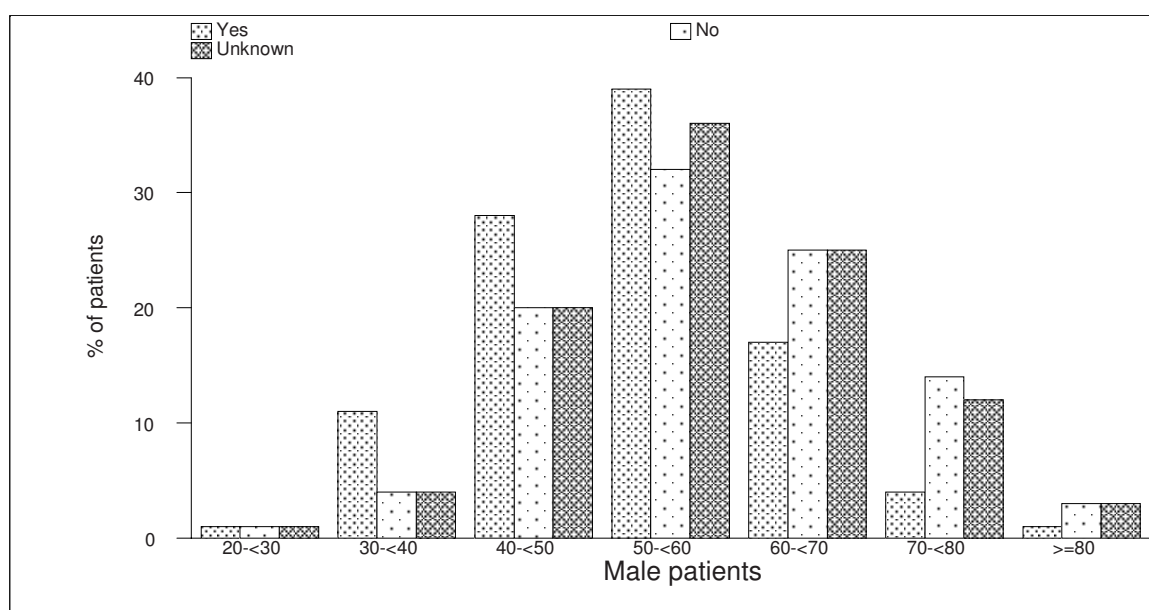


Figure 2.3.5b Age-gender distribution for female patients with ACS by family history, Malaysia 2006

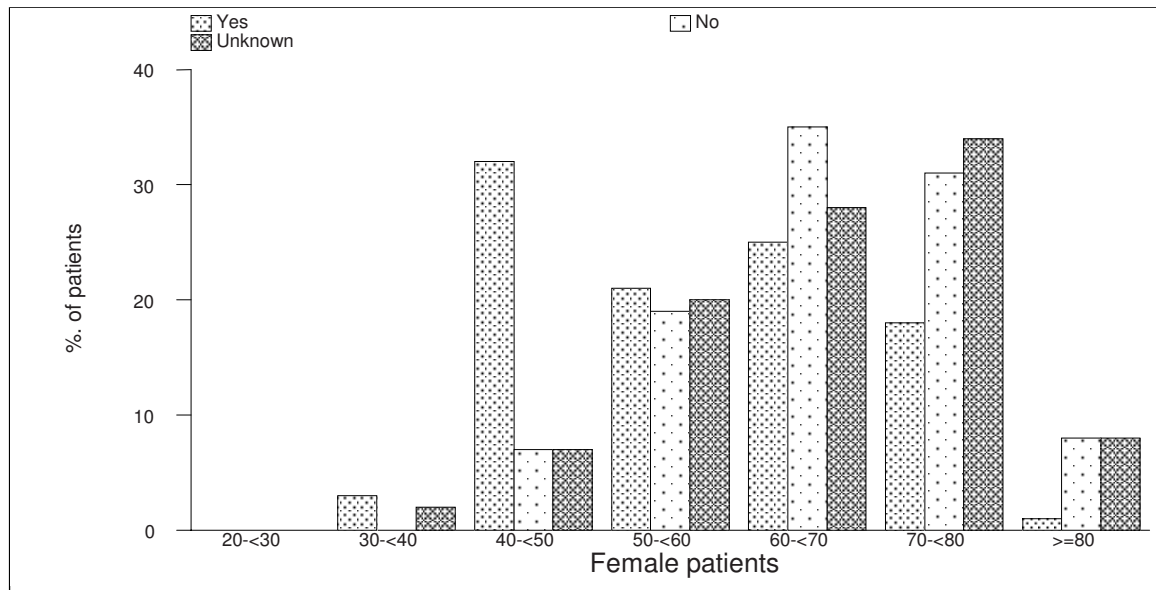


Table 2.3.6 Age-gender distribution for patients with ACS by smoking status, Malaysia 2006

Gender	Age group	Smoking status							
		Never		Former (quit more than 30 days)		Current (any tobacco use within last 30 days)		Unknown	
		No.	%	No.	%	No.	%	No.	%
Male	20 - <30	2	0	1	0	19	2	0	0
	30 - <40	12	2	19	3	97	9	3	3
	40 - <50	112	18	103	14	316	28	10	11
	50 - <60	205	33	249	33	397	36	37	43
	60 - <70	189	31	205	27	198	18	24	28
	70 - <80	82	13	138	18	75	7	11	13
	≥80	16	3	38	5	9	1	2	2
Total	618	100	753	100	1111	100	87	100	
Female	20 - <30	0	0	0	0	1	4	0	0
	30 - <40	12	2	0	0	0	0	0	0
	40 - <50	75	10	2	4	2	7	1	5
	50 - <60	148	20	6	12	5	19	7	32
	60 - <70	234	31	18	35	10	37	3	14
	70 - <80	229	30	17	33	8	30	11	50
	≥80	54	7	9	17	1	4	0	0
Total	752	100	52	100	27	100	22	100	

Note: Percentage is to the nearest decimal point.

Figure 2.3.6a Age-gender distribution for male patients with ACS by smoking status, Malaysia 2006

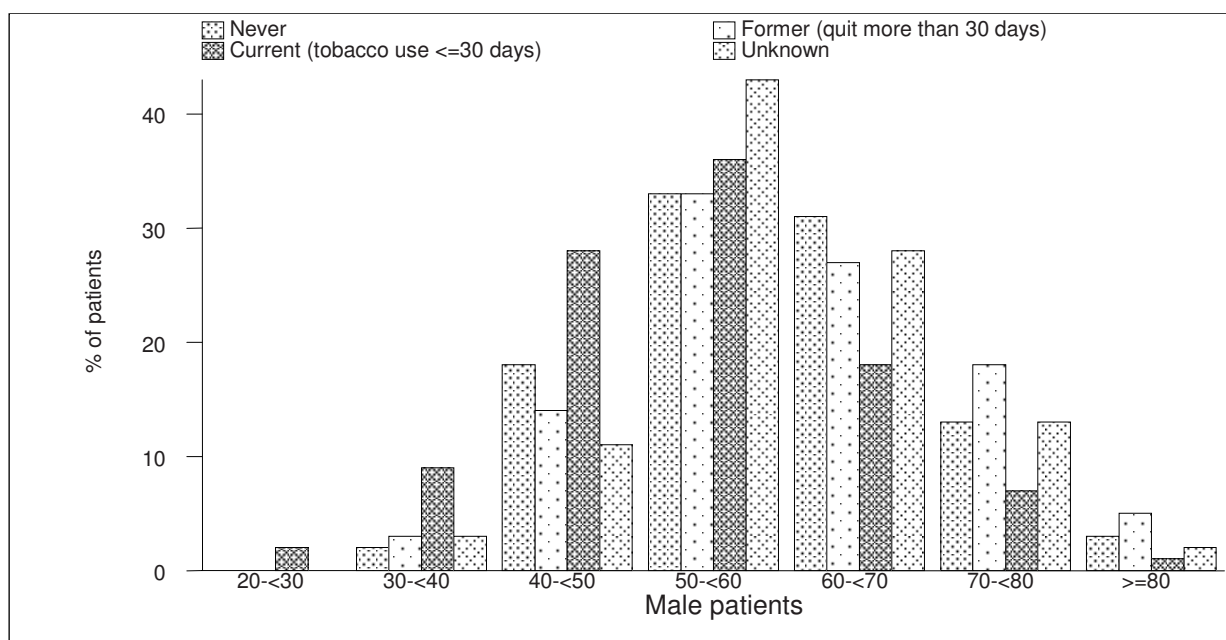


Figure 2.3.6b Age-gender distribution for female patients with ACS by smoking status, Malaysia 2006

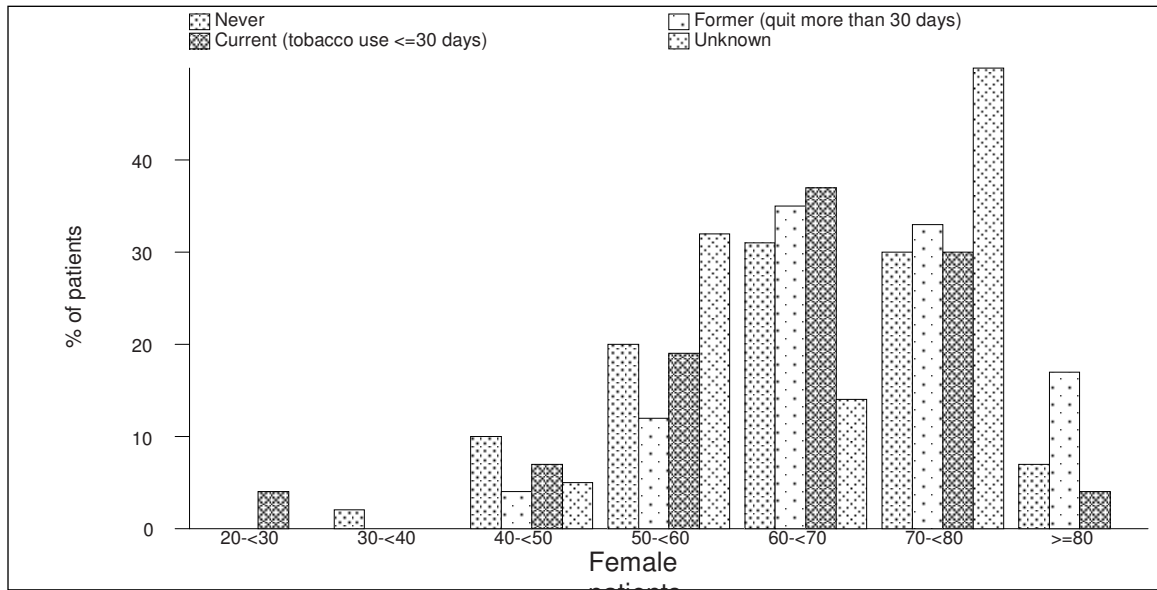


Table 2.4 Pre-morbid distribution for patients with ACS, Malaysia 2006

		Dyslipidaemia																	
		Yes						No						Not known					
		Hypertension						Hypertension						Hypertension					
		Yes		No		Unknown		Yes		No		Unknown		Yes		No		Unknown	
		No.	%	No.	%	No.	%	No.	%	No.	%	No.	%	No.	%	No.	%	No.	%
Diabetes	Yes	540	16	65	2	29	1	231	7	119	3	2	0	373	11	54	2	84	2
	No	280	8	104	3	1	0	241	7	296	9	0	0	163	5	121	4	20	1
	Unknown	73	2	2	0	37	1	8	0	3	0	2	0	175	5	22	1	377	11

** The percentage is based on the grand total (N=3422)

Note: Percentage is to the nearest decimal point.

Figure 2.4a Pre-morbid distribution for diabetic patients with ACS, Malaysia 2006

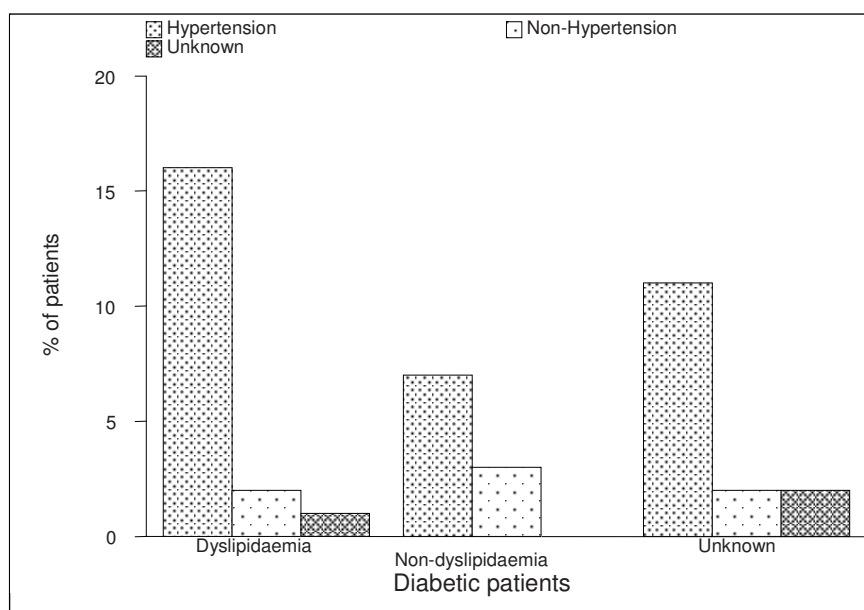


Figure 2.4b Pre-morbid distribution for non-diabetic patients with ACS, Malaysia 2006

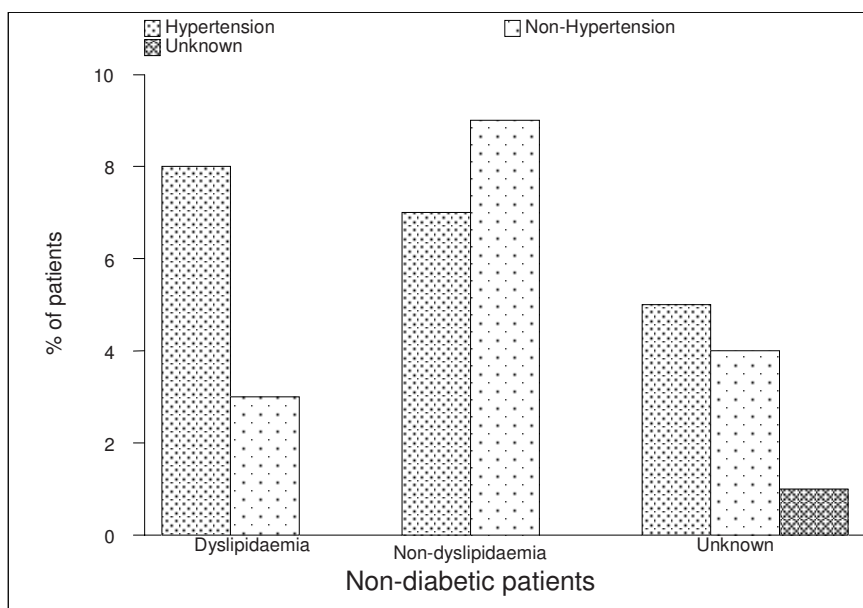


Table 2.5 Presence of cumulative risk factors

Presence of cumulative Risk factors *	Total=3422	
	No.	%
None	143	4
1 risk factor	634	19
2 risk factor	987	29
3 risk factor	938	27
> 3 risk factor	720	21

* Risk factors are defined as presence of dyslipidaemia, hypertension, diabetes, family history of premature cardiovascular disease, smoking, and obesity.

Note: Percentage is to the nearest decimal point.

Figure 2.5 Distribution of presence of cumulative risk factors

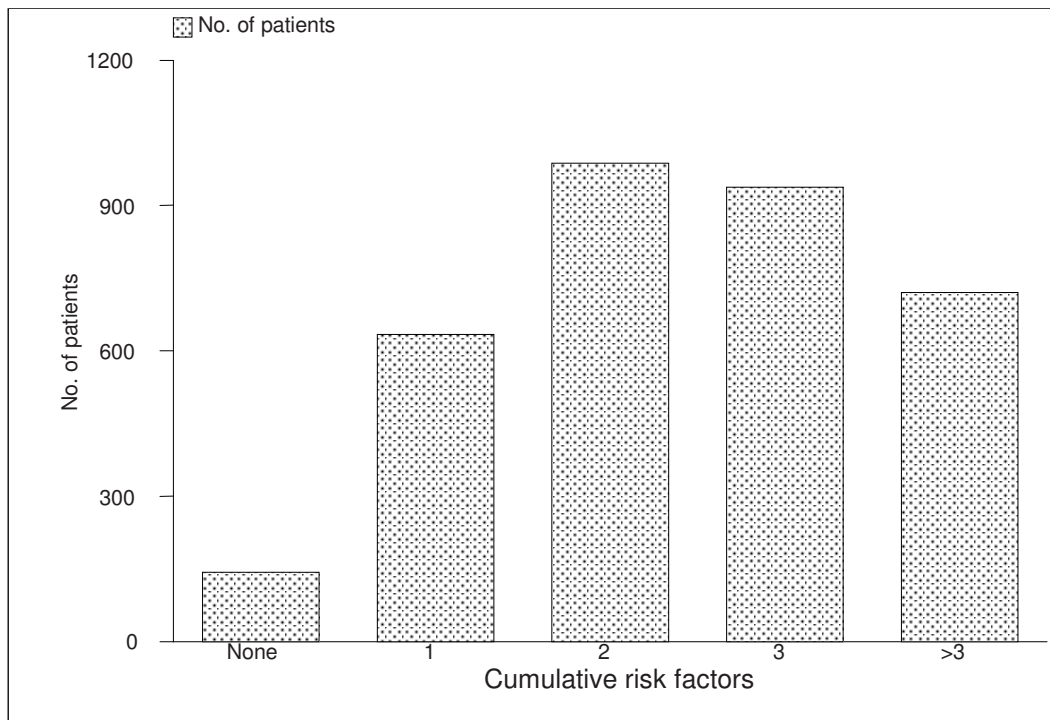


Table 2.6 Summary of type of cardiac presentation for patients with ACS, Malaysia 2006

	Total=3422
Acute coronary syndrome stratum, no. %	
• STEMI	1445 (42)
• NSTEMI	1132 (33)
• UA	845 (25)

Note: Percentage is to the nearest decimal point.

Figure 2.6 Stratum distribution for patients with ACS, Malaysia 2006

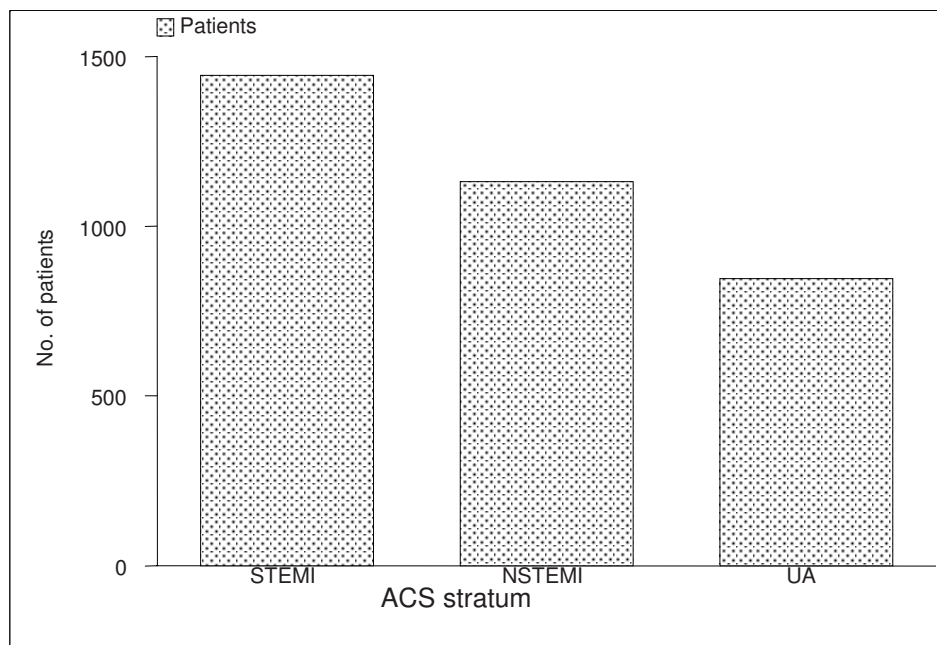


Table 2.7 Characteristics of patients with ACS by ACS stratum, Malaysia 2006

	STEMI N=1445	NSTEMI N=1132	UA N=845
1. DEMOGRAPHICS			
1.1 Age, years			
• Mean, SD	56 (12)	62 (11)	60 (11)
• Median (min, max)	56 (21, 93)	63 (23, 100)	60 (32, 92)
1.2 Age group, no. %			
• 20 - <30	22 (2)	1 (0)	0 (0)
• 30 - <40	91 (6)	27 (2)	25 (3)
• 40 - <50	343 (24)	139 (12)	139 (16)
• 50 - <60	460 (32)	330 (29)	264 (31)
• 60 - <70	318 (22)	334 (30)	229 (27)
• 70 - <80	180 (12)	244 (22)	147 (17)
• ≥80	31 (2)	57 (5)	41 (5)
1.3 Gender, no. %			
• Male	1230 (85)	779 (69)	560 (66)
• Female	215 (15)	353 (31)	285 (34)
1.4 Ethnic group, no. %			
• Malay	780 (54)	514 (45)	390 (46)
• Chinese	301 (21)	265 (23)	220 (26)
• Indian	286 (20)	303 (27)	210 (25)
• Orang Asli	0 (0)	0 (0)	0 (0)
• Kadazan	1 (0)	1 (0)	0 (0)
• Melanau	0 (0)	0 (0)	0 (0)
• Murut	0 (0)	0 (0)	0 (0)
• Bajau	1 (0)	0 (0)	0 (0)
• Bidayuh	16 (1)	9 (1)	3 (0)
• Iban	21 (1)	19 (2)	8 (1)
• Other Malaysian	12 (1)	14 (1)	11 (1)
• Foreigner	27 (2)	7 (1)	3 (0)
2. OTHER CORONARY RISK FACTORS			
2.1 Smoking, no. %			
• Never	417 (29)	551 (49)	402 (48)
• Former (quit >30 days)	272 (19)	295 (26)	238 (28)
• Current (any tobacco use within last 30 days)	723 (50)	259 (23)	156 (18)
• Unknown	33 (2)	27 (2)	49 (6)
2.2 Family history of premature cardiovascular disease, no. %			
• Yes	168 (12)	127 (11)	109 (13)
• No	742 (51)	550 (49)	392 (46)
• Not known	535 (37)	455 (40)	344 (41)

	STEMI N=1445	NSTEMI N=1132	UA N=845
2.3 Antropometric			
BMI			
• N	831	698	397
• Mean, SD	25.69 (4.27)	25.45 (4.37)	26.46 (4.81)
• Median, (min, max)	25.14 (13.15, 60.39)	25.05 (14.87, 59.94)	25.806 (14.872, 45.986)
BMI, kg/m², no. %			
• <18.5	21 (3)	24 (3)	13 (3)
• 18.5-23	181 (22)	174 (25)	71 (18)
• > 23	629 (76)	500 (72)	313 (79)
WHR			
• N	643	454	297
• Mean, SD	0.97 (0.08)	0.97 (0.09)	0.96 (0.10)
• Median, (min, max)	0.96 (0.54, 1.63)	0.96 (0.67, 1.61)	0.96 (0.46, 1.85)
WHR, no. %			
• Men	550	329	212
• ≤ 1.0	405 (74)	230 (70)	151 (71)
• >1.0	145 (26)	99 (30)	61 (29)
• Women	93	125	85
• ≤ 0.85	9 (10)	18 (14)	8 (9)
• >0.85	84 (90)	107 (86)	77 (91)
Waist circumference, cm			
• N	690	494	318
• Mean, SD	88.8 (14.1)	89.6 (14.9)	91.7 (14.1)
• Median, (min, max)	90.0 (36.0, 131.0)	90.0 (36.0, 160.0)	92.0 (37.5, 152.0)
Waist circumference, cm, no. %			
• Men	592	350	220
• ≤ 90	307 (52)	184 (53)	95 (43)
• > 90	285 (48)	166 (47)	125 (57)
• Women	98	144	98
• ≤ 80	20 (20)	30 (21)	18 (18)
• > 80	78 (80)	114 (79)	80 (82)
2.4 Co-morbidity			
Dyslipidaemia, no. %			
• Yes	278 (19)	464 (41)	389 (46)
• No	458 (32)	247 (22)	197 (23)
• Not known	709 (49)	421 (37)	259 (31)
Hypertension, no. %			
• Yes	681 (47)	789 (70)	614 (73)
• No	433 (30)	202 (18)	151 (18)
• Not known	331 (23)	141 (12)	80 (9)

	STEMI N=1445	NSTEMI N=1132	UA N=845
Diabetes, no. %			
• Yes	525 (36)	579 (51)	393 (47)
• No	538 (37)	364 (32)	324 (38)
• Not known	382 (26)	189 (17)	128 (15)
Fasting blood glucose, mmol/L			
• N	1149	812	600
• Mean (SD)	8.7 (4.0)	8.0 (4.0)	7.4 (3.7)
• Median (min, max)	7.1 (3.2, 29.8)	6.6 (3.0, 27.8)	6.1 (3.1, 29.9)
Myocardial infarction history, no. %			
• Yes	144 (10)	216 (19)	202 (24)
• No	876 (61)	579 (51)	392 (46)
• Not known	425 (29)	337 (30)	251 (30)
Documented CAD > 50% stenosis, no. %			
• Yes	74 (5)	230 (20)	204 (24)
• No	851 (59)	527 (47)	356 (42)
• Not known	520 (36)	375 (33)	285 (34)
Chronic angina (onset more than 2 weeks ago), no. %			
• Yes	103 (7)	189 (17)	210 (25)
• No	920 (64)	651 (58)	441 (52)
• Not known	422 (29)	292 (26)	194 (23)
New onset angina (less than 2 weeks), no. %			
• Yes	628 (43)	538 (48)	366 (43)
• No	479 (33)	378 (33)	303 (36)
• Not known	338 (23)	216 (19)	176 (21)
Heart failure, no. %			
• Yes	48 (3)	153 (14)	83 (10)
• No	1008 (70)	724 (64)	557 (66)
• Not known	389 (27)	255 (23)	205 (24)
Chronic lung disease, no. %			
• Yes	34 (2)	57 (5)	39 (5)
• No	1016 (70)	810 (72)	605 (72)
• Not known	395 (27)	265 (23)	201 (24)
Renal disease, no. %			
• Yes	58 (4)	142 (13)	53 (6)
• No	989 (68)	729 (64)	587 (69)
• Not known	398 (28)	261 (23)	205 (24)
Cerebrovascular disease, no. %			
• Yes	46 (3)	65 (6)	38 (4)
• No	1011 (70)	806 (71)	603 (71)
• Not known	388 (27)	261 (23)	204 (24)

	STEMI N=1445	NSTEMI N=1132	UA N=845
Peripheral vascular disease, no. %			
• Yes	4 (0)	25 (2)	8 (1)
• No	1040 (72)	830 (73)	622 (74)
• Not known	401 (28)	277 (24)	215 (25)
None of the above, no. %			
• Yes	125 (9)	36 (3)	18 (2)
• No	1320 (91)	1096 (97)	827 (98)
• Not known	0 (0)	0 (0)	0 (0)
Coronary artery disease*, no. %			
• Yes	779 (54)	787 (70)	633 (75)
• No	306 (21)	146 (13)	80 (9)
• Not known	360 (25)	199 (18)	132 (16)

**Coronary artery disease is defined as "Yes" on any of the following co-morbidities: 1) History of myocardial infarction,, 2) Documented CAD >50% stenosis, 3) Chronic angina (onset more than 2 weeks ago), 4) New onset angina (less than 2 weeks).*

Note: Percentage is to the nearest decimal point.

Figure 2.7a Age group (years) distribution for patients with ACS by ACS stratum, Malaysia 2006

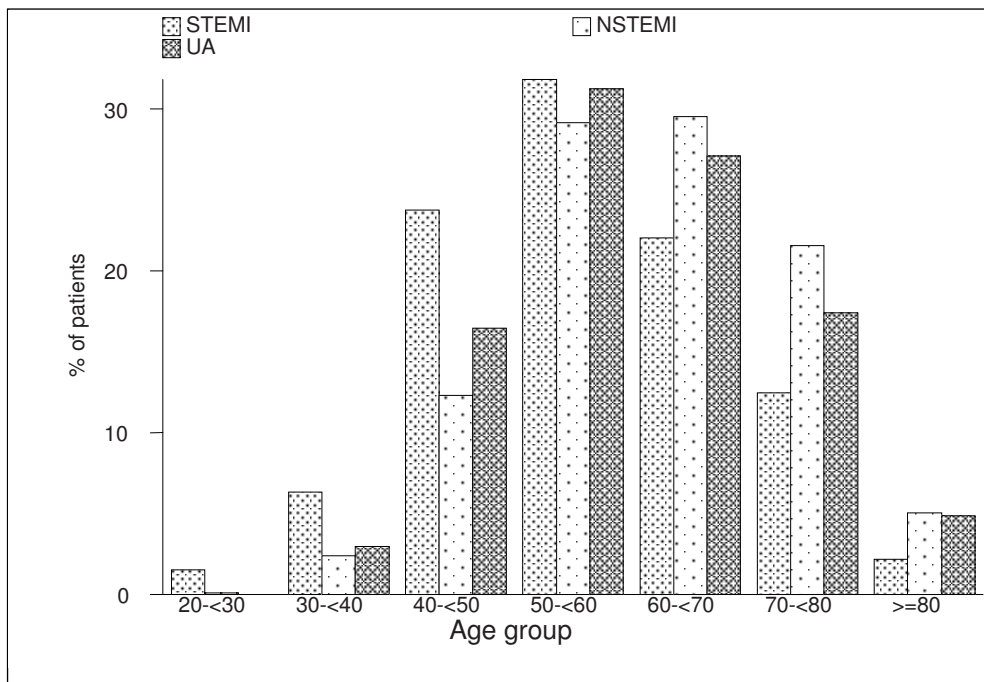


Figure 2.7b Gender distribution for patients with ACS by ACS stratum, Malaysia 2006

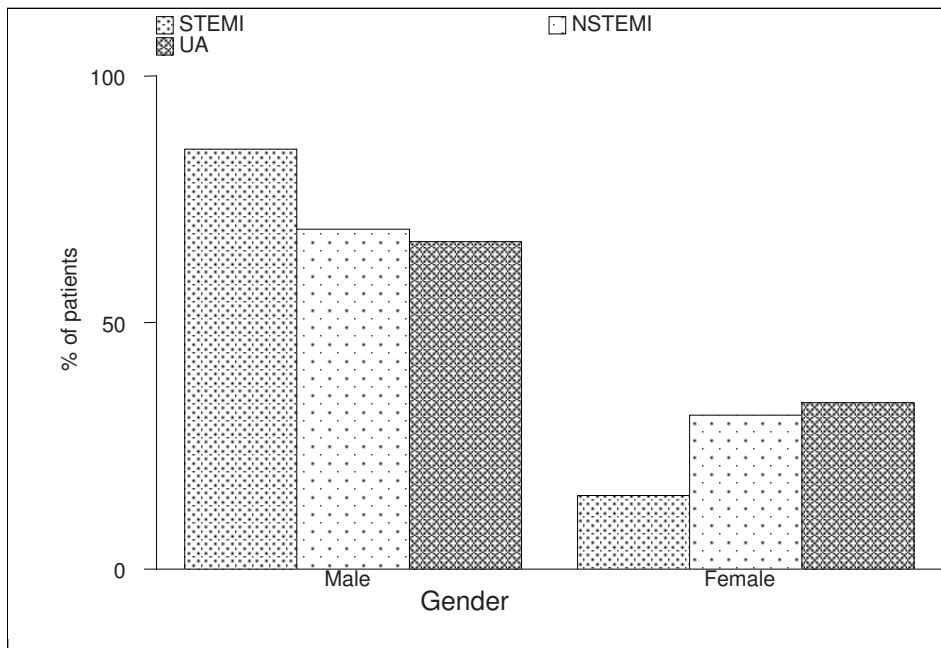


Figure 2.7c Ethnic group distribution for patients with ACS by ACS stratum, Malaysia 2006

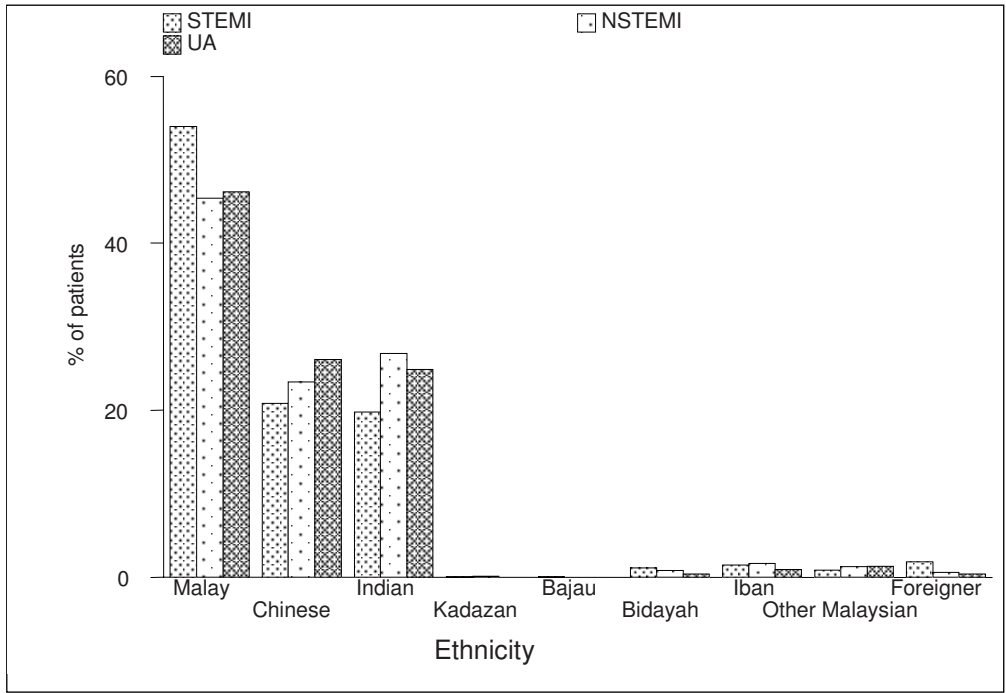


Figure 2.7d Smoking status for patients with ACS by ACS stratum, Malaysia 2006

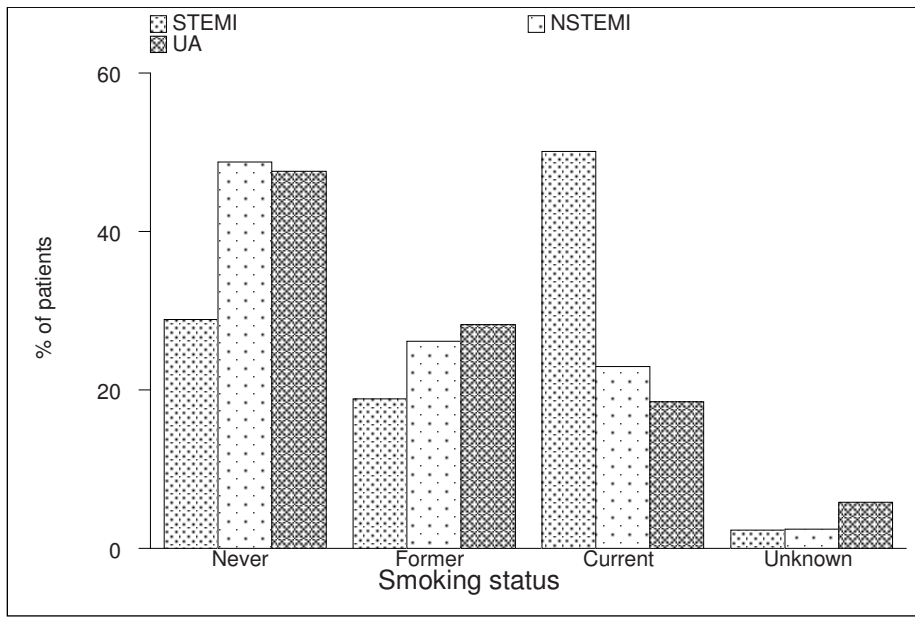


Figure 2.7e Family history of premature cardiovascular disease for patients with ACS by ACS stratum, Malaysia 2006

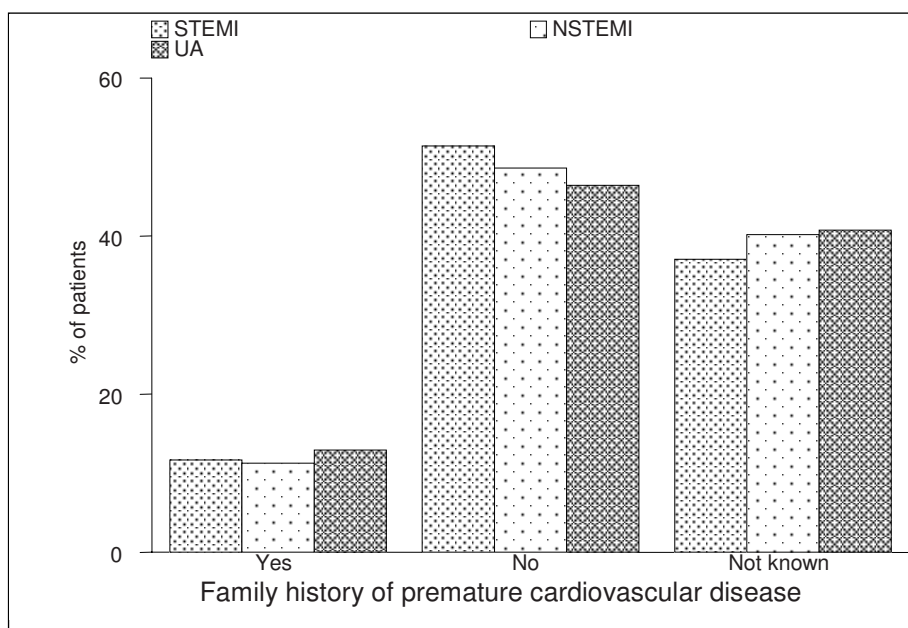


Figure 2.7f BMI for patients with ACS by ACS stratum, Malaysia 2006

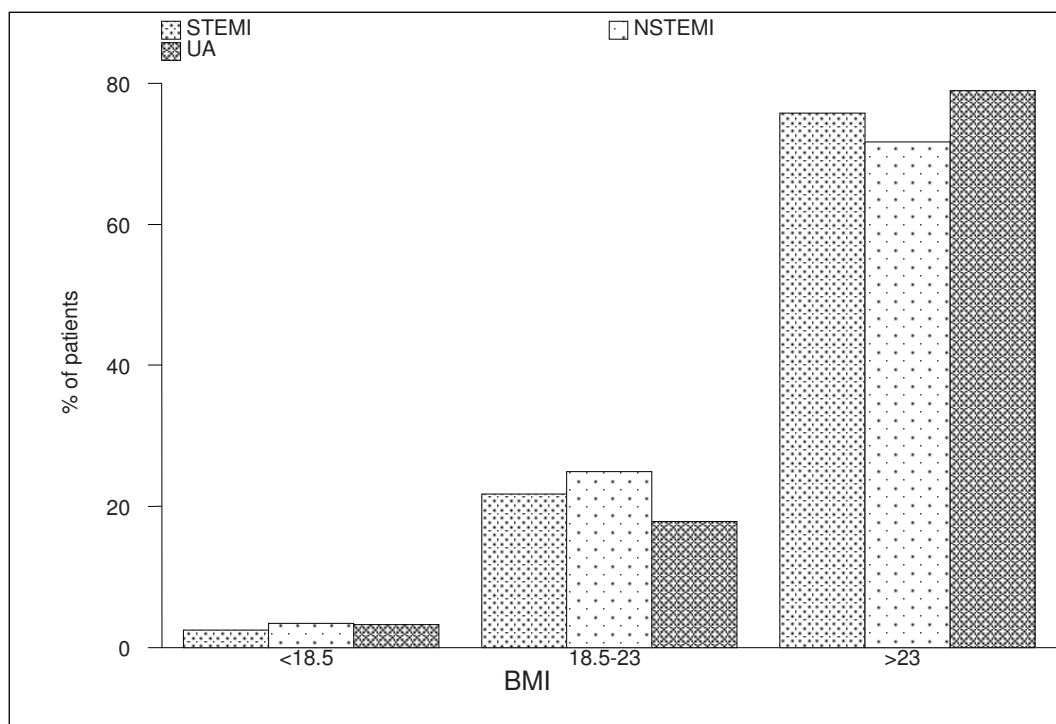


Figure 2.7g WHR for patients with ACS by ACS stratum, Malaysia 2006

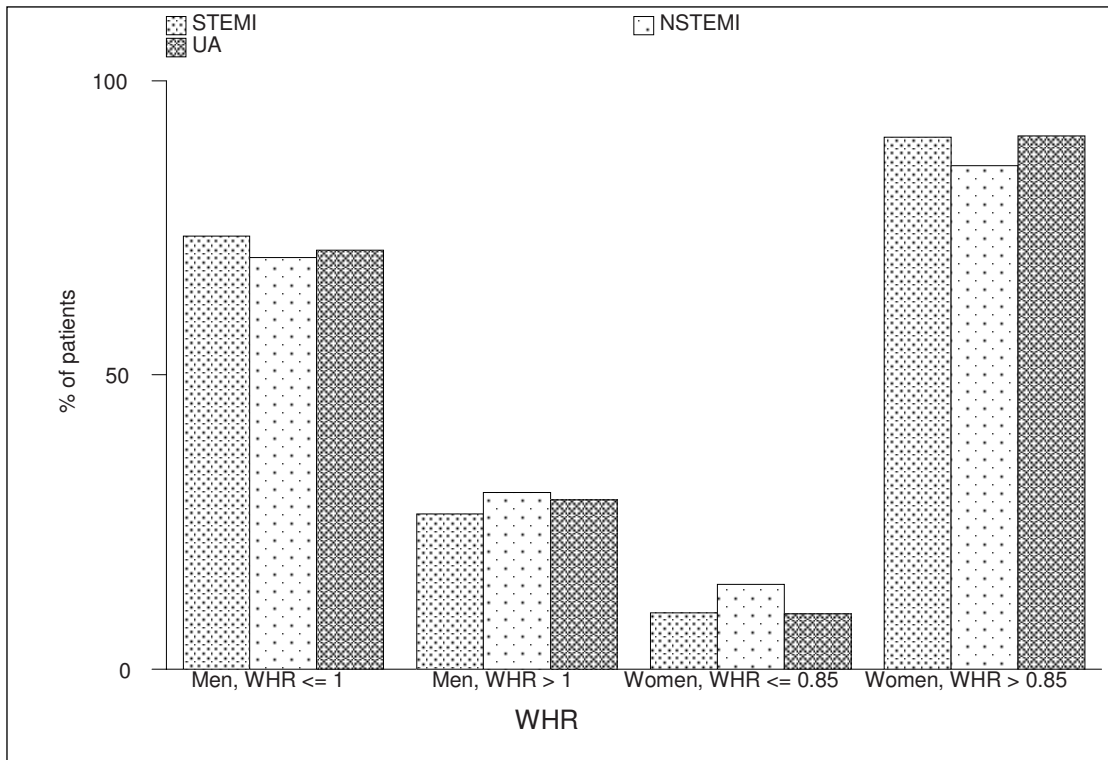


Figure 2.7h Waist circumference (cm) for patients with ACS by ACS stratum, Malaysia 2006

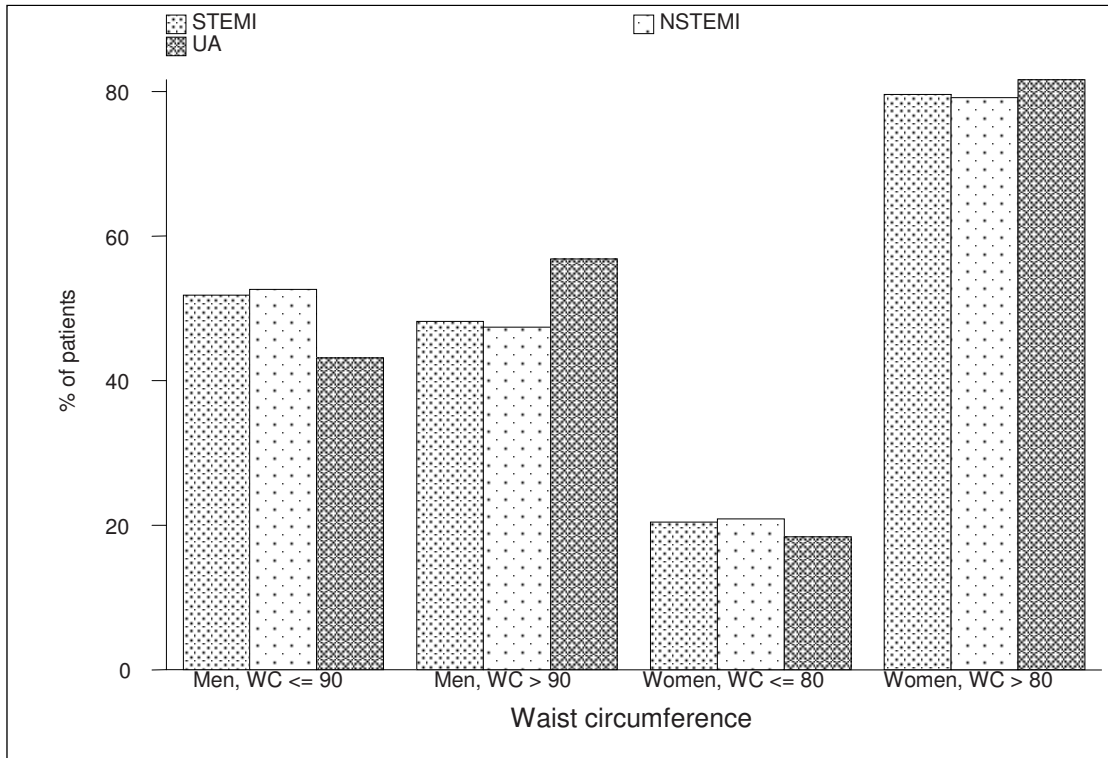
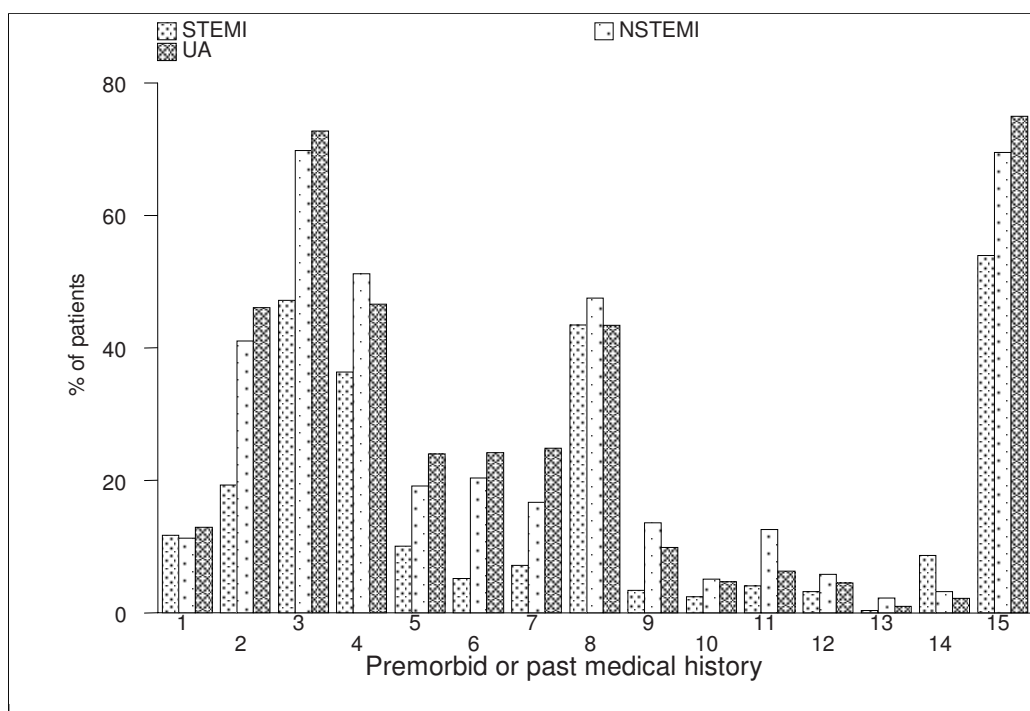


Figure 2.7i Co-morbidities for patients with ACS by ACS stratum, Malaysia 2006



	Pre-morbid or past medical history
1	Family history of premature cardiovascular disease
2	Dyslipidaemia
3	Hypertension
4	Diabetes
5	Myocardial infarction history
6	Documented CAD > 50% stenosis
7	Chronic angina (onset more than 2 weeks ago)
8	New onset angina (less than 2 weeks)
9	Heart failure
10	Chronic lung disease
11	Renal disease
12	Cerebrovascular disease
13	Peripheral vascular disease
14	None of the above
15	Coronary artery disease

Table 2.7.1 Age-gender distribution of patients with ACS by ACS stratum, Malaysia 2006

Gender	Age group	ACS stratum					
		STEMI		NSTEMI		UA	
		No.	%	No.	%	No.	%
Male	20 - <30	21	2	1	0	0	0
	30 - <40	87	7	23	3	21	4
	40 - <50	321	26	122	16	98	18
	50 - <60	414	34	274	35	200	36
	60 - <70	256	21	218	28	142	25
	70 - <80	113	9	112	14	81	14
	≥80	18	1	29	4	18	3
	Total	1230	100	779	100	560	100
Female	20 - <30	1	0	0	0	0	0
	30 - <40	4	2	4	1	4	1
	40 - <50	22	10	17	5	41	14
	50 - <60	46	21	56	16	64	22
	60 - <70	62	29	116	33	87	31
	70 - <80	67	31	132	37	66	23
	≥80	13	6	28	8	23	8
	Total	215	100	353	100	285	100

Note: Percentage is to the nearest decimal point.

Figure 2.7.1a Age-gender distribution for male patients with ACS by ACS stratum, Malaysia 2006

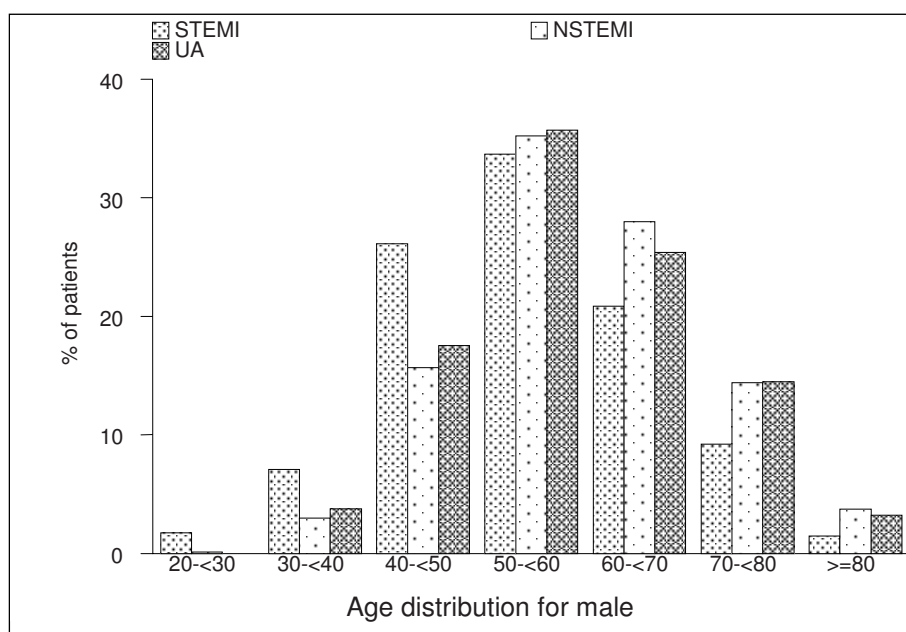
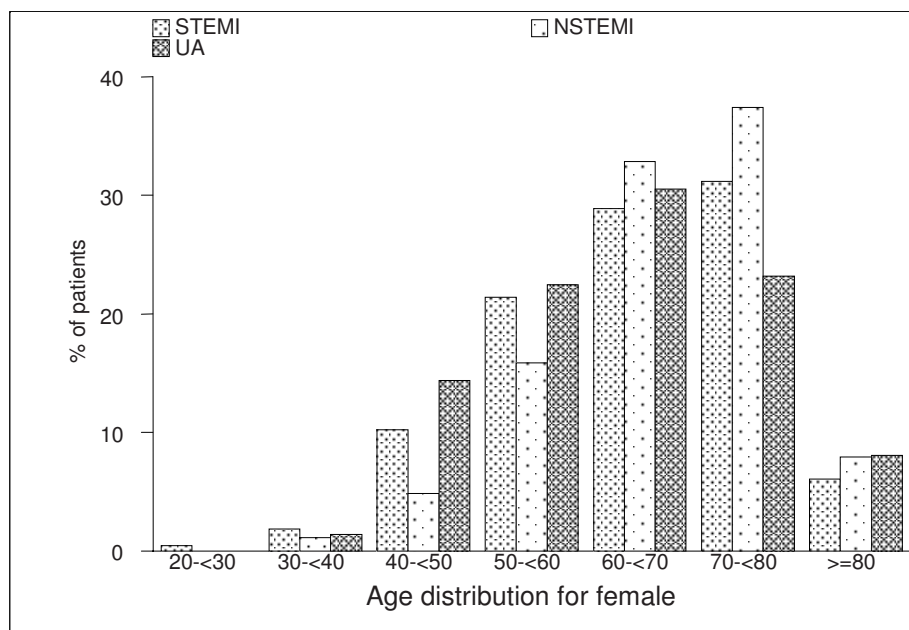


Figure 2.7.1b Age-gender distribution for female patients with ACS by ACS stratum, Malaysia 2006



CHAPTER 3

CLINICAL PRESENTATION AND INVESTIGATION

Saravanan Krishnan

Lu Hou Tee

Lee Chuey Yan

This chapter presents the results of clinical presentation and investigations of acute coronary syndrome patients registered in the ACS registry.

As discussed in Chapter 2, patients were classified according to ACS presentations of STEMI, NSTEMI and UA. A higher percentage of patients presented as STEMI (42%), followed by NSTEMI (32%) and UA (25%). The lower percentage of patients presented as NSTEMI and UA in this cohort could be due to underreporting as some of these patients were admitted to the general medical wards or non-cardiac wards.

There is no significant difference in the number of distinct episodes of angina in the past 24 hours among patients with STEMI (70%), NSTEMI (62%) and UA (65%) (Table 3.1).

The Killip classification was used to categorize the presence and severity of heart failure at the time of initial presentation ¹. The results show that 56% of STEMI patients were categorized in Killip class I and were followed by Killip class II (20%); both Killip class III and Killip class IV collectively made up 9%. Killip classification was also used among the NSTEMI and UA patients as it was documented as a powerful independent predictor of all-cause mortality in NSTEMI patients ¹. For NSTEMI and UA patients, both class III and IV made up 10 % and 2% respectively.

The result shows that more than 90% of patients in each category had gone through either one of the following cardiac marker tests, which are Creatine Kinase-MB, Creatine Kinase, Troponin T or Troponin I.

Among STEMI patients, 85% (N=801) of them showed positive results for CK-MB test, 75% (N=1325) for CK; 99% (N=183) and 93% (N=58) for Troponin T and Troponin I, respectively. While among NSTEMI patients, 52% (N=874) showed positive results for CK-MB test, 38% (N=970) for CK; 98% (N=446) and 93% (N=167) for Troponin T and Troponin I, respectively. Troponin testing was conducted on only a small number of patients, as a few hospitals in the NCVD-ACS registry have the test available.

Among the younger patients (aged from 20 to less than 40 years old) who presented with acute coronary syndrome, majority of them presented with STEMI (68%). (Table 3.2.1) The incidence of systolic hypertension among older patients was found to be higher as expected. The result also shows that the peak cardiac biomarkers were higher among the younger patients. The total cholesterol, LDL and triglycerides levels were higher among the younger patients. This is a reflection of the dietary trends among the young. As expected, a higher percentage of older patients presented with Killip class II (21%) and Killip class III and IV (9%) compared to Killip class II (7%) and Killip class III and IV (3%) in the younger age group.

The incidence of STEMI was higher in men (48%) compared to women (25%). However, more women were reported to have NSTEMI (41%) and UA (33%) compared to men, who accounted for 30% and 22% respectively.

The Left Ventricular Ejection Fraction (LV EF) is poorer among the elderly and diabetic patients. These patients were likely to have chronic diffuse ischemic disease. A large proportion of patients do not know their pre-morbid dyslipidemia status. There was a higher incidence of STEMI among patients who do not have diabetes, hypertension and elevated lipid levels. However, this finding is most likely misleading, as a large proportion of patients in this group remained undiagnosed for their co-morbidities.

Summary Points:

- Out of the 3422 patients with ACS 42% presented with STEMI, 33% with NSTEMI and 25% UA.
- Most of the patients in STEMI group were in Killip class I and II, 5% of STEMI patients were in Killip class IV compared to 3% in NSTEMI group and none in UA.
- There were no significant differences in baseline blood pressure (BP), heart rate (HR), cholesterol or blood sugar levels between the three groups.
- STEMI patients have a lower number of established risk factors with respect to dyslipidaemia, hypertension and diabetes mellitus compared to NSTEMI and UA patients.

Reference:

- 1 Khot U.N., J. Gang, Moliterno D.J. et.al. 2003. Prognostics Importance of Physical Examination for Heart Failure in Non-ST Elevation Acute Coronary Syndromes: The enduring value of Killip classification. *Journal of the American Medical Association (JAMA)*.290: 2174-2181

Table 3.1 Cardiac presentations of patients with ACS by ACS stratum, Malaysia 2006

	STEMI N=1445	NSTEMI N=1132	UA N=845
Systolic blood pressure, mmHg			
• N	1428	1116	836
• Mean, SD	134 (28)	145 (30)	146 (28)
• Median, (min, max)	130 (60, 230)	142 (66, 230)	142 (60, 224)
Diastolic blood pressure, mmHg			
• N	1402	1087	823
• Mean, SD	80 (17)	82 (17)	82 (15)
• Median, (min, max)	80 (26, 120)	82 (19, 120)	82 (34, 120)
Heart rate at presentation, beats/min			
• N	1434	1122	838
• Mean, SD	82 (21)	86 (23)	81 (19)
• Median, (min, max)	80 (30, 180)	84 (29, 180)	79 (40, 166)
Number of distinct episodes of angina in past 24 hours, no. %			
• 0-2	933 (65)	626 (55)	459 (54)
• >2	68 (5)	74 (7)	93 (11)
• Missing	444 (31)	432 (38)	293 (35)
Killip classification code, no. %			
• I	802 (56)	489 (43)	377 (45)
• II	288 (20)	206 (18)	98 (12)
• III	62 (4)	76 (7)	15 (2)
• IV	66 (5)	30 (3)	4 (0)
• Not stated/inadequately described	227 (16)	331 (29)	351 (42)
Patients with any cardiac marker done, no. %	1402 (97)	1114 (98)	762 (90)
Peak CK-MB, Unit/L, no. %	801	874	397
• >25	681 (85)	457 (52)	0 (0)
Peak CK, Unit/L, no. %	1325	970	722
• >2x reference upper limits	999 (75)	364 (38)	0 (0)
Peak TnT, no. %	183	446	27
• Positive or >0.01	181 (99)	436 (98)	0 (0)
Peak TnI, no. %	58	167	55
• Positive or > reference upper limit/0.04	54 (93)	156 (93)	0 (0)
Total cholesterol, mmol/L			
• N	1175	788	508
• Mean, SD	5.4 (1.3)	5.2 (1.3)	5.1 (1.3)
• Median, (min, max)	5.3 (3.0, 14.3)	5.1 (3.0, 11.0)	5.0 (3.0, 10.8)

	STEMI N=1445	NSTEMI N=1132	UA N=845
HDL-C, mmol/L			
• N	1155	807	514
• Mean, SD	1.13 (0.38)	1.13 (0.37)	1.12 (0.38)
• Median, (min, max)	1.10 (0.50, 4.94)	1.10 (0.50, 4.24)	1.07 (0.50, 4.50)
LDL-C, mmol/L			
• N	1144	793	491
• Mean, SD	3.45 (1.22)	3.16 (1.20)	3.09 (1.19)
• Median, (min, max)	3.40 (1.10, 10.10)	3.03 (1.00, 8.87)	2.99 (1.00, 8.90)
Triglycerides, mmol/L			
• N	1025	687	439
• Mean, SD	2.05 (1.26)	2.06 (1.18)	2.25 (1.54)
• Median, (min, max)	1.70 (1.00, 13.50)	1.70 (1.00, 11.10)	1.80 (1.00, 14.00)
Left ventricular ejection fraction, %			
• N	948	569	199
• Mean, SD	47 (11)	47 (14)	50 (17)
• Median, (min, max)	47 (10, 79)	48 (9, 79)	52 (8, 80)

**** Not all participating centre performed Troponin T or I tests.**

Note: Percentage is to the nearest decimal point.

Figure 3.1.1 Number of distinct angina episodes for patients with ACS by ACS stratum, Malaysia 2006

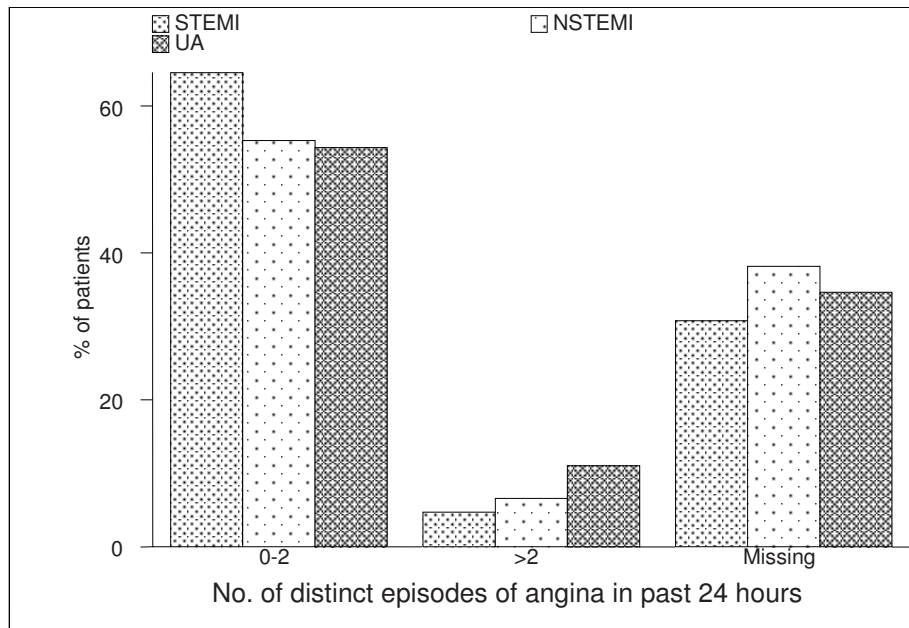


Figure 3.1.2 Killip classification code for patients with ACS by ACS stratum, Malaysia 2006

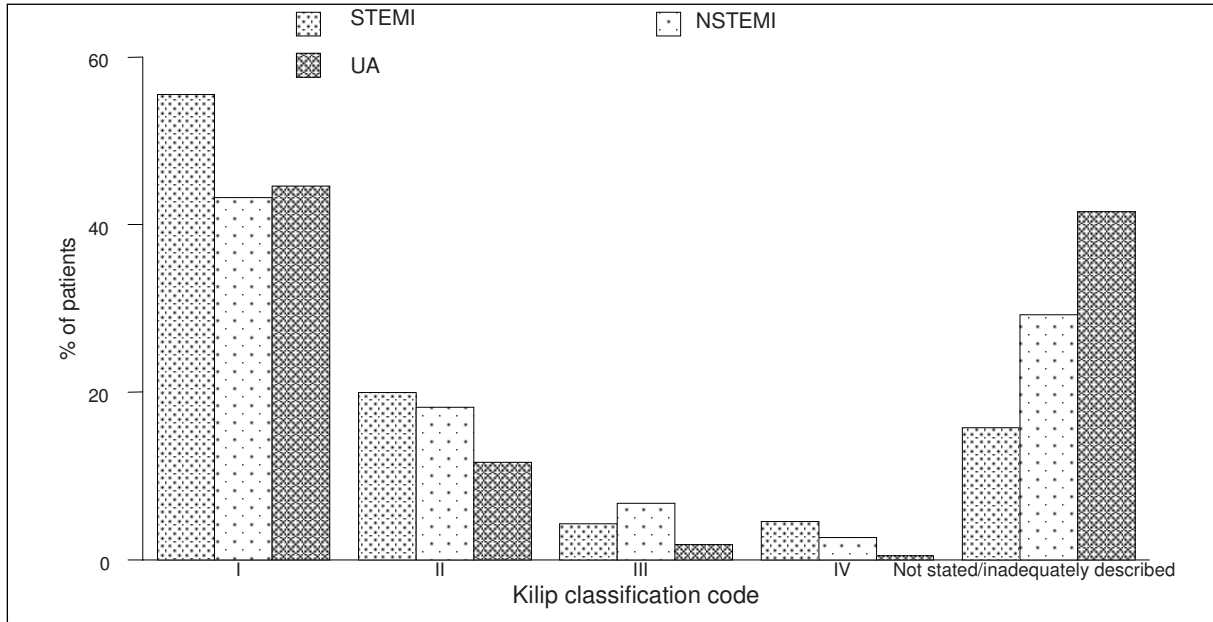


Table 3.2.1 Cardiac presentation of patients with ACS by age group (years), Malaysia 2006

	Age group*		
	Young N=166	Middle-age N=1675	Elderly N=1581
Acute coronary syndrome stratum, no. %			
• STEMI	113 (68)	803 (48)	529 (33)
• NSTEMI	28 (17)	469 (28)	635 (40)
• UA	25 (15)	403 (24)	417 (26)
Systolic blood pressure, mmHg			
• N	165	1657	1558
• Mean, SD	133 (25)	139 (28)	143 (31)
• Median, (min, max)	130 (77, 220)	137 (60, 230)	140 (60, 230)
Diastolic blood pressure, mmHg			
• N	158	1620	1534
• Mean, SD	81 (17)	83 (16)	80 (17)
• Median, (min, max)	80 (38, 118)	82 (35, 120)	80 (19, 120)
Heart rate at presentation, beats/min			
• N	165	1663	1566
• Mean, SD	83 (19)	82 (20)	84 (23)
• Median, (min, max)	82 (44, 153)	80 (30, 171)	82 (29, 180)
Number of distinct episodes of angina in past 24 hours, no. %			
• 0-2	116 (70)	1021 (61)	881 (56)
• >2	9 (5)	118 (7)	108 (7)
• Missing	41 (25)	536 (32)	592 (37)
Killip classification code, no. %			
• I	121 (73)	900 (54)	647 (41)
• II	11 (7)	249 (15)	332 (21)
• III	2 (1)	59 (4)	92 (6)
• IV	4 (2)	47 (3)	49 (3)
• Not stated/inadequately described	28 (17)	420 (25)	461 (29)
Peak CK-MB, Unit/L, no. %	101	994	977
• >25	70 (69)	563 (57)	505 (52)
Peak CK, Unit/L, no. %	144	1511	1362
• >2x reference upper limits	95 (66)	742 (49)	526 (39)
Peak TnT, no. %	23	307	326
• Positive or >0.01	22 (96)	285 (93)	310 (95)
Peak Tnl, no. %	17	112	151
• Positive or > reference upper limit/0.04	13 (76)	77 (69)	120 (79)

	Age group*		
	Young N=166	Middle-age N=1675	Elderly N=1581
Total cholesterol, mmol/L			
• N	143	1257	1071
• Mean, SD	5.7 (1.7)	5.5 (1.3)	5.0 (1.2)
• Median, (min, max)	5.4 (3.1, 14.3)	5.4 (3.0, 10.8)	4.9 (3.0, 11.0)
HDL-C, mmol/L			
• N	141	1268	1067
• Mean, SD	1.08 (0.40)	1.08 (0.35)	1.19 (0.40)
• Median, (min, max)	1.00 (0.60, 3.50)	1.01 (0.50, 4.94)	1.13 (0.50, 4.50)
LDL-C, mmol/L			
• N	141	1218	1069
• Mean, SD	3.67 (1.50)	3.41 (1.22)	3.08 (1.14)
• Median, (min, max)	3.46 (1.50, 10.10)	3.40 (1.00, 9.30)	3.00 (1.00, 8.87)
Triglycerides, mmol/L			
• N	132	1154	865
• Mean, SD	2.39 (1.47)	2.24 (1.36)	1.85 (1.14)
• Median, (min, max)	2.10 (1.00, 13.50)	1.90 (1.00, 13.00)	1.56 (1.00, 14.00)
Left ventricular ejection fraction, %			
• N	96	863	757
• Mean, SD	51 (12)	48 (13)	46 (13)
• Median, (min, max)	50 (12, 75)	50 (8, 80)	45 (10, 80)

*Young is defined as age from 20 to less than 40 years, middle-age is defined as age between 40 to less than 60 years and elderly is defined as 60 years and above

Note: Percentage is to the nearest decimal point.

Figure 3.2.1a Stratum distribution for patients with ACS by age group (years), Malaysia 2006

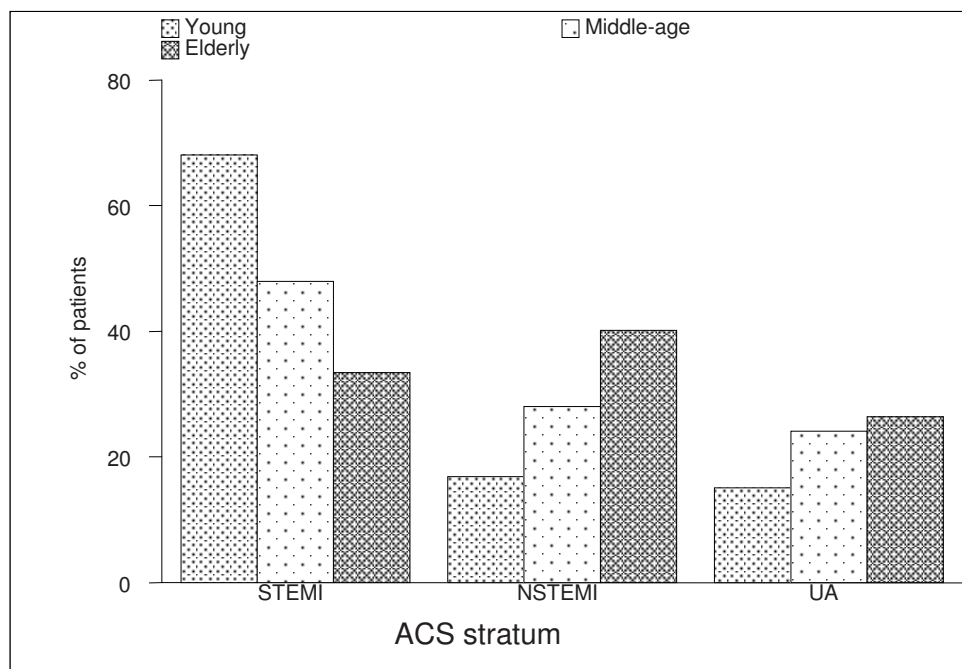


Figure 3.2.1b Number of distinct angina episodes for patients with ACS by age group (years), Malaysia 2006

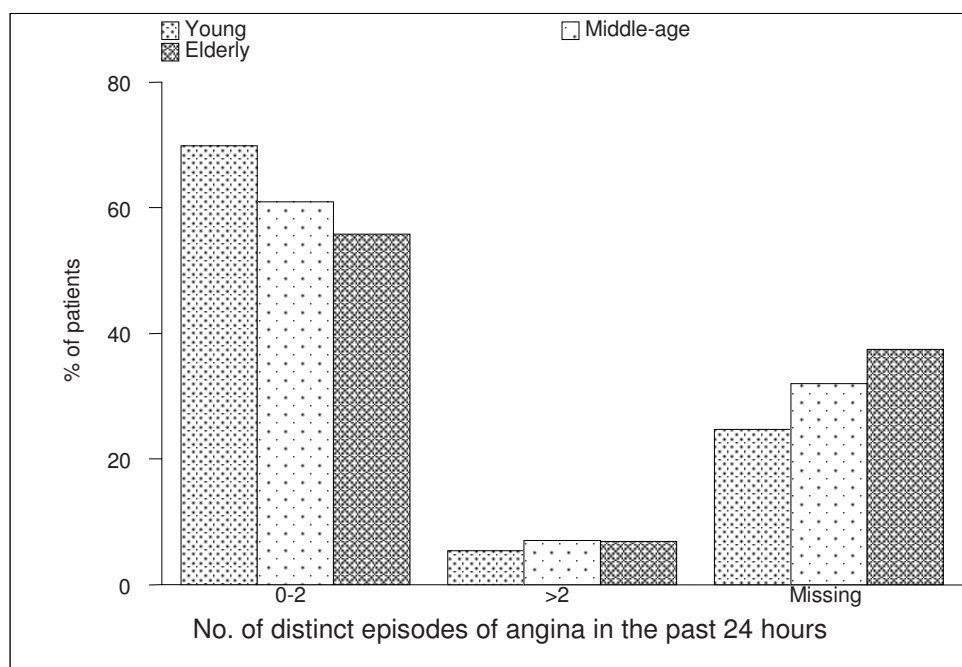


Figure 3.2.1c Killip classification code for patients with ACS by age group (years), Malaysia 2006

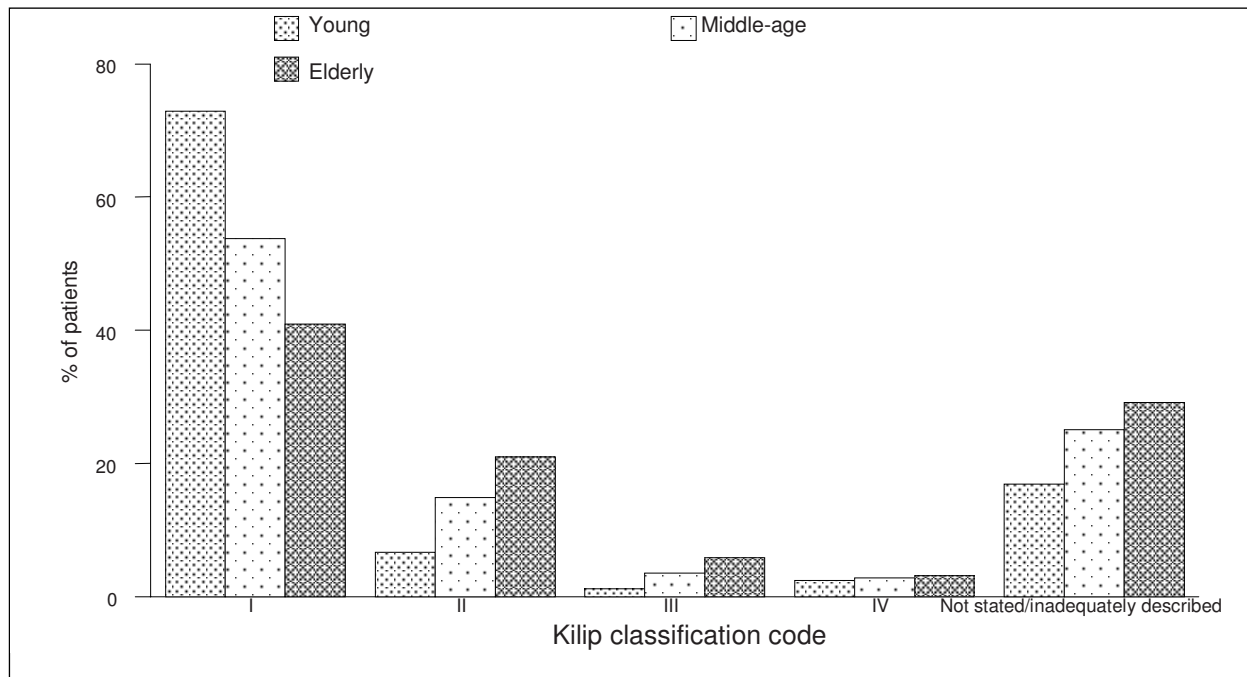


Table 3.2.2 Cardiac presentation of patients with ACS by gender, Malaysia 2006

	Male N=2569	Female N=853
Acute coronary syndrome stratum, no. %		
• STEMI	1230 (48)	215 (25)
• NSTEMI	779 (30)	353 (41)
• UA	560 (22)	285 (33)
Systolic blood pressure, mmHg		
• N	2542	838
• Mean, SD	138 (28)	147 (31)
• Median, (min, max)	136 (60, 230)	144 (60, 229)
Diastolic blood pressure, mmHg		
• N	2484	828
• Mean, SD	82 (17)	80 (17)
• Median, (min, max)	80 (30, 120)	80 (19, 120)
Heart rate at presentation, beats/min		
• N	2548	846
• Mean, SD	82 (21)	88 (22)
• Median, (min, max)	80 (30, 180)	86 (29, 166)
Number of distinct episodes of angina in past 24 hours, no. %		
• 0-2	1564 (61)	454 (53)
• >2	176 (7)	59 (7)
• Missing	829 (32)	340 (40)
Killip classification code, no. %		
• I	1319 (51)	349 (41)
• II	429 (17)	163 (19)
• III	107 (4)	46 (5)
• IV	79 (3)	21 (2)
• Not stated/inadequately described	635 (25)	274 (32)
Peak CK-MB, Unit/L, no. %	1541	531
• >25	908 (59)	230 (43)
Peak CK, Unit/L, no. %	2311	706
• >2x reference upper limits	1147 (50)	216 (31)
Peak TnT, no. %	491	165
• Positive or >0.01	464 (95)	153 (93)
Peak Tnl, no. %	180	100
• Positive or > reference upper limit/0.04	134 (74)	76 (76)

	Male N=2569	Female N=853
Total cholesterol, mmol/L		
• N	1907	564
• Mean, SD	5.3 (1.3)	5.3 (1.4)
• Median, (min, max)	5.1 (3.0, 14.3)	5.2 (3.0, 11.0)
HDL-C, mmol/L		
• N	1904	572
• Mean, SD	1.10 (0.37)	1.22 (0.40)
• Median, (min, max)	1.06 (0.50, 4.94)	1.20 (0.50, 4.24)
LDL-C, mmol/L		
• N	1857	571
• Mean, SD	3.30 (1.20)	3.21 (1.28)
• Median, (min, max)	3.24 (1.00, 10.10)	3.04 (1.00, 8.90)
Triglycerides, mmol/L		
• N	1661	490
• Mean, SD	2.12 (1.31)	1.99 (1.24)
• Median, (min, max)	1.75 (1.00, 13.50)	1.61 (1.00, 14.00)
Left ventricular ejection fraction, %		
• N	1338	378
• Mean, SD	47 (13)	49 (14)
• Median, (min, max)	47 (8, 79)	50 (15, 80)

Note: Percentage is to the nearest decimal point.

Figure 3.2.2a Stratum distribution for patients with ACS by gender, Malaysia 2006

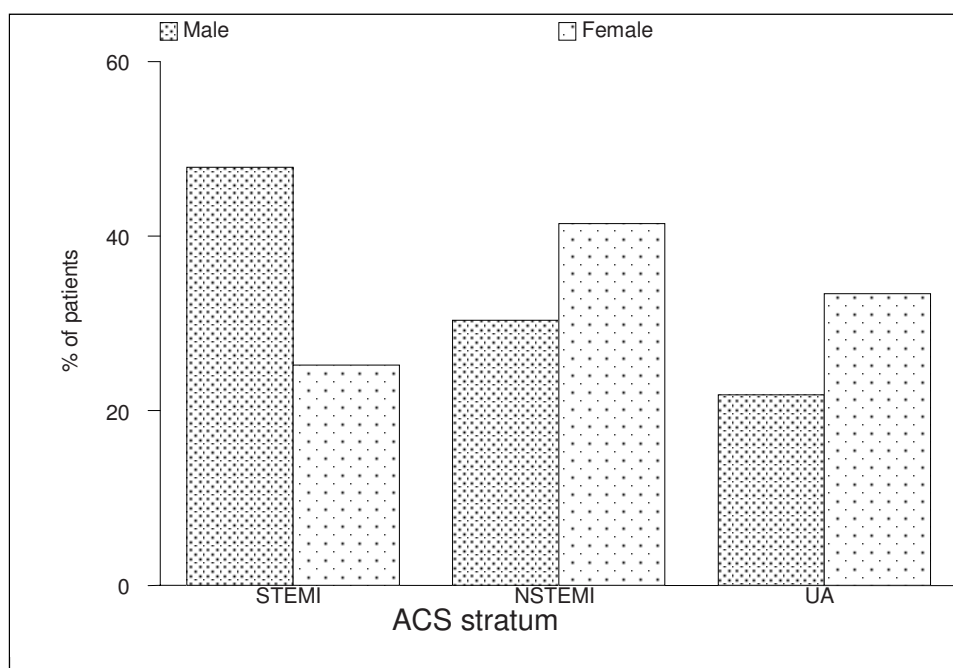


Figure 3.2.2b Number of distinct angina episodes for patients with ACS by gender, Malaysia 2006

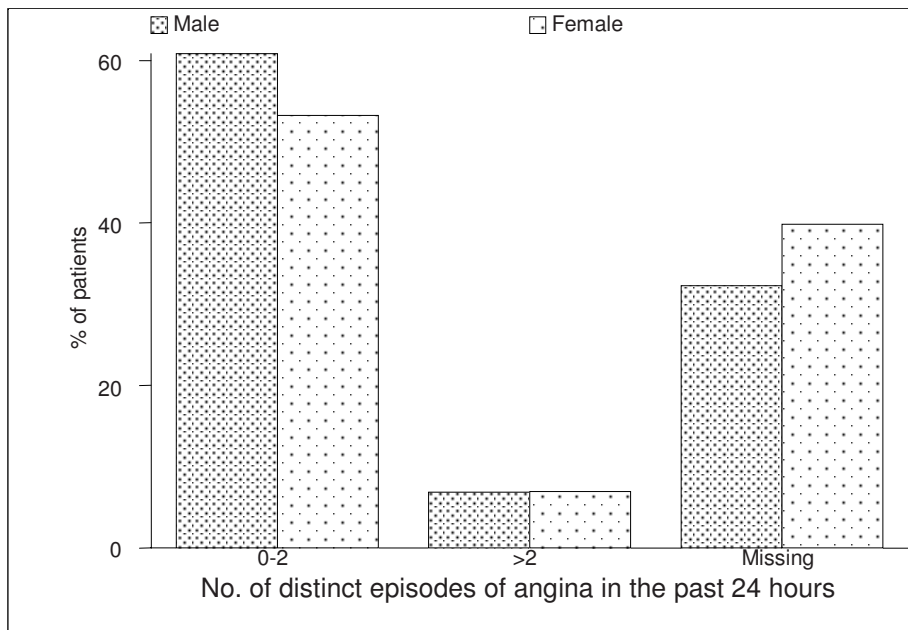


Figure 3.2.2c Killip classification code for patients with ACS by gender, Malaysia 2006

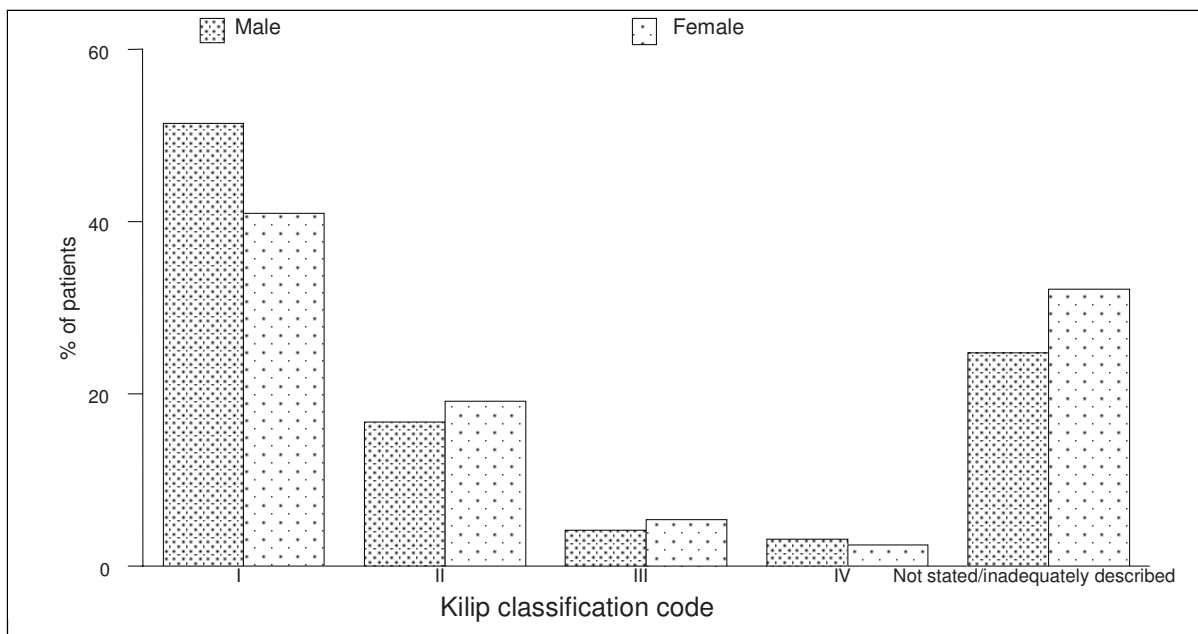


Table 3.2.3 Cardiac presentation of patients with ACS by pre-morbid diabetes, Malaysia 2006

	Pre-morbid diabetes		
	Diabetic N=1497	Non-diabetic N=1226	Not known N=699
Acute coronary syndrome stratum, no. %			
• STEMI	525 (35)	538 (44)	382 (55)
• NSTEMI	579 (39)	364 (30)	189 (27)
• UA	393 (26)	324 (26)	128 (18)
Systolic blood pressure, mmHg			
• N	1475	1215	690
• Mean, SD	142 (30)	140 (29)	136 (28)
• Median, (min, max)	140 (66, 229)	137 (60, 230)	134 (64, 219)
Diastolic blood pressure, mmHg			
• N	1457	1180	675
• Mean, SD	81 (16)	81 (17)	81 (17)
• Median, (min, max)	80 (19, 120)	81 (26, 120)	80 (30, 120)
Heart rate at presentation, beats/min			
• N	1486	1216	692
• Mean, SD	86 (21)	80 (21)	81 (22)
• Median, (min, max)	85 (29, 180)	78 (35, 171)	79 (39, 171)
Number of distinct episodes of angina in past 24 hours, no. %			
• 0-2	862 (58)	669 (55)	487 (70)
• >2	109 (7)	69 (6)	57 (8)
• Missing	526 (35)	488 (40)	155 (22)
Killip classification code, no. %			
• I	700 (47)	619 (50)	349 (50)
• II	290 (19)	193 (16)	109 (16)
• III	84 (6)	39 (3)	30 (4)
• IV	51 (3)	22 (2)	27 (4)
• Not stated/inadequately described	372 (25)	353 (29)	184 (26)
Peak CK-MB, Unit/L, no. %	966	740	366
• >25	469 (49)	421 (57)	248 (68)
Peak CK, Unit/L, no. %	1354	1035	628
• >2x reference upper limits	526 (39)	489 (47)	348 (55)
Peak TnT, no. %	333	207	116
• Positive or >0.01	312 (94)	193 (93)	112 (97)
Peak Tnl, no. %	130	92	58
• Positive or > reference upper limit/0.04	104 (80)	67 (73)	39 (67)

	Pre-morbid diabetes		
	Diabetic N=1497	Non-diabetic N=1226	Not known N=699
Total cholesterol, mmol/L			
• N	1067	884	520
• Mean, SD	5.2 (1.3)	5.3 (1.3)	5.4 (1.3)
• Median, (min, max)	5.0 (3.0, 10.8)	5.3 (3.0, 14.3)	5.3 (3.0, 10.4)
HDL-C, mmol/L			
• N	1076	886	514
• Mean, SD	1.10 (0.39)	1.16 (0.39)	1.14 (0.34)
• Median, (min, max)	1.04 (0.50, 4.90)	1.10 (0.50, 4.94)	1.10 (0.50, 4.20)
LDL-C, mmol/L			
• N	1039	880	509
• Mean, SD	3.10 (1.22)	3.37 (1.21)	3.48 (1.18)
• Median, (min, max)	2.95 (1.00, 9.30)	3.31 (1.00, 10.10)	3.37 (1.02, 8.20)
Triglycerides, mmol/L			
• N	952	745	454
• Mean, SD	2.25 (1.44)	1.96 (1.11)	1.98 (1.23)
• Median, (min, max)	1.82 (1.00, 14.00)	1.67 (1.00, 12.90)	1.62 (1.00, 13.00)
Left ventricular ejection fraction, %			
• N	736	609	371
• Mean, SD	46 (13)	49 (13)	48 (13)
• Median, (min, max)	45 (8, 80)	50 (10, 80)	49 (10, 78)

Note: Percentage is to the nearest decimal point.

Figure 3.2.3a Stratum distribution for patients with ACS by pre-morbid diabetes, Malaysia 2006

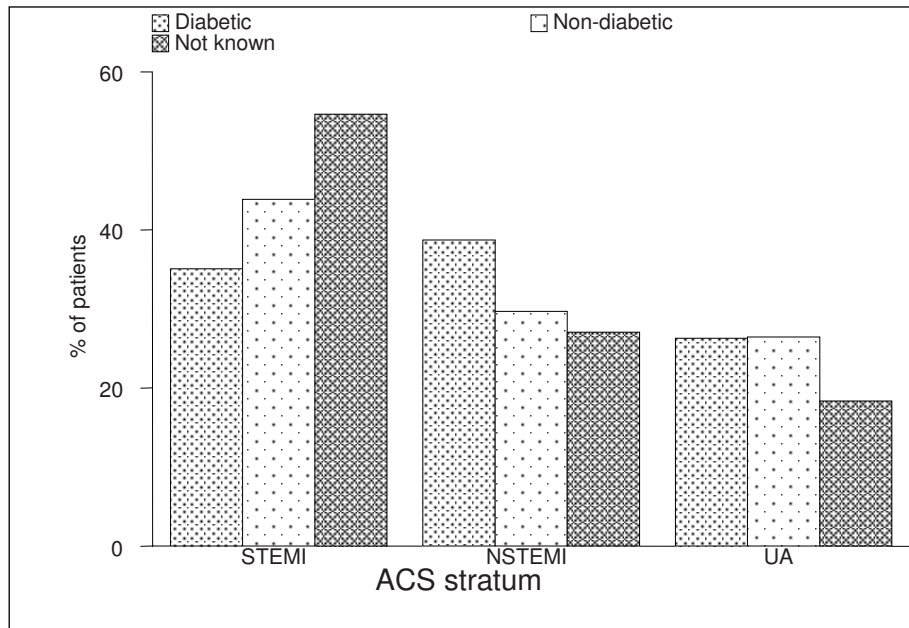


Figure 3.2.3b Number of distinct angina episodes for patients with ACS by pre-morbid diabetes, Malaysia 2006

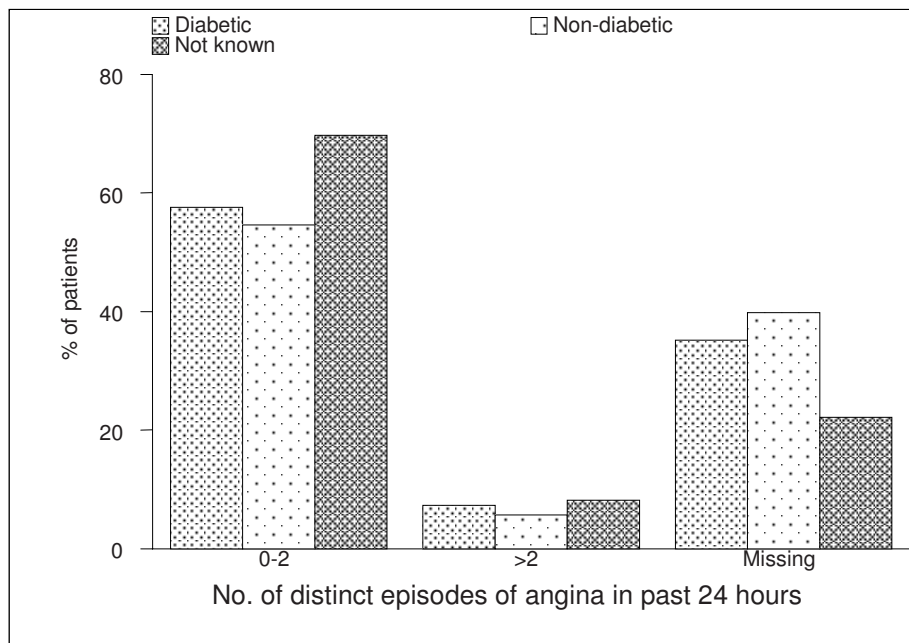


Figure 3.2.3c Killip classification code for patients with ACS by pre-morbid diabetes, Malaysia 2006

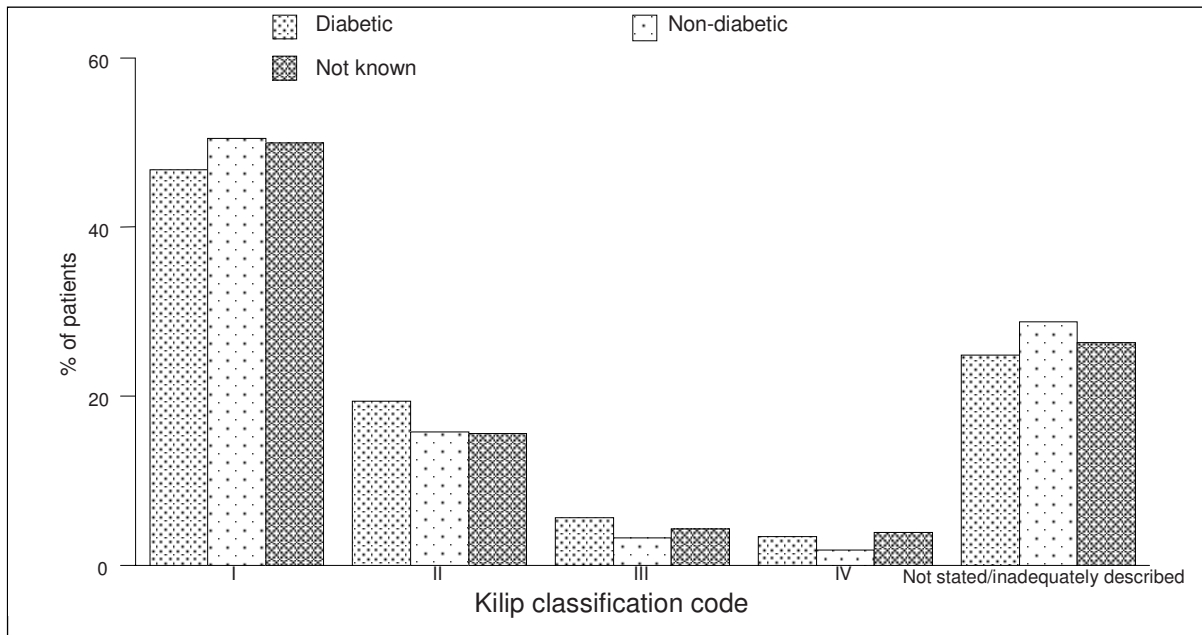


Table 3.2.4 Cardiac presentation of patients with ACS by pre-morbid hypertension, Malaysia 2006

	Pre-morbid hypertension		
	Hypertensive N=2084	Non- hypertensive N=786	Not known N=552
Acute coronary syndrome stratum, no. %			
• STEMI	681 (33)	433 (55)	331 (60)
• NSTEMI	789 (38)	202 (26)	141 (26)
• UA	614 (29)	151 (19)	80 (14)
Systolic blood pressure, mmHg			
• N	2052	784	544
• Mean, SD	146 (30)	132 (26)	131 (26)
• Median, (min, max)	144 (60, 230)	130 (60, 226)	130 (64, 223)
Diastolic blood pressure, mmHg			
• N	1999	773	540
• Mean, SD	83 (17)	78 (16)	79 (17)
• Median, (min, max)	83 (26, 120)	79 (19, 120)	80 (30, 119)
Heart rate at presentation, beats/min			
• N	2069	779	546
• Mean, SD	84 (22)	82 (20)	82 (22)
• Median, (min, max)	82 (29, 180)	80 (36, 157)	80 (39, 171)
Number of distinct episodes of angina in past 24 hours, no. %			
• 0-2	1206 (58)	427 (54)	385 (70)
• >2	165 (8)	36 (5)	34 (6)
• Missing	713 (34)	323 (41)	133 (24)
Killip classification code, no. %			
• I	992 (48)	411 (52)	265 (48)
• II	374 (18)	121 (15)	97 (18)
• III	94 (5)	33 (4)	26 (5)
• IV	45 (2)	24 (3)	31 (6)
• Not stated/inadequately described	579 (28)	197 (25)	133 (24)
Peak CK-MB, Unit/L, no. %	1314	455	303
• >25	633 (48)	287 (63)	218 (72)
Peak CK, Unit/L, no. %	1845	682	490
• >2x reference upper limits	680 (37)	387 (57)	296 (60)
Peak TnT, no. %	429	134	93
• Positive or >0.01	400 (93)	128 (96)	89 (96)

	Pre-morbid hypertension		
	Hypertensive N=2084	Non- hypertensive N=786	Not known N=552
Peak Tnl, no. %	189	43	48
• Positive or > reference upper limit/0.04	134 (71)	37 (86)	39 (81)
Total cholesterol, mmol/L			
• N	1462	599	410
• Mean, SD	5.2 (1.3)	5.4 (1.3)	5.4 (1.4)
• Median, (min, max)	5.1 (3.0, 11.9)	5.4 (3.0, 14.3)	5.3 (3.0, 10.4)
HDL-C, mmol/L			
• N	1481	591	404
• Mean, SD	1.13 (0.40)	1.11 (0.35)	1.13 (0.34)
• Median, (min, max)	1.10 (0.50, 4.94)	1.07 (0.50, 3.50)	1.10 (0.53, 4.20)
LDL-C, mmol/L			
• N	1442	587	399
• Mean, SD	3.15 (1.18)	3.48 (1.24)	3.45 (1.26)
• Median, (min, max)	3.04 (1.00, 8.90)	3.50 (1.10, 10.10)	3.36 (1.02, 8.20)
Triglycerides, mmol/L			
• N	1275	518	358
• Mean, SD	2.12 (1.34)	2.06 (1.19)	2.05 (1.28)
• Median, (min, max)	1.71 (1.00, 14.00)	1.70 (1.00, 12.90)	1.67 (1.00, 13.50)
Left ventricular ejection fraction, %			
• N	980	441	295
• Mean, SD	48 (14)	48 (12)	46 (13)
• Median, (min, max)	48 (8, 80)	47 (20, 77)	46 (10, 79)

Note: Percentage is to the nearest decimal point.

Figure 3.2.4a Stratum distribution for patients with ACS by pre-morbid hypertension, Malaysia 2006

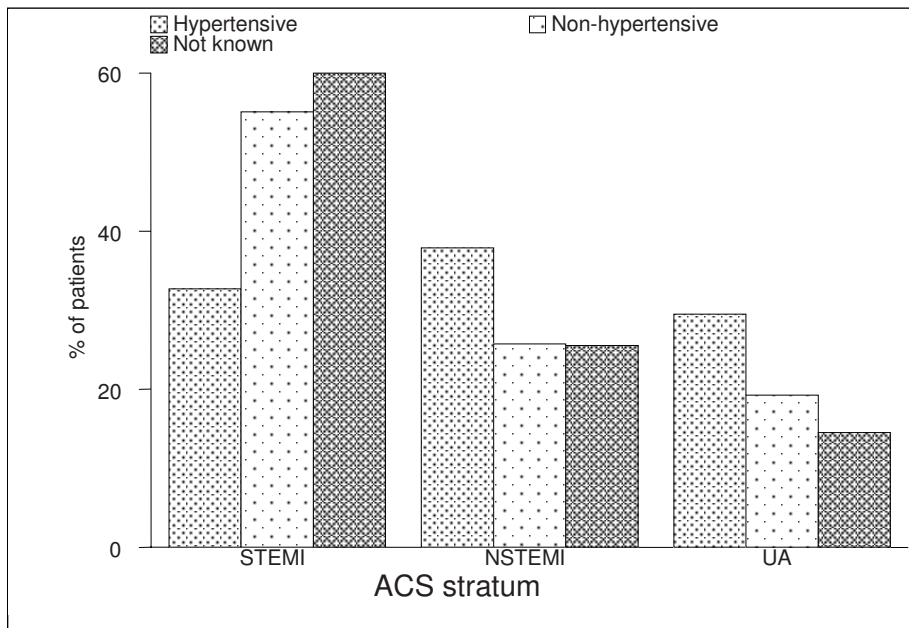


Figure 3.2.4b Number of distinct angina episodes for patients with ACS by pre-morbid hypertension, Malaysia 2006

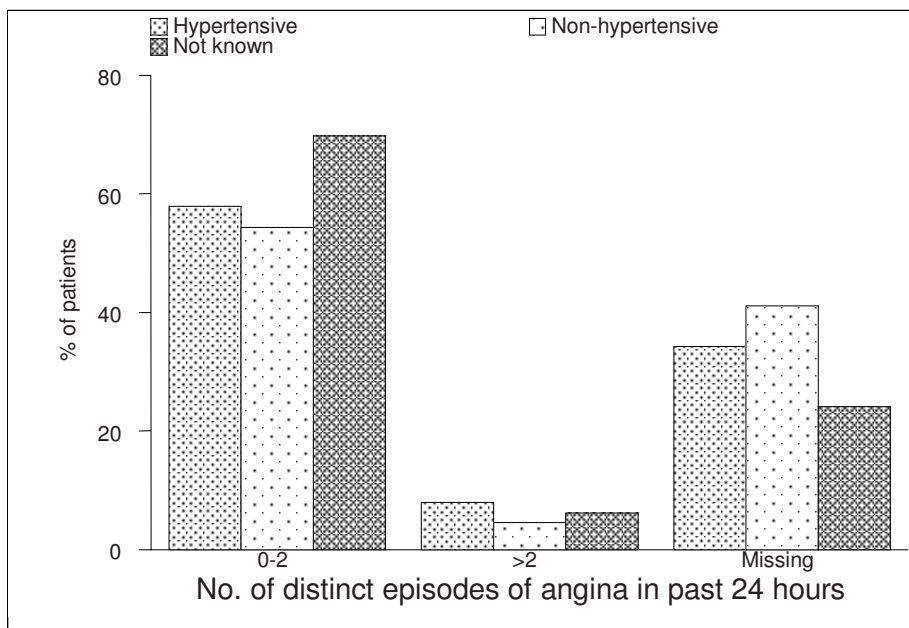


Figure 3.2.4c Killip classification code for patients with ACS by pre-morbid hypertension, Malaysia 2006

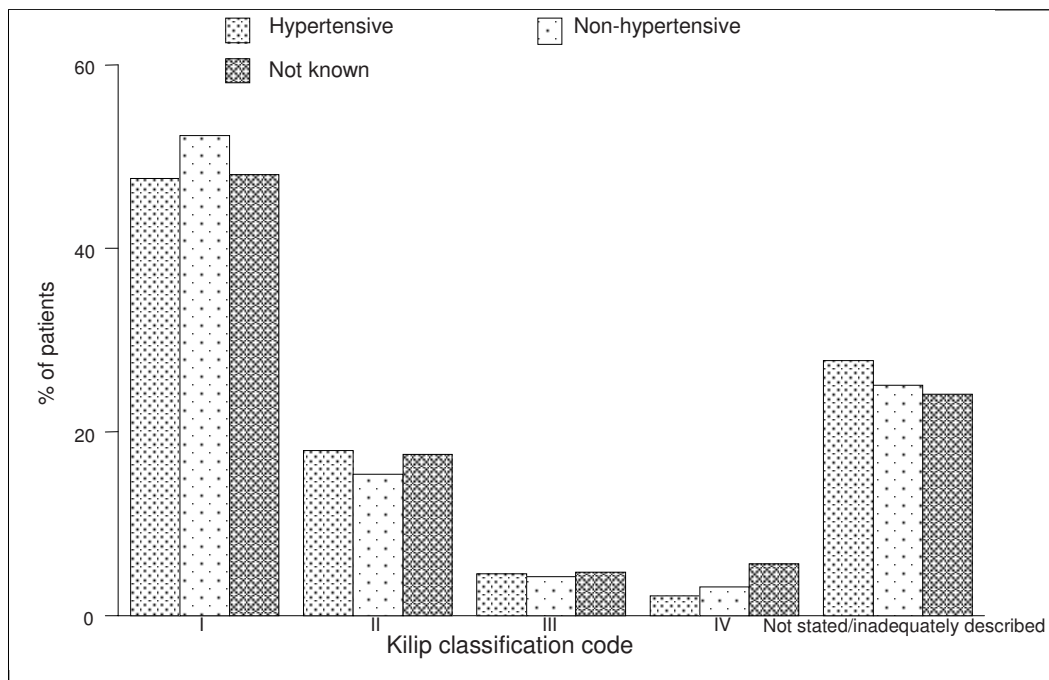


Table 3.2.5 Cardiac presentation of patients with ACS by pre-morbid dyslipidaemia, Malaysia 2006

	Pre-morbid dyslipidaemia		
	Yes N=1131	No N=902	Not known N=1389
Acute coronary syndrome stratum, no. %			
• STEMI	278 (25)	458 (51)	709 (51)
• NSTEMI	464 (41)	247 (27)	421 (30)
• UA	389 (34)	197 (22)	259 (19)
Systolic blood pressure, mmHg			
• N	1123	891	1366
• Mean, SD	144 (29)	140 (28)	138 (30)
• Median, (min, max)	140 (70, 229)	138 (60, 230)	135 (60, 230)
Diastolic blood pressure, mmHg			
• N	1095	876	1341
• Mean, SD	82 (16)	81 (16)	81 (17)
• Median, (min, max)	81 (19, 120)	80 (30, 120)	80 (28, 120)
Heart rate at presentation, beats/min			
• N	1123	894	1377
• Mean, SD	83 (21)	83 (21)	83 (22)
• Median, (min, max)	80 (32, 180)	80 (30, 180)	80 (29, 171)
Number of distinct episodes of angina in past 24 hours, no. %			
• 0-2	652 (58)	475 (53)	891 (64)
• >2	100 (9)	37 (4)	98 (7)
• Missing	379 (34)	390 (43)	400 (29)
Killip classification code, no. %			
• I	573 (51)	426 (47)	669 (48)
• II	208 (18)	126 (14)	258 (19)
• III	48 (4)	43 (5)	62 (4)
• IV	24 (2)	19 (2)	57 (4)
• Not stated/inadequately described	278 (25)	288 (32)	343 (25)
Peak CK-MB, Unit/L, no. %			
• >25	749	514	809
	352 (47)	316 (61)	470 (58)
Peak CK, Unit/L, no. %			
• >2x reference upper limits	1029	765	1223
	319 (31)	393 (51)	651 (53)
Peak TnT, no. %			
• Positive or >0.01	263	148	245
	240 (91)	141 (95)	236 (96)
Peak TnI, no. %			
• Positive or > reference upper limit/0.04	77	56	147
	52 (68)	46 (82)	112 (76)

	Pre-morbid dyslipidaemia		
	Yes N=1131	No N=902	Not known N=1389
Total cholesterol, mmol/L			
• N	777	675	1019
• Mean, SD	5.2 (1.3)	5.3 (1.3)	5.4 (1.3)
• Median, (min, max)	5.0 (3.0, 10.8)	5.2 (3.0, 14.3)	5.3 (3.0, 10.4)
HDL-C, mmol/L			
• N	784	675	1017
• Mean, SD	1.13 (0.36)	1.16 (0.44)	1.11 (0.34)
• Median, (min, max)	1.10 (0.50, 4.50)	1.10 (0.50, 4.94)	1.10 (0.50, 4.24)
LDL-C, mmol/L			
• N	763	669	996
• Mean, SD	3.13 (1.18)	3.31 (1.26)	3.37 (1.21)
• Median, (min, max)	3.00 (1.00, 8.90)	3.24 (1.00, 10.10)	3.30 (1.02, 8.20)
Triglycerides, mmol/L			
• N	685	581	885
• Mean, SD	2.12 (1.22)	2.06 (1.28)	2.10 (1.37)
• Median, (min, max)	1.80 (1.00, 13.00)	1.70 (1.00, 12.90)	1.70 (1.00, 14.00)
Left ventricular ejection fraction, %			
• N	549	470	697
• Mean, SD	47 (14)	48 (12)	48 (13)
• Median, (min, max)	47 (10, 80)	50 (9, 80)	49 (8, 80)

Note: Percentage is to the nearest decimal point.

Figure 3.2.5a Stratum distribution for patients with ACS by pre-morbid dyslipidaemia, Malaysia 2006

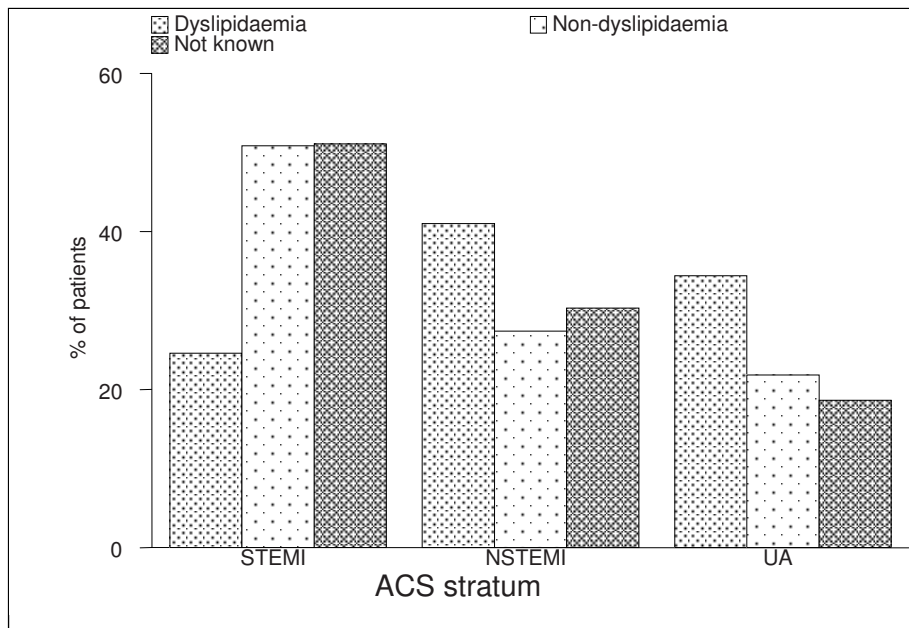


Figure 3.2.5b Number of distinct angina episodes for patients with ACS by pre-morbid dyslipidaemia, Malaysia 2006

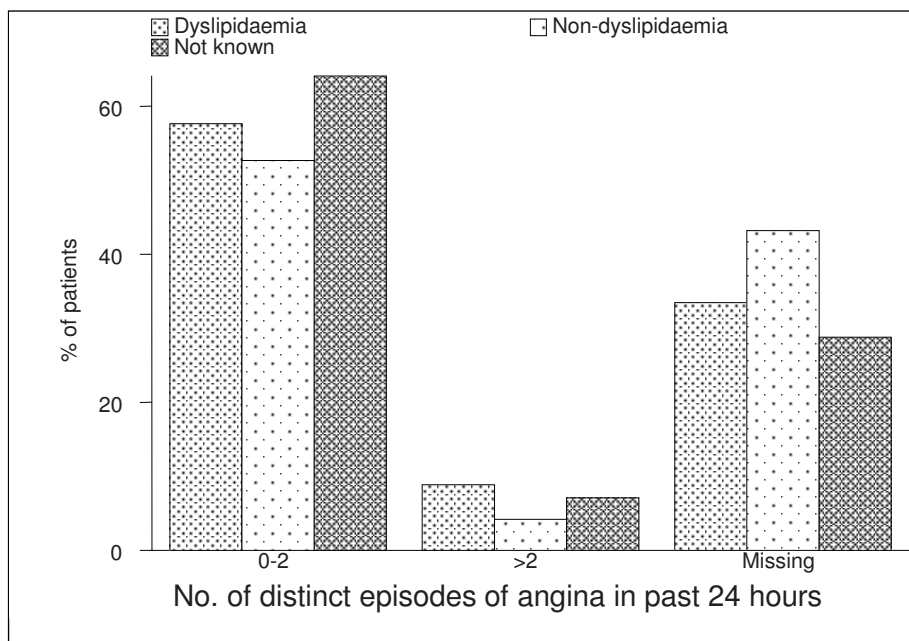
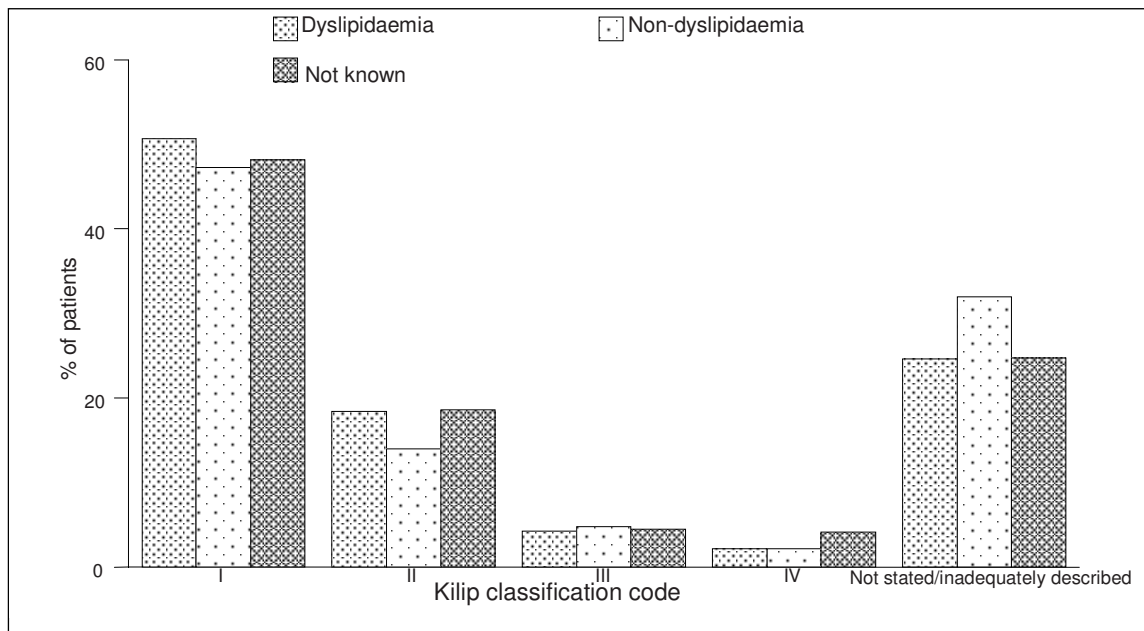


Figure 3.2.5c Killip classification code for patients with ACS by pre-morbid dyslipidaemia, Malaysia 2006



CHAPTER 4

TREATMENT

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This chapter summarizes management pattern of patients admitted with acute coronary syndrome registered in the ACS registry 2006.

Admission days

In general the total admission days in hospital is similar across the ACS strata; that is STEMI 6 days \pm 3, NSTEMI 6 days \pm 4 and UA 5 days \pm 4. However, there is an increasing trend of total admission days with increasing age (STEMI; young is 5 days \pm 2, middle age is 6 days \pm 3, elderly is 7 days \pm 4 and NSTEMI/UA; young is 5 \pm 2 days, middle age is 5 \pm 3 days, elderly is 6 \pm 4 days).

A similar pattern is also seen in the number of days in CCU where the days spent after STEMI increases as the age increases with a maximum of 21 days in the middle age and 26 days in the elderly age groups, as compared to 10 days in the young age group. A similar pattern is also seen in the number of days spent in intensive care unit (ICU)/ coronary intensive care unit (CICU) post STEMI, with a generally higher number of days for the middle age and elderly age group with a maximum of 17 days, as compared to only 4 days maximum in the young age group. The young age group patients also spent fewer days in CCU/ICU/CICU post NSTEMI/UA (Table 4.3.1).

In general, no differences are seen in total admission days or number of days admitted to CCU between the different ethnic groups presented with ACS. However, a longer admission into ICU/CICU was seen among the Malays (mean 4 days \pm 4) following STEMI, as compared to other ethnic groups (Chinese 2 days \pm 2, Indian and other ethnic groups 2 days \pm 1). In NSTEMI/UA, the Malay group spent the longest time in CCU/ICU/CICU (24 days) while the lowest was in ethnic groups other than Malays, Chinese and Indian (10 days).

Treatment of STEMI (Table 4.1)

Out of the total 1,445 patients with STEMI, 70% received fibrinolytic therapy. The highest proportion of patients who received fibrinolytic therapy was seen in the young age group (aged 20 \geq X > 40, 78%) versus the middle age (aged 40 \geq X >60, 75%) and elderly (aged \geq 60, 62%). The proportion of females treated with fibrinolysis was also lower than males (67% v 71% respectively). Amongst the different ethnic groups, the Malays received the most fibrinolysis treatment (78%) with the lowest proportion being the Indians (66%). However, out of the total of 117 patients who proceeded directly to primary angioplasty, the highest proportion was in the Indian population (13%), followed by Chinese (8%), Malays (7%) and other ethnic groups (6%).

Most of the patients presenting with STEMI were treated conservatively during the same admission with only 308 (21%) having percutaneous coronary intervention (PCI). Most

patients were treated and stabilized medically and referred to a tertiary centre later on as outpatients for further management. There was no difference in the number of PCI among the different age groups. Male patients had a higher proportion of PCI (22%) as compared to the females (16%). When comparing the different ethnic groups that presented with STEMI, the Indian population had the highest proportion of PCI (29%) with the Malay population (18%) accounting for the lowest proportion.

Only a small number (n=10) underwent CABG during the same admission following a STEMI, with mostly in the elderly age group (n=7).

Treatment of NSTEMI/UA

Out of the total of 1,977 patients presenting with NSTEMI/UA, the highest number of patients was in the elderly age group (n=1052), followed by middle age (n=872) and young age group (n=53). The majority of patients were treated medically and only 14% of NSTEMI and 9% of UA patients had PCI. There were more number of younger age group patients (19%) compared to middle age group (14%) and elderly age group (11%) who went for PCI. A small number of middle age (n=13, 1%) and elderly patients (n=22, 2%) were transferred to another centre for further intervention. A small number of middle age (n=25) and elderly (n=32) patients had CABG performed during the same admission.

The number of PCI appeared to be lower in females (n=60, 9%) than male (n=182, 14%). There was no significant difference seen in the proportion of patients undergoing PCI among different ethnic groups. The proportion of patients transferred to another centre was also similar among the different ethnic groups.

Pharmacological treatment of ACS

Aspirin and statins were used in more than 90% of patients in all the ACS groups with no difference seen in STEMI patients of different genders, age groups and ethnic groups. However, there was a downward trend in the use of aspirin during admission with NSTEMI/UA as the age increased (young 94%, middle age 92% and elderly 89%). Female patients who had NSTEMI/UA also received less treatment with aspirin (88%) as compared to male patients (91%). ADP antagonist use was slightly lower in the UA group (50%) as compared to the STEMI (60%) or NSTEMI groups (64%).

Due to religious beliefs, the use of LMWH in the Malay population was the lowest in both ACS strata (28% in STEMI, 55% in NSTEMI/UA) as compared to the rest of the ethnic groups (Table 4.2.3 and 4.3.3). The use of GP receptor inhibitor was very low in all ACS strata, the highest being in STEMI (n=77, 5%), NSTEMI (n=47, 4%) and UA (n=19, 2%).

Hypoglycaemic agents were mostly used among the Indian population for all ACS presentations, reaching 41% in STEMI and 42% in NSTEMI/UA. Similarly insulin therapy in the Indian patients was also the highest (37% in all ACS) as compared to the other ethnic groups. This may reflect the prevalence of diabetes in the different ethnic groups.

In all the ACS groups, diuretics were used the most amongst the elderly age group (STEMI 37% and NSTEMI 42%). The lowest usage of diuretics was seen in the young patients presenting with STEMI (2%). This is in proportion with the high proportion of Killips Class 1 in the young age group as described in the earlier chapter.

Summary Points:

- Patients with ACS stayed an average of 6 days in hospital which included approximately 3 days in CCU. There is an increasing trend of longer duration of hospitalization with increasing age.
- For the STEMI patients 70% received thrombolysis and only 8% proceeded directly to primary angioplasty.
- The highest proportion of patients who received thrombolytic therapy was seen in the young age group, male and Malays.
- Twenty-percent of STEMI patients had PCI during the same admission. Males have a higher proportion of PCI compared to females and Indians have the highest proportion of PCI while Malays had the lowest.
- For NSTEMI/UA, majority of the patients were medically treated. Only 14% of NSTEMI and 9% of UA patients had PCI on the same admission.
- Prescription and utilization of adjunctive proven pharmacological therapy were high in all groups.

Table 4.1 Summary of treatments for patients with ACS by ACS stratum, Malaysia 2006

	STEMI N=1445	NSTEMI N=1132	UA N=845
Total admission days*			
• N	1420	1104	830
• Mean, SD	6 (3)	6 (4)	5 (4)
• Median, (min, max)	5 (1,28)	5 (1,30)	4 (1,29)
Number of days on CCU			
• N	1093	450	140
• Mean, SD	3 (3)	4 (3)	3 (3)
• Median, (min, max)	3 (1,26)	3 (1,24)	2 (1,20)
Number of days on ICU/CICU			
• N	87	110	27
• Mean, SD	3 (3)	4 (3)	4 (2)
• Median, (min, max)	2 (1,17)	3 (1,23)	4 (1,9)
Fibrinolytic therapy, no. %			
• Given	1018 (70)	NA	NA
• Not given–proceeded directly to primary angioplasty	117 (8)	NA	NA
• Not given–Contraindicated	70 (5)	NA	NA
• Not given–Missed thrombolysis	193 (13)	NA	NA
• Not given–Others**	47 (3)	NA	NA
Cardiac catheterization, no. %			
• Yes	298 (21)	251 (22)	106 (13)
• No	1106 (77)	858 (76)	727 (86)
• Number transferred to another centre	39 (3)	23 (2)	12 (1)
Percutaneous coronary intervention, no. %			
• Yes	308 (21)	162 (14)	80 (9)
• No	1137 (79)	970 (86)	765 (91)
CABG, no. %			
• Yes	10 (1)	42 (4)	15 (2)
• No	1435 (99)	1090 (96)	830 (98)
Pre-admission aspirin use, no. %			
• Yes	227 (16)	465 (41)	372 (44)
• No	965 (67)	468 (41)	257 (30)
• Unknown	253 (18)	199 (18)	216 (26)

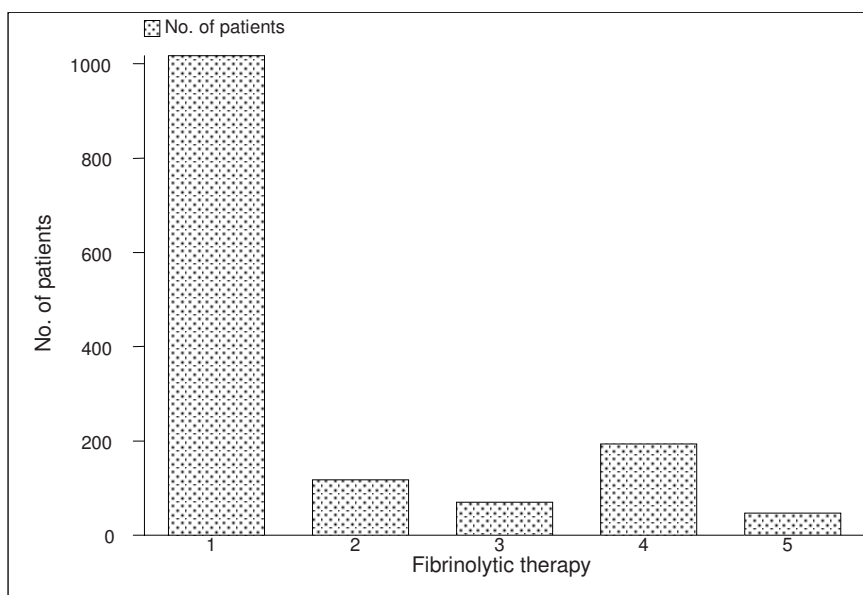
	STEMI N=1445	NSTEMI N=1132	UA N=845
Pharmacological therapy given during admission, no. %			
• Aspirin	1368 (95)	1018 (90)	765 (91)
• ADP antagonist	868 (60)	719 (64)	422 (50)
• GP receptor inhibitor	77 (5)	47 (4)	19 (2)
• Unfractionated heparin	181 (13)	203 (18)	197 (23)
• LMWH	446 (31)	767 (68)	537 (64)
• Beta blocker	951 (66)	737 (65)	587 (69)
• ACE inhibitor	865 (60)	597 (53)	510 (60)
• Angiotensin II receptor blocker	66 (5)	131 (12)	70 (8)
• Statin	1333 (92)	1022 (90)	769 (91)
• Other lipid lowering agent	54 (4)	79 (7)	54 (6)
• Diuretics	393 (27)	464 (41)	241 (29)
• Calcium antagonist	94 (7)	253 (22)	195 (23)
• Oral hypoglycaemic agent	373 (26)	364 (32)	236 (28)
• Insulin	379 (26)	320 (28)	183 (22)
• Anti-arrhythmic agent	135 (9)	72 (6)	49 (6)

**Total admission days is derived as Outcome date – Admission date + 1*

***Not given–Others includes missing and refusal*

Note: Percentage is to the nearest decimal point.

Figure 4.1.1 Fibrinolytic therapy for patients with STEMI by ACS stratum, Malaysia 2006



1. Given, 2. Not given—proceeded directly to primary angioplasty, 3. Not given—Contraindicated, 4. Not given—Missed thrombolysis, 5. Not given—Others**

Figure 4.1.2 Cardiac catheterization for patients with ACS by ACS stratum, Malaysia 2006

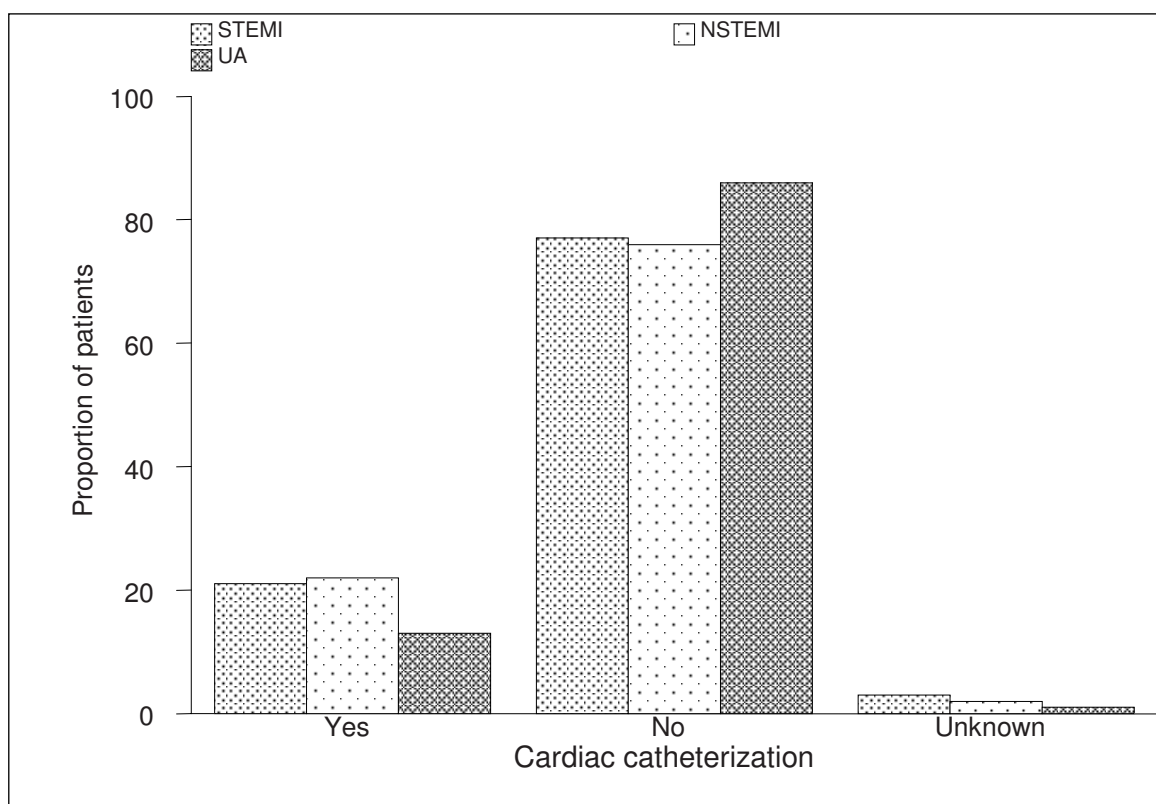


Figure 4.1.3 Percutaneous coronary intervention for patients with ACS by ACS stratum, Malaysia 2006

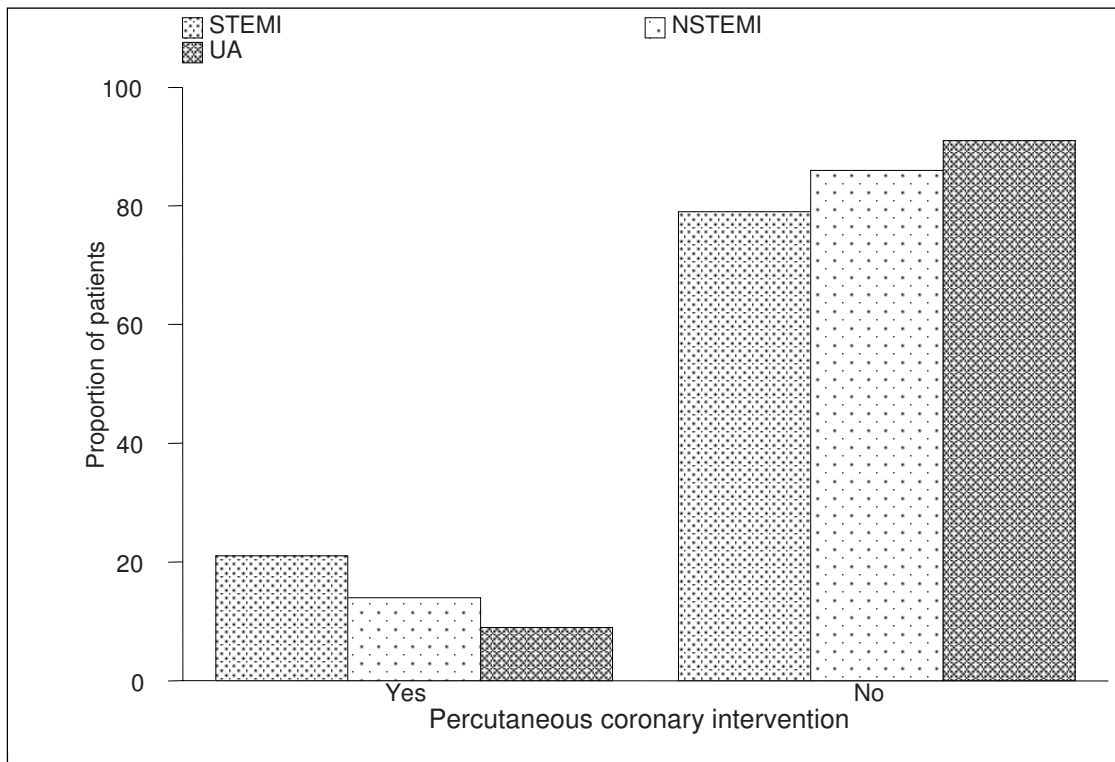


Figure 4.1.4 CABG for patients with ACS by ACS stratum, Malaysia 2006

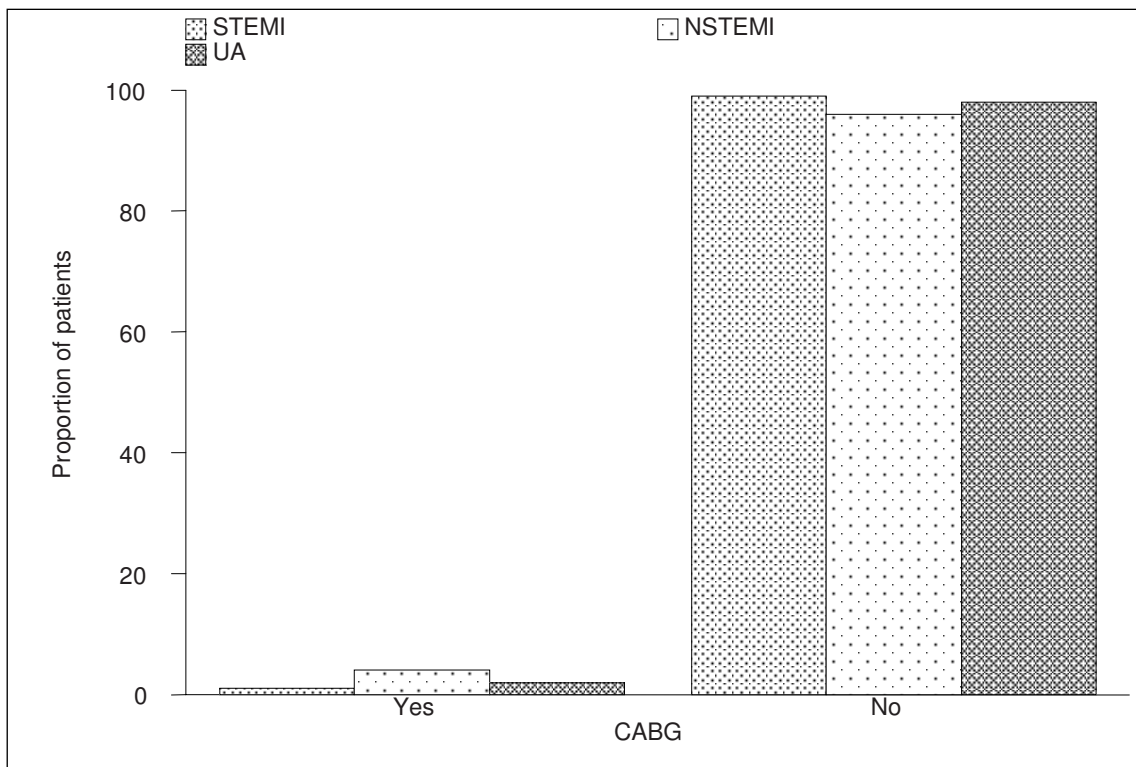


Table 4.2.1 Treatments for patients with STEMI by age group (years), Malaysia 2006

	Age group*		
	Young N=113	Middle-age N=803	Elderly N=529
Total admission days**			
• N	110	796	514
• Mean, SD	5 (2)	6 (3)	7 (4)
• Median, (min, max)	5 (1,20)	5 (1,28)	5 (1,28)
Number of days on CCU			
• N	86	614	393
• Mean, SD	3 (2)	3 (2)	4 (3)
• Median, (min, max)	3 (1,10)	3 (1,21)	3 (1,26)
Number of days on ICU/CICU			
• N	7	48	32
• Mean, SD	2 (1)	3 (3)	4 (4)
• Median, (min, max)	2 (1,4)	2 (1,17)	3 (1,17)
Fibrinolytic therapy, no. %			
• Given	88 (78)	601 (75)	329 (62)
• Not given—proceeded directly to primary angioplasty	7 (6)	59 (7)	51 (10)
• Not given—Contraindicated	3 (3)	31 (4)	36 (7)
• Not given—Missed thrombolysis	10 (9)	94 (12)	89 (17)
• Not given – Others***	5 (4)	18 (2)	24 (5)
Cardiac catheterization, no. %			
• Yes	27 (24)	171 (21)	100 (19)
• No	81 (72)	612 (76)	413 (78)
• No-Transferred to another centre	5 (4)	19 (2)	15 (3)
Percutaneous coronary intervention, no. %			
• Yes	25 (22)	180 (22)	103 (19)
• No	88 (78)	623 (78)	426 (81)
CABG, no. %			
• Yes	0 (0)	3 (0)	7 (1)
• No	113 (100)	800 (100)	522 (99)
Pre-admission aspirin use, no. %			
• Yes	10 (9)	111 (14)	106 (20)
• No	89 (79)	536 (67)	340 (64)
• Unknown	14 (12)	156 (19)	83 (16)

	Age group*		
	Young N=113	Middle-age N=803	Elderly N=529
Pharmacological therapy given during admission, no. %			
• ASA	110 (97)	760 (95)	498 (94)
• ADP antagonist	69 (61)	452 (56)	347 (66)
• GP receptor inhibitor	7 (6)	42 (5)	28 (5)
• Unfractionated heparin	10 (9)	105 (13)	66 (12)
• LMWH	44 (39)	232 (29)	170 (32)
• Beta blocker	86 (76)	555 (69)	310 (59)
• ACE inhibitor	68 (60)	506 (63)	291 (55)
• Angiotensin II receptor blocker	6 (5)	39 (5)	21 (4)
• Statin	105 (93)	750 (93)	478 (90)
• Other lipid lowering agent	10 (9)	29 (4)	15 (3)
• Diuretics	13 (12)	184 (23)	196 (37)
• Calcium antagonist	3 (3)	46 (6)	45 (9)
• Oral hypoglycaemic agent	22 (19)	229 (29)	122 (23)
• Insulin	22 (19)	221 (28)	136 (26)
• Anti-arrhythmic agent	7 (6)	71 (9)	57 (11)

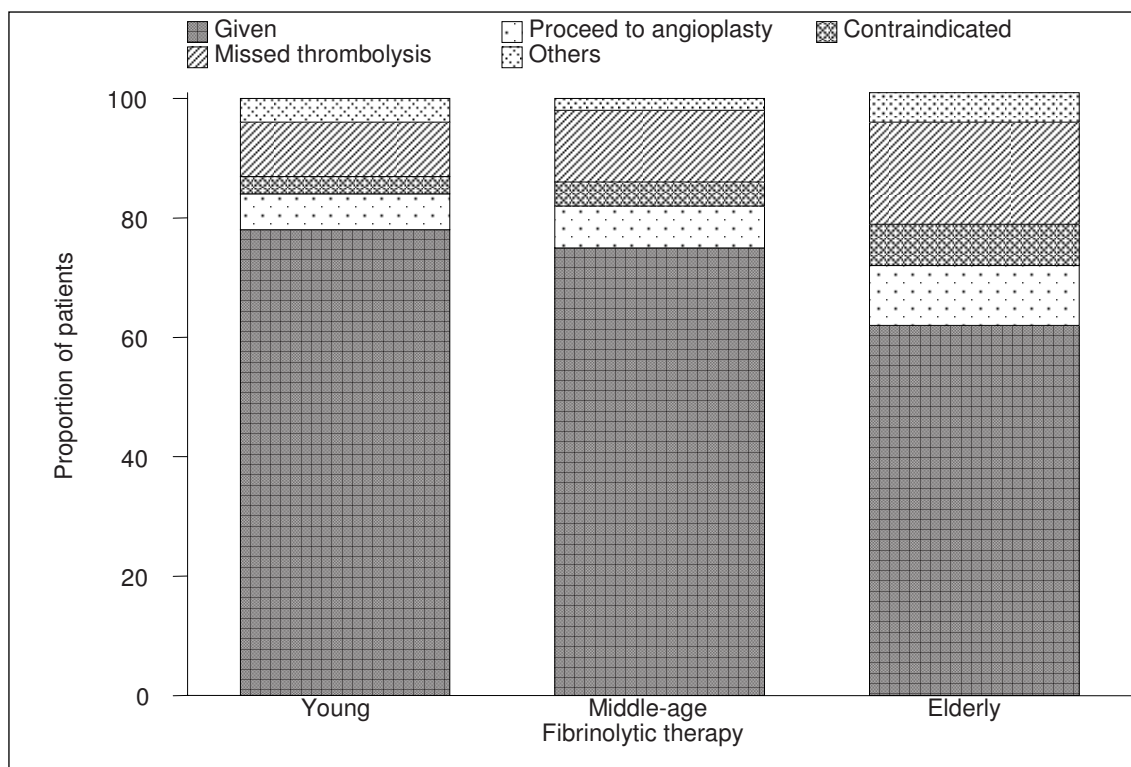
*Young is defined as age from 20 to less than 40 years, middle-age is defined as age between 40 to less than 60 years and elderly is defined as 60 years and above.

**Total admission days is derived as Outcome date – Admission date + 1

***Not given—Others includes missing and refusal

Note: Percentage is to the nearest decimal point.

Figure 4.2.1a Fibrinolytic therapy for patients with STEMI by age group, Malaysia 2006



1. Given, 2. Not given—proceeded directly to primary angioplasty, 3. Not given—Contraindicated, 4. Not given—Missed thrombolysis, 5. Not given—Others**

** Others includes patients who refused the fibrinolytic therapy and missing

Note: Percentage is to the nearest decimal point.

Figure 4.2.1b Cardiac catheterization for patients with STEMI by age group, Malaysia 2006

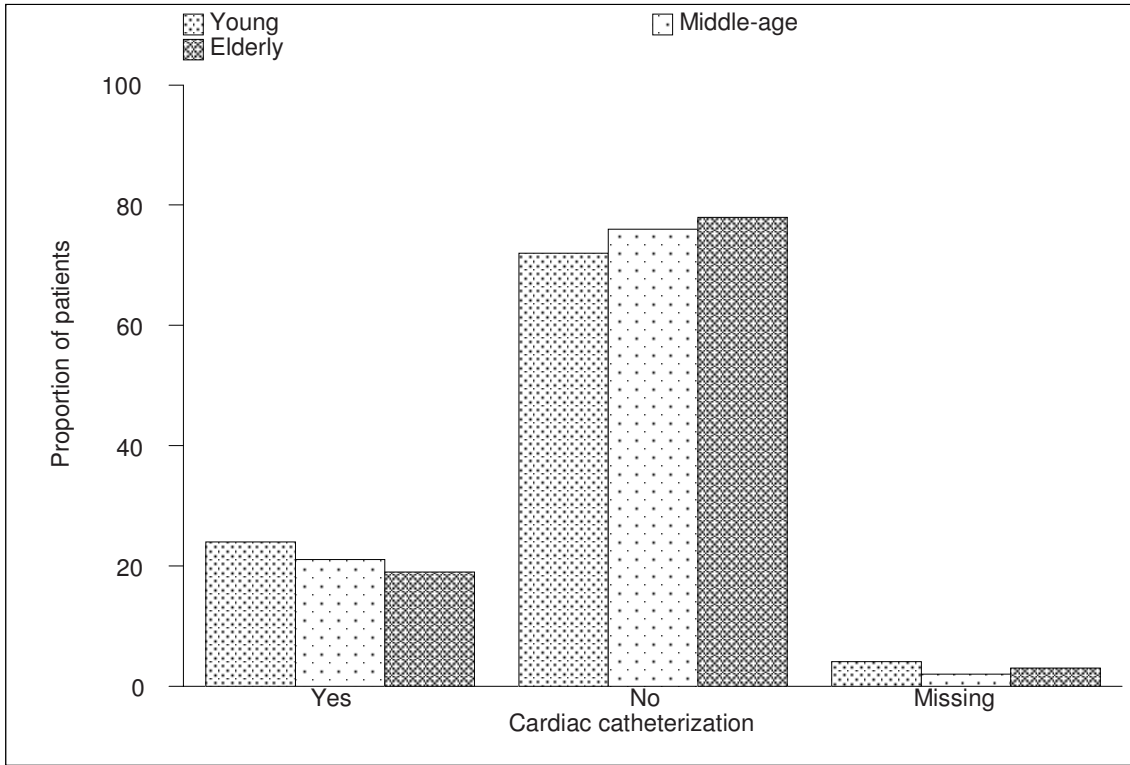


Figure 4.2.1c Percutaneous coronary intervention for patients with STEMI by age group, Malaysia 2006

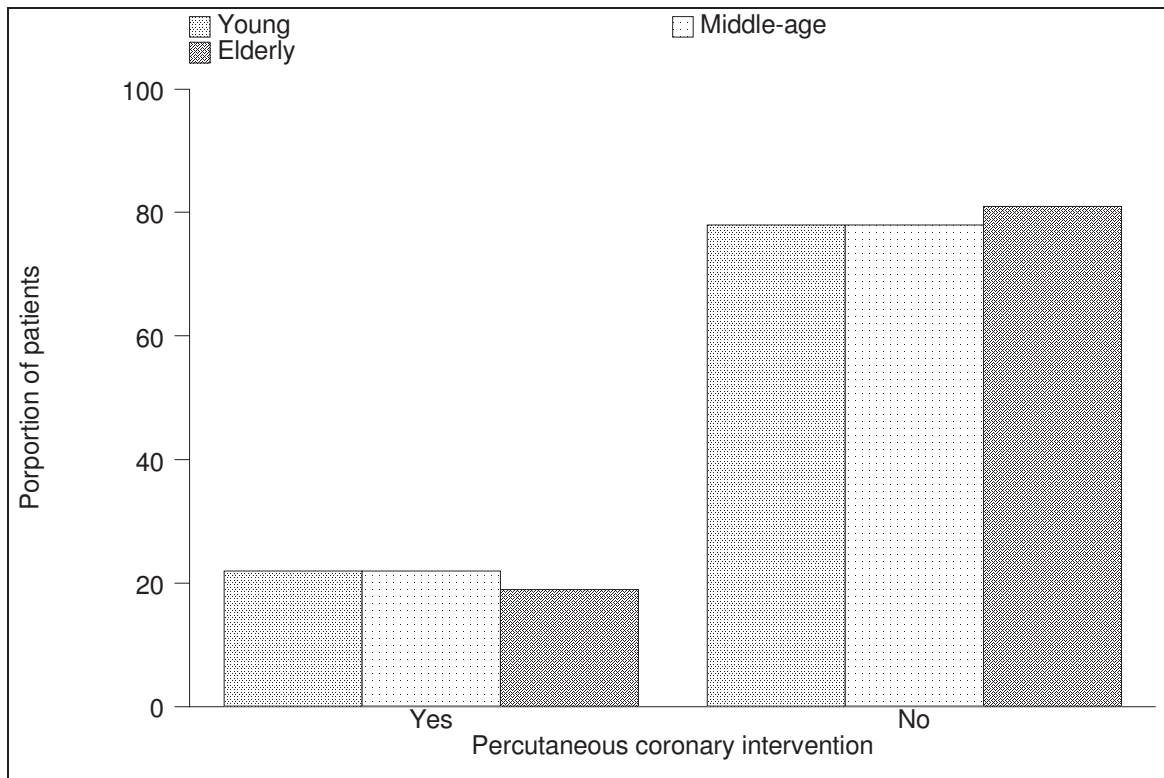


Figure 4.2.1d CABG for patients with STEMI by age group, Malaysia 2006

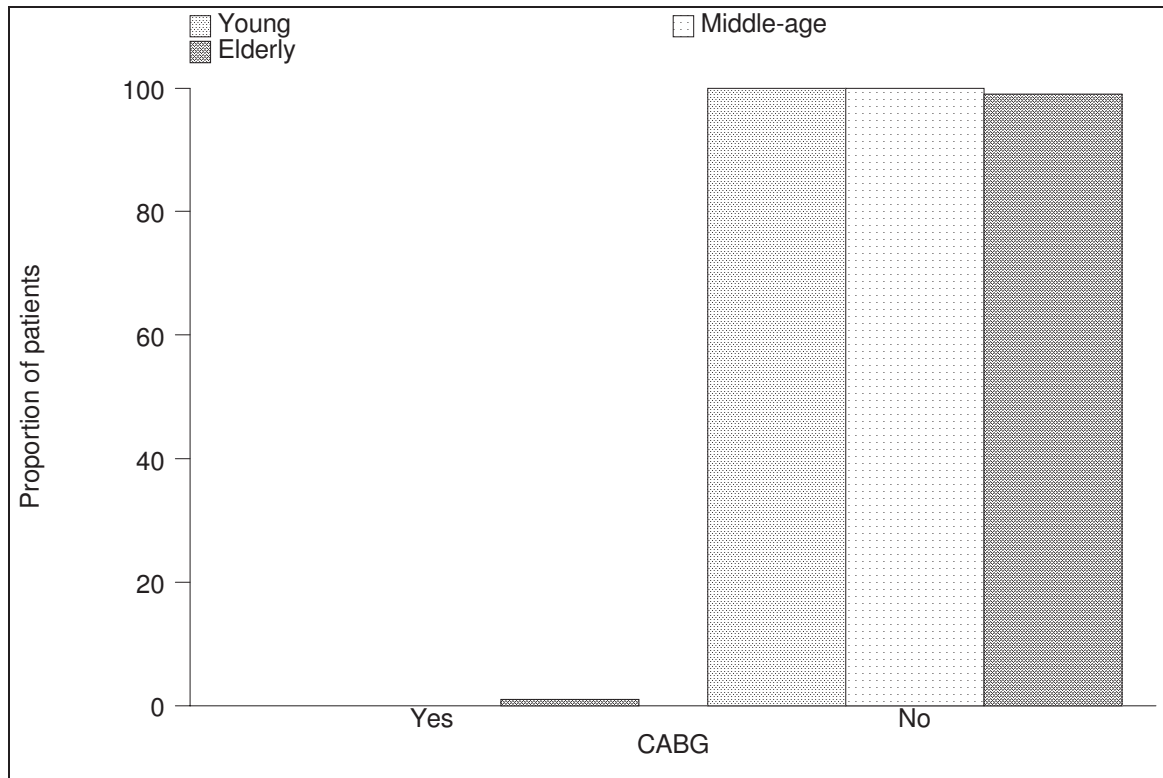


Table 4.2.2 Treatments for patients with STEMI by gender, Malaysia 2006

	Male N=1230	Female N=215
Total admission days*		
• N	1211	209
• Mean, SD	6 (3)	7 (4)
• Median, (min, max)	5 (1,28)	5 (1,28)
Number of days on CCU		
• N	936	157
• Mean, SD	3 (3)	4 (3)
• Median, (min, max)	3 (1,26)	3 (1,21)
Number of days on ICU/CICU		
• N	68	19
• Mean, SD	3 (3)	3 (2)
• Median, (min, max)	2 (1,17)	2 (1,7)
Fibrinolytic therapy, no. %		
• Given	875 (71)	143 (67)
• Not given—proceeded directly to primary angioplasty	99 (8)	18 (8)
• Not given-Contraindicated	59 (5)	11 (5)
• Not given-Missed thrombolysis	156 (13)	37 (17)
• Not given—Others**	41 (3)	6 (3)
Cardiac catheterization, no. %		
• Yes	266 (22)	32 (15)
• No	929 (76)	177 (82)
• No-Transferred to another centre	33 (3)	6 (3)
Percutaneous coronary intervention, no. %		
• Yes	273 (22)	35 (16)
• No	957 (78)	180 (84)
CABG, no. %		
• Yes	9 (1)	1 (0)
• No	1221 (99)	214 (100)
Pre-admission aspirin use, no. %		
• Yes	189 (15)	38 (18)
• No	824 (67)	141 (66)
• Unknown	217 (18)	36 (17)

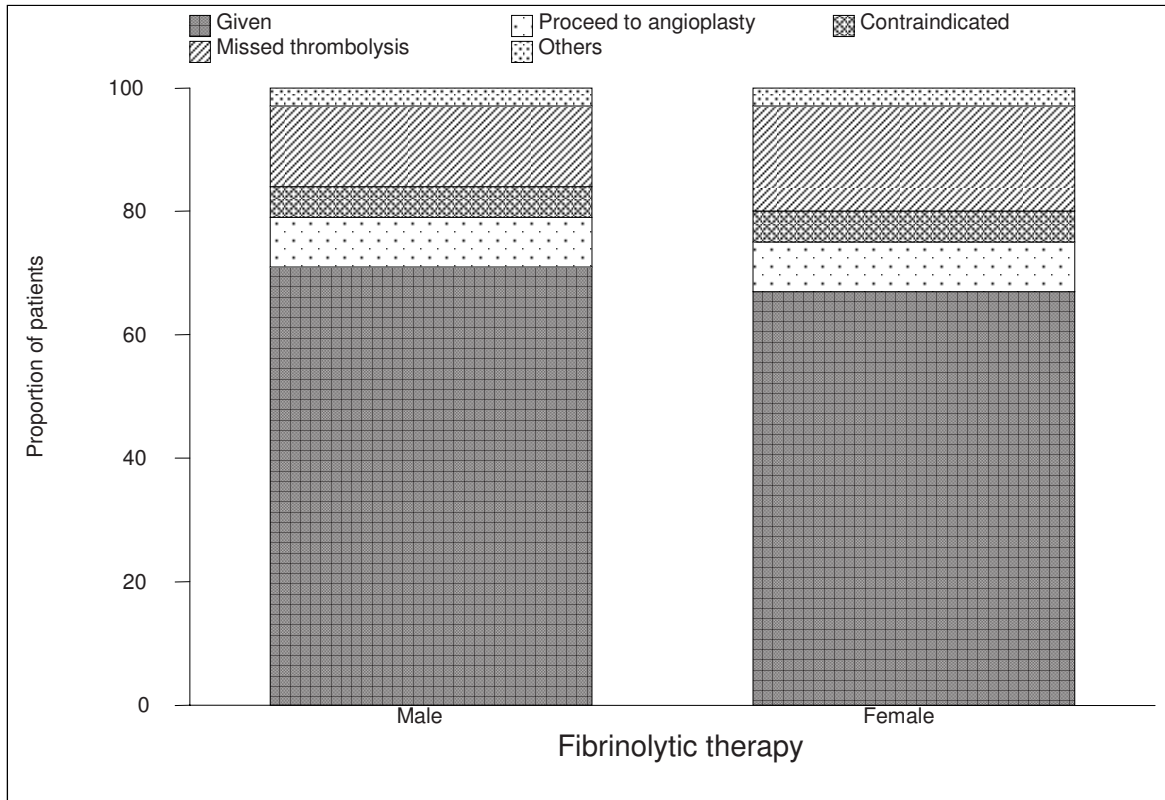
	Male N=1230	Female N=215
Pharmacological therapy given during admission, no. %		
• ASA	1164 (95)	204 (95)
• ADP antagonist	726 (59)	142 (66)
• GP receptor inhibitor	65 (5)	12 (6)
• Unfractionated heparin	156 (13)	25 (12)
• LMWH	371 (30)	75 (35)
• Beta blocker	812 (66)	139 (65)
• ACE inhibitor	759 (62)	106 (49)
• Angiotensin II receptor blocker	56 (5)	10 (5)
• Statin	1136 (92)	197 (92)
• Other lipid lowering agent	45 (4)	9 (4)
• Diuretics	316 (26)	77 (36)
• Calcium antagonist	79 (6)	15 (7)
• Oral hypoglycaemic agent	307 (25)	66 (31)
• Insulin	290 (24)	89 (41)
• Anti-arrhythmic agent	112 (9)	23 (11)

*Total admission days is derived as Outcome date – Admission date + 1

**Not given–Others includes missing and refusal

Note: Percentage is to the nearest decimal point.

Figure 4.2.2a Fibrinolytic therapy for patients with STEMI by gender, Malaysia 2006



1. Given, 2. Not given–proceeded directly to primary angioplasty, 3. Not given–Contraindicated, 4. Not given–Missed thrombolysis, 5. Not given–Others**

** Others includes patients who refused the fibrinolytic therapy and missing
Note: Percentage is to the nearest decimal point.

Figure 4.2.2b Cardiac catheterization for patients with STEMI by gender, Malaysia 2006

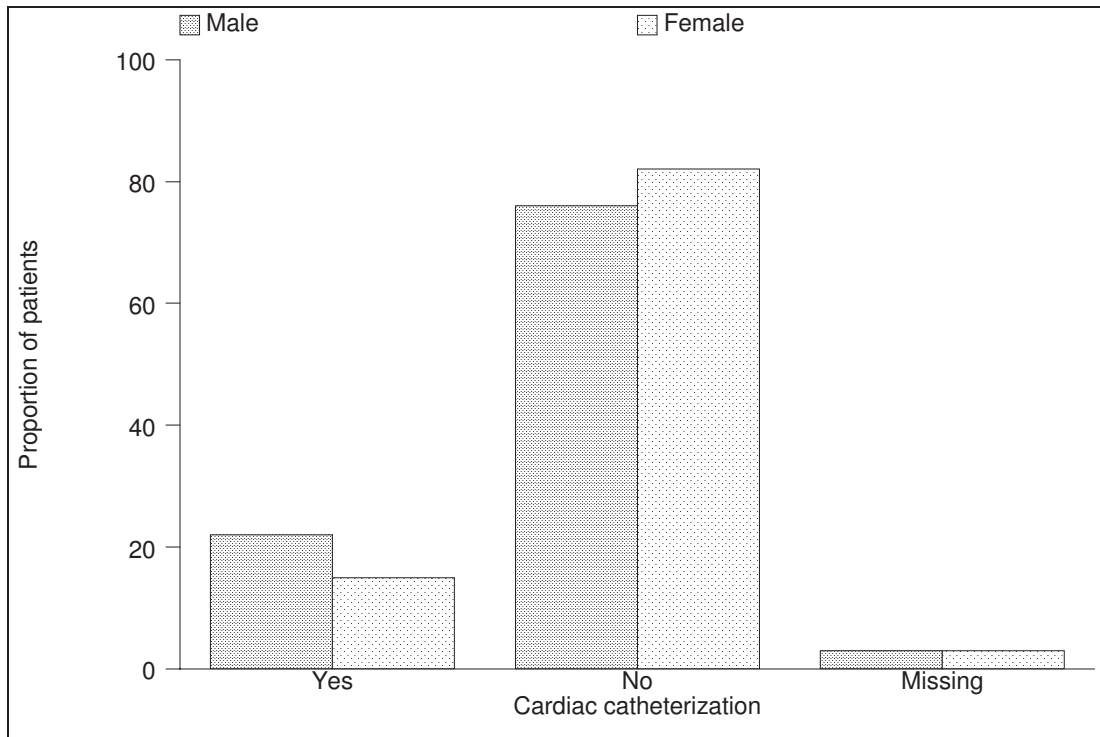


Figure 4.2.2c Percutaneous coronary intervention for patients with STEMI by gender, Malaysia 2006

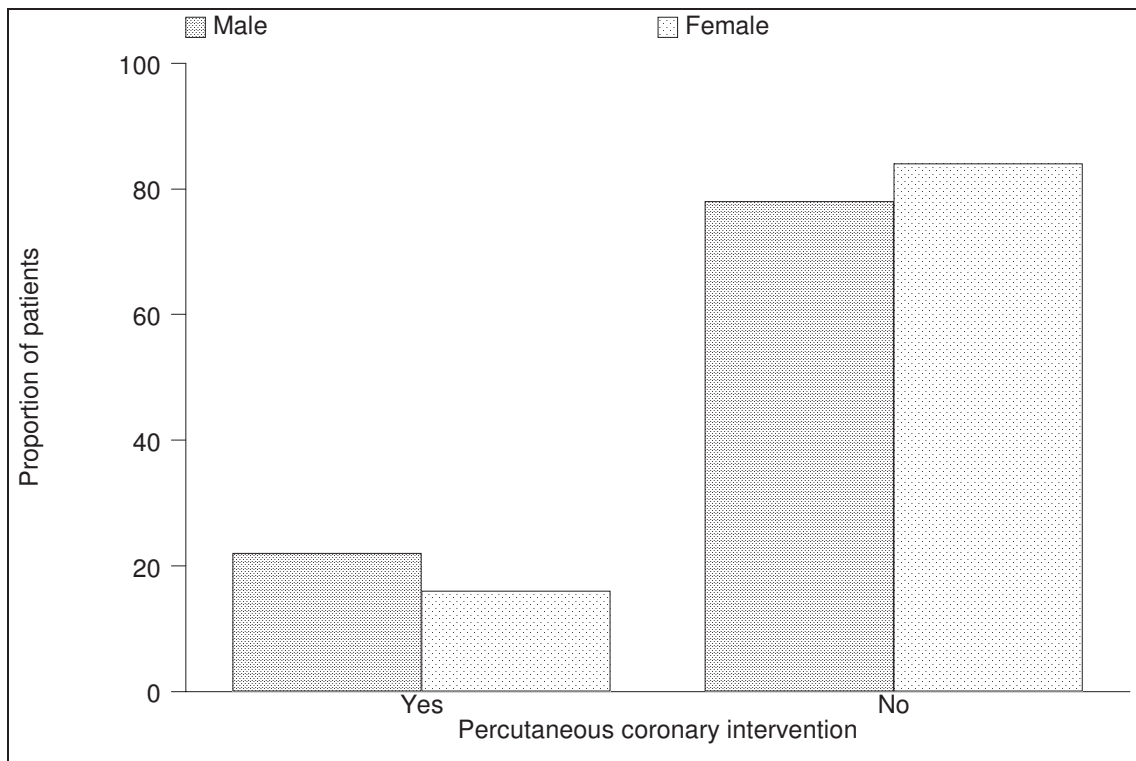


Figure 4.2.2d CABG for patients with STEMI by gender, Malaysia 2006

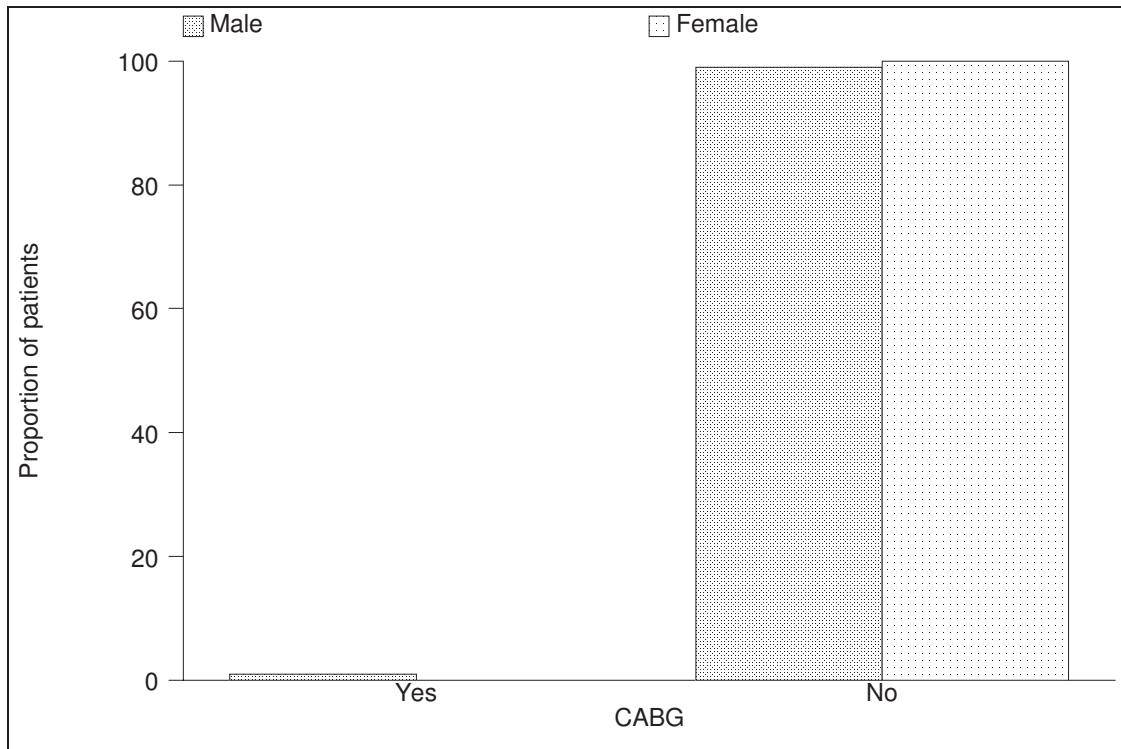


Table 4.2.3 Treatments for patients with STEMI by ethnic group, Malaysia 2006

	Malay N=780	Chinese N=301	Indian N=286	Others* N=78
Total admission days**				
• N	764	294	284	78
• Mean, SD	6 (3)	6 (3)	6 (4)	5 (3)
• Median, (min, max)	5 (1,28)	5 (1,28)	5 (1,27)	5 (1,25)
Number of days on CCU				
• N	614	225	187	67
• Mean, SD	4 (2)	3 (2)	4 (3)	3 (2)
• Median, (min, max)	3 (1,21)	3 (1,15)	3 (1,26)	3 (1,10)
Number of days on ICU/CICU				
• N	43	23	3	18
• Mean, SD	4 (4)	2 (2)	2 (1)	2 (1)
• Median, (min, max)	2 (1,17)	2 (1,8)	2 (1,2)	2 (1,4)
Fibrinolytic therapy, no. %				
• Given	567 (73)	206 (68)	188 (66)	57 (73)
• Not given—proceeded directly to primary angioplasty	52 (7)	24 (8)	36 (13)	5 (6)
• Not given- Contraindicated	39 (5)	14 (5)	11 (4)	6 (8)
• Not given-Missed thrombolysis	99 (13)	43 (14)	42 (15)	9 (12)
• Not given— Others***	23 (3)	14 (5)	9 (3)	1 (1)
Cardiac catheterization, no. %				
• Yes	141 (18)	64 (21)	75 (26)	18 (23)
• No	617 (79)	231 (77)	203 (71)	55 (71)
• No-Transferred to another centre	20 (3)	6 (2)	8 (3)	5 (6)
Percutaneous coronary intervention, no. %				
• Yes	143 (18)	66 (22)	83 (29)	16 (21)
• No	637 (82)	235 (78)	203 (71)	62 (79)
CABG, no. %				
• Yes	8 (1)	2 (1)	0 (0)	0 (0)
• No	772 (99)	299 (99)	286 (100)	78 (100)
Pre-admission aspirin use, no. %				
• Yes	112 (14)	44 (15)	66 (23)	5 (6)
• No	554 (71)	197 (65)	164 (57)	50 (64)
• Unknown	114 (15)	60 (20)	56 (20)	23 (29)

	Malay N=780	Chinese N=301	Indian N=286	Others* N=78
Pharmacological therapy given during admission, no. %				
• ASA	731 (94)	293 (97)	271 (95)	73 (94)
• ADP antagonist	441 (57)	194 (64)	183 (64)	50 (64)
• GP receptor inhibitor	44 (6)	10 (3)	19 (7)	4 (5)
• Unfrac heparin	120 (15)	21 (7)	38 (13)	2 (3)
• LMWH	218 (28)	96 (32)	104 (36)	28 (36)
• Beta blocker	504 (65)	209 (69)	193 (67)	45 (58)
• ACE inhibitor	478 (61)	165 (55)	184 (64)	38 (49)
• Angiotensin II receptor blocker	30 (4)	14 (5)	18 (6)	4 (5)
• Statin	714 (92)	281 (93)	265 (93)	73 (94)
• Other lipid lowering agent	31 (4)	10 (3)	11 (4)	2 (3)
• Diuretics	234 (30)	64 (21)	80 (28)	15 (19)
• Calcium antagonist	55 (7)	10 (3)	21 (7)	8 (10)
• Oral hypoglycaemic agent	178 (23)	69 (23)	116 (41)	10 (13)
• Insulin	184 (24)	73 (24)	107 (37)	15 (19)
• Anti-arrhythmic agent	65 (8)	37 (12)	25 (9)	8 (10)

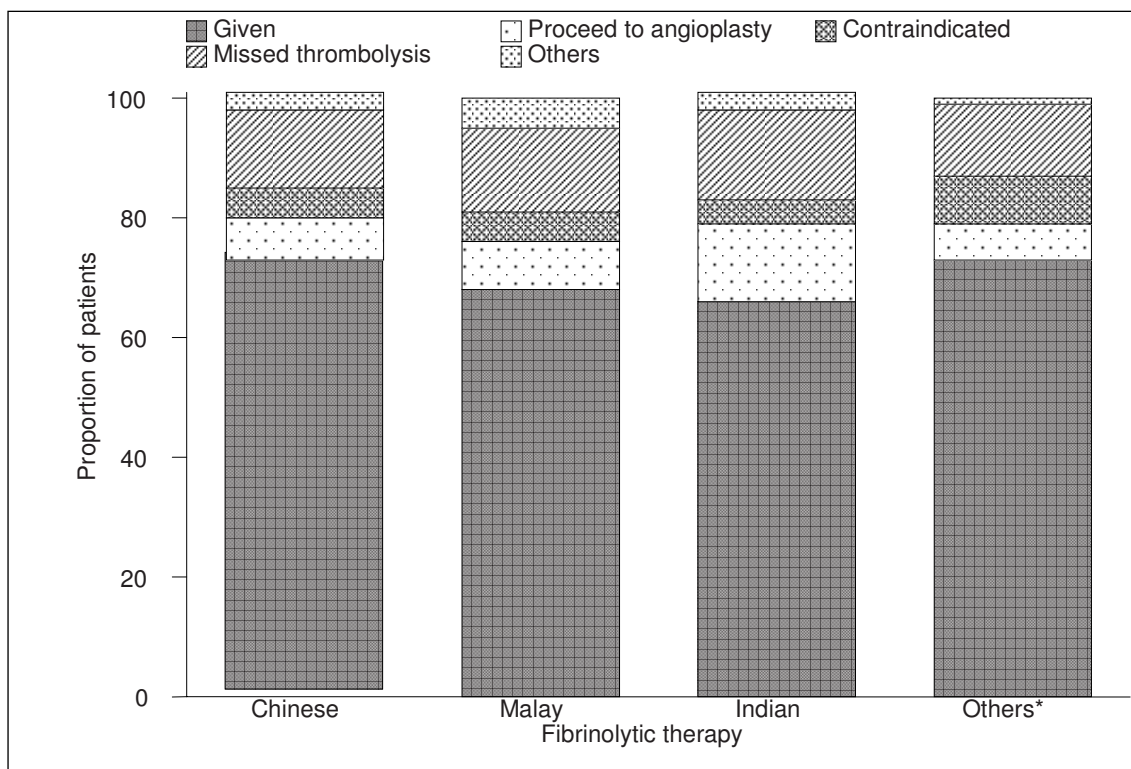
*Others includes Orang asli, Kadazan, Melanau, Murut, Bajau, Bidayuh, Iban, other Malaysian and foreigner

**Total admission days is derived as Outcome date – Admission date + 1

***Not given–Others includes missing and refusal

Note: Percentage is to the nearest decimal point.

Figure 4.2.3a Fibrinolytic therapy for patients with STEMI by ethnic group, Malaysia 2006



1. Given, 2. Not given—proceeded directly to primary angioplasty, 3. Not given—Contraindicated, 4. Not given—Missed thrombolysis, 5. Not given—Others**

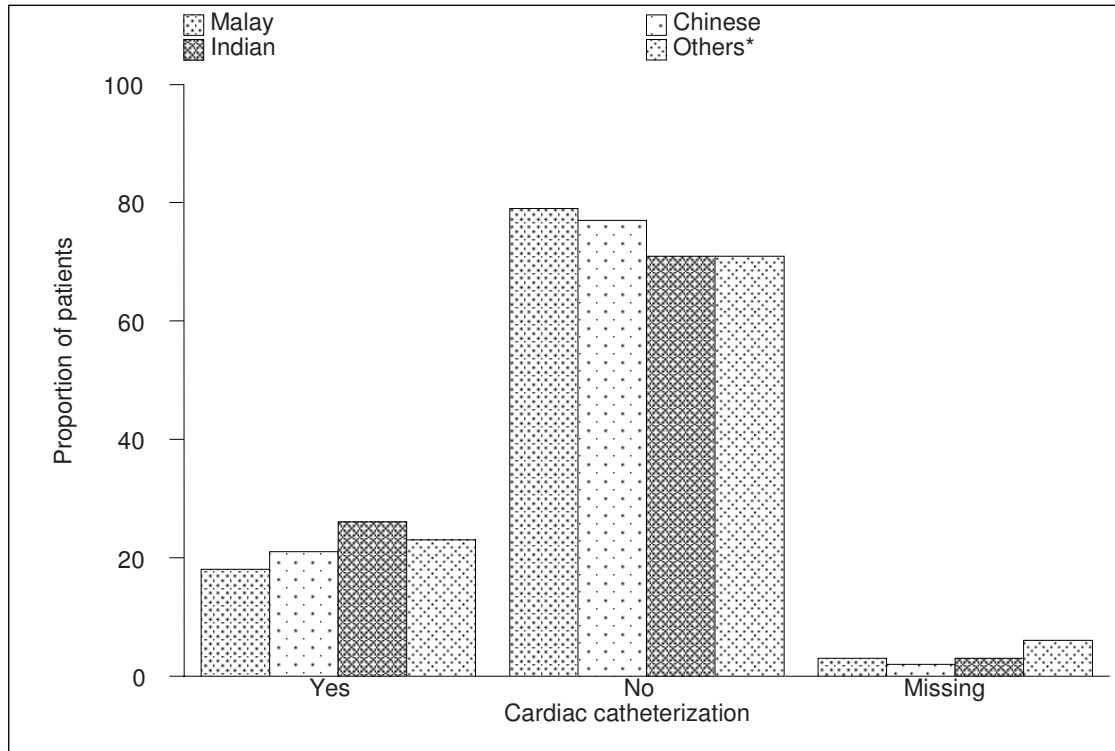
**Others include Orang asli, Kadazan, Melanau, Murut, Bajau, Bidayuh, Iban, other Malaysian and foreigner

Note:

1. Percentage is to the nearest decimal point.

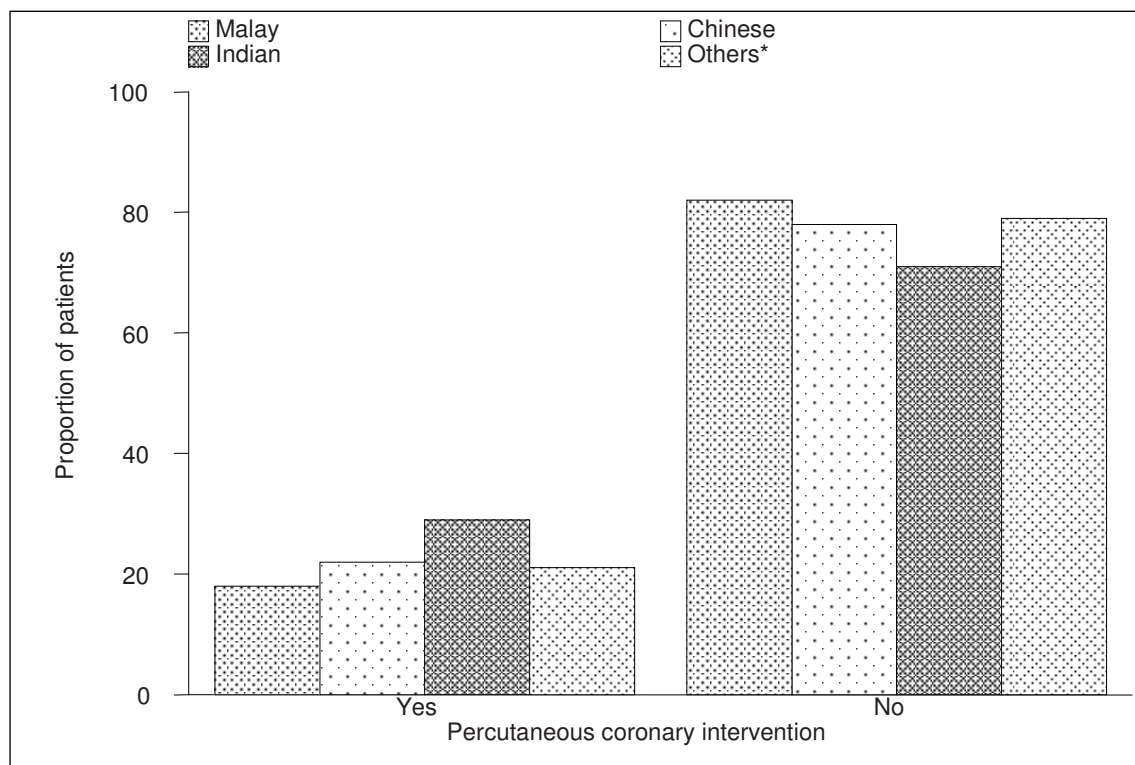
2. Others includes patients who refused the fibrinolytic therapy and missing

Figure 4.2.3b Cardiac catheterization for patients with STEMI by ethnic group, Malaysia 2006



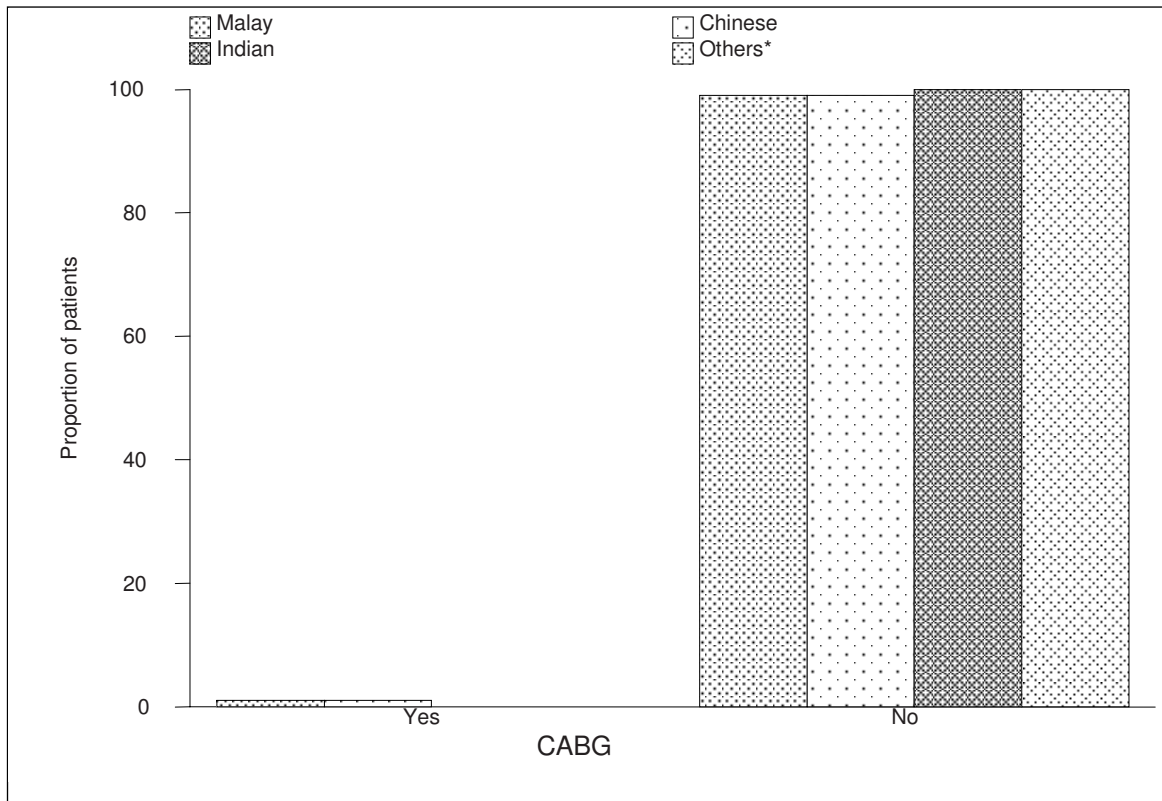
* Others includes Orang asli, Kadazan, Melanau, Murut, Bajau, Bidayuh, Iban, other Malaysian and foreigner

Figure 4.2.3c Percutaneous coronary intervention for patients with STEMI by ethnic group, Malaysia 2006



*Others includes Orang asli, Kadazan, Melanau, Murut, Bajau, Bidayuh, Iban, other Malaysian and foreigner

Figure 4.2.3d CABG on admission for patients with STEMI by ethnic group, Malaysia 2006



**Others includes Orang asli, Kadazan, Melanau, Murut, Bajau, Bidayuh, Iban, other Malaysian and foreigner

Table 4.3.1 Treatments for patients with NSTEMI/UA by age group (years), Malaysia 2006

	Age group*		
	Young N=53	Middle-age N=872	Elderly N=1052
Total admission days**			
• N	53	856	1025
• Mean, SD	5 (2)	5 (3)	6 (4)
• Median, (min, max)	4 (2,15)	4 (1,27)	5 (1,30)
Number of days on CCU			
• N	14	246	330
• Mean, SD	2 (1)	3 (3)	3 (3)
• Median, (min, max)	2 (1,5)	3 (1,20)	3 (1,24)
Number of days on ICU/CICU			
• N	1	62	74
• Mean, SD	1 (.)	4 (3)	4 (3)
• Median, (min, max)	1 (1,1)	3 (1,12)	3 (1,23)
Cardiac catheterization, no. %			
• Yes	11 (21)	164 (19)	182 (17)
• No	42 (79)	695 (80)	848 (81)
• No-Transferred to another centre	0 (0)	13 (1)	22 (2)
Percutaneous coronary intervention, no. %			
• Yes	10 (19)	120 (14)	112 (11)
• No	43 (81)	752 (86)	940 (89)
CABG, no. %			
• Yes	0 (0)	25 (3)	32 (3)
• No	53 (100)	847 (97)	1020 (97)
Pre-admission aspirin use			
• Yes	13 (25)	341 (39)	483 (46)
• No	25 (47)	343 (39)	357 (34)
• Unknown	15 (28)	188 (22)	212 (20)

	Age group*		
	Young N=53	Middle-age N=872	Elderly N=1052
Pharmacological therapy given during admission, no. %			
• ASA	50 (94)	801 (92)	932 (89)
• ADP antagonist	28 (53)	493 (57)	620 (59)
• GP receptor inhibitor	0 (0)	27 (3)	39 (4)
• Unfractionated heparin	14 (26)	200 (23)	186 (18)
• LMWH	28 (53)	566 (65)	710 (67)
• Beta blocker	34 (64)	613 (70)	677 (64)
• ACE inhibitor	26 (49)	503 (58)	578 (55)
• Angiotensin II receptor blocker	4 (8)	75 (9)	122 (12)
• Statin	48 (91)	804 (92)	939 (89)
• Other lipid lowering agent	4 (8)	64 (7)	65 (6)
• Diuretics	14 (26)	249 (29)	442 (42)
• Calcium antagonist	9 (17)	156 (18)	283 (27)
• Oral hypoglycaemic agent	8 (15)	262 (30)	330 (31)
• Insulin	12 (23)	218 (25)	273 (26)
• Anti-arrhythmic agent	3 (6)	45 (5)	73 (7)

*Young is defined as age from 20 to less than 40 years, middle-age is defined as age between 40 to less than 60 years and elderly is defined as 60 years and above.

**Total admission days is derived as Outcome date – Admission date + 1

Note: Percentage is to the nearest decimal point.

Figure 4.3.1a Cardiac catheterization for patients with NSTEMI/UA by age group (years), Malaysia 2006

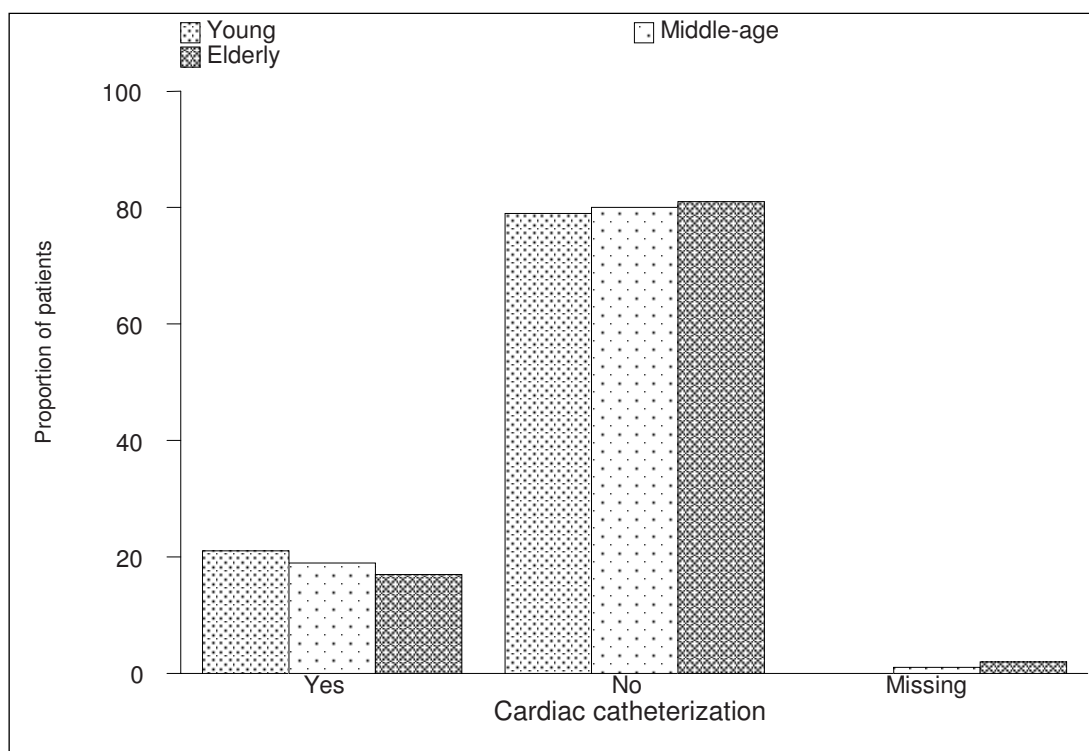


Figure 4.3.1b Percutaneous coronary intervention for patients with NSTEMI/UA by age group (years), Malaysia 2006

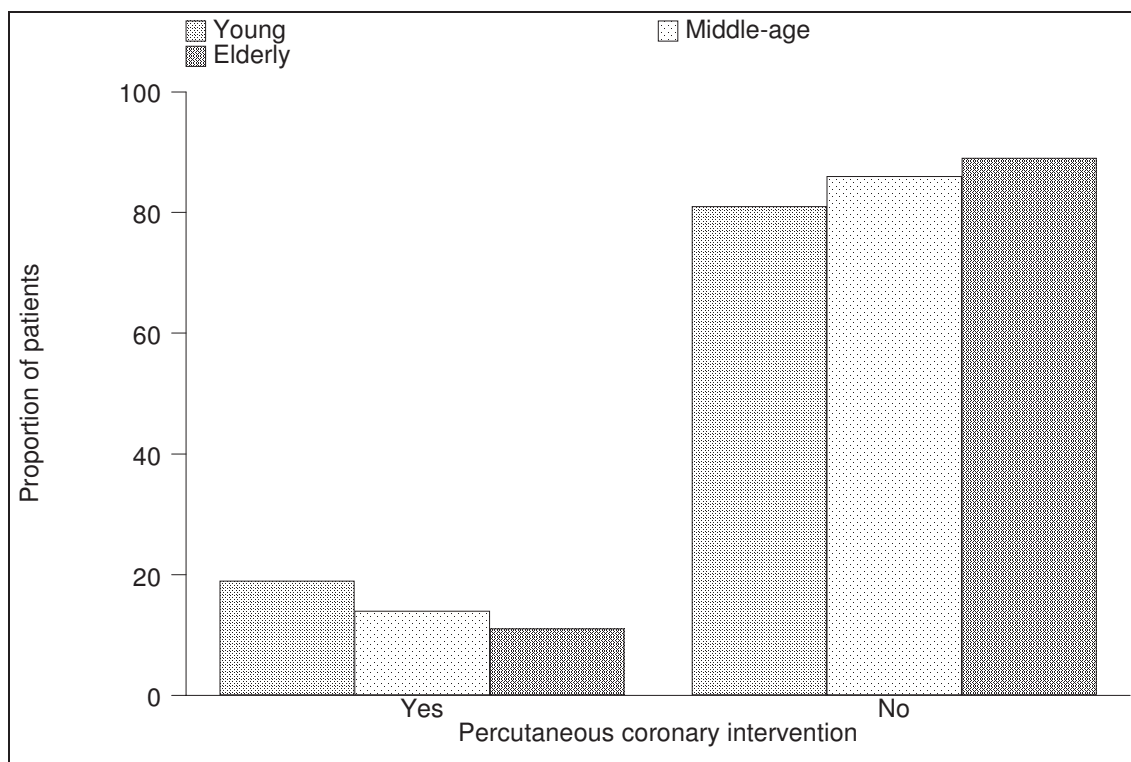


Figure 4.3.1c CABG for patients with NSTEMI/UA by age group (years), Malaysia 2006

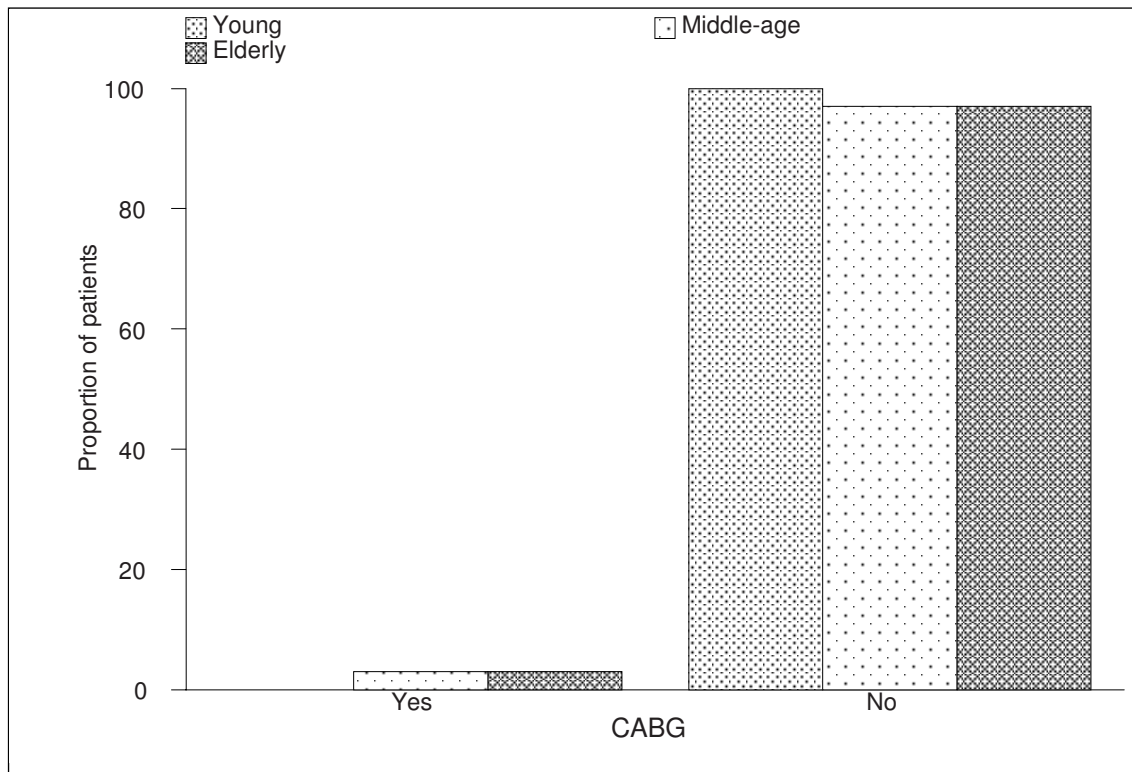


Table 4.3.2 Treatments for patients with NSTEMI/UA by gender, Malaysia 2006

	Male N=1339	Female N=638
Total admission day*		
• N	1315	619
• Mean, SD	6 (4)	6 (4)
• Median, (min, max)	4 (1,29)	5 (1,30)
Number of days on CCU		
• N	425	165
• Mean, SD	3 (3)	4 (3)
• Median, (min, max)	3 (1,24)	3 (1,19)
Number of days on ICU/CICU		
• N	96	41
• Mean, SD	3 (2)	5 (4)
• Median, (min, max)	3 (1,11)	4 (1,23)
Cardiac catheterization, no. %		
• Yes	264 (20)	93 (15)
• No	1056 (79)	529 (83)
• No-Transferred to another centre	19 (1)	16 (3)
Percutaneous coronary intervention, no. %		
• Yes	182 (14)	60 (9)
• No	1157 (86)	578 (91)
CABG, no. %		
• Yes	42 (3)	15 (2)
• No	1297 (97)	623 (98)
Pre-admission aspirin use		
• Yes	570 (43)	267 (42)
• No	471 (35)	254 (40)
• Unknown	298 (22)	117 (18)

	Male N=1339	Female N=638
Pharmacological therapy given during admission, no. %		
• ASA	1224 (91)	559 (88)
• ADP antagonist	813 (61)	328 (51)
• GP receptor inhibitor	50 (4)	16 (3)
• Unfractionated heparin	283 (21)	117 (18)
• LMWH	878 (66)	426 (67)
• Beta blocker	902 (67)	422 (66)
• ACE inhibitor	772 (58)	335 (53)
• Angiotensin II receptor blocker	119 (9)	82 (13)
• Statin	1225 (91)	566 (89)
• Other lipid lowering agent	82 (6)	51 (8)
• Diuretics	453 (34)	252 (39)
• Calcium antagonist	263 (20)	185 (29)
• Oral hypoglycaemic agent	381 (28)	219 (34)
• Insulin	319 (24)	184 (29)
• Anti-arrhythmic agent	85 (6)	36 (6)

*Total admission days is derived as Outcome date – Admission date + 1

Note: Percentage is to the nearest decimal point.

Figure 4.3.2a Cardiac catheterization for patients with NSTEMI/UA by gender, Malaysia 2006

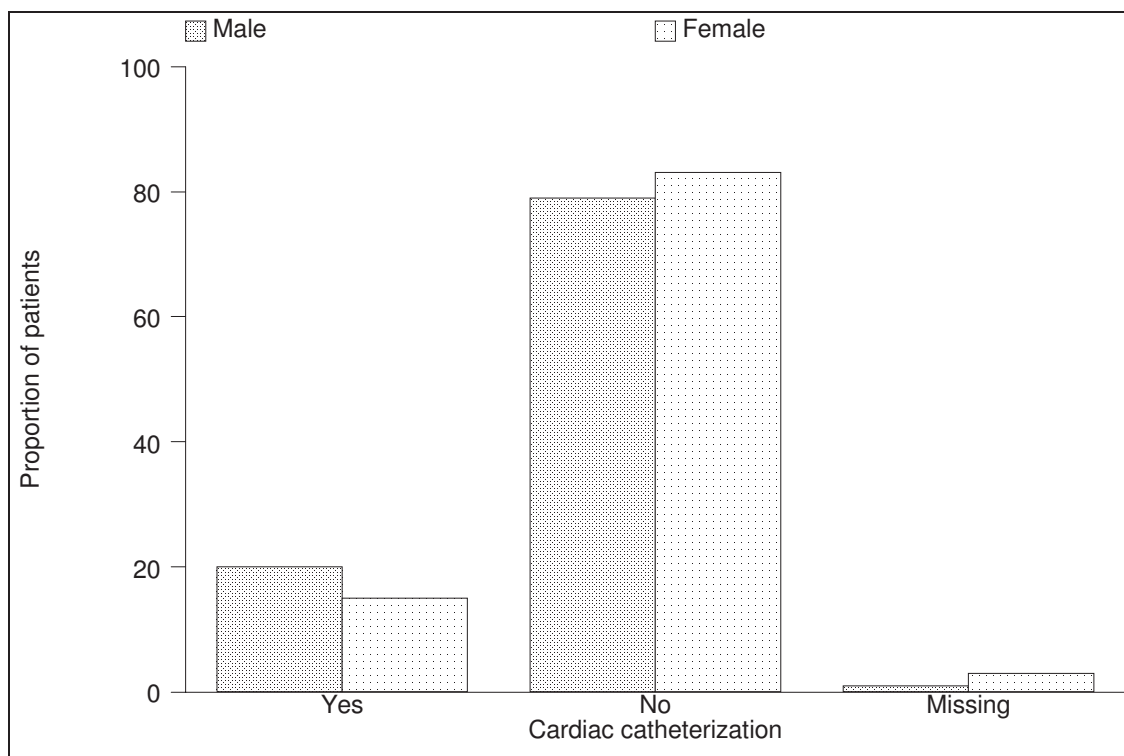


Figure 4.3.2b Percutaneous coronary intervention for patients with NSTEMI/UA by gender, Malaysia 2006

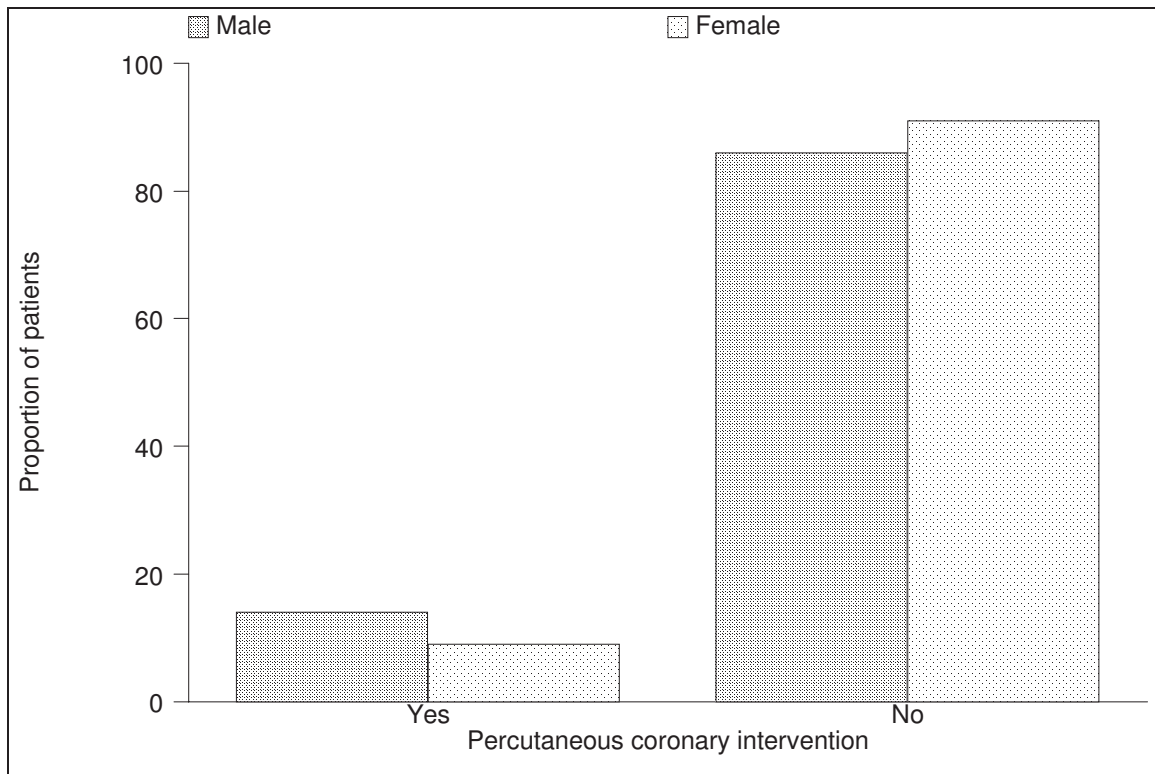


Figure 4.3.2c CABG for patients with NSTEMI/UA by gender, Malaysia 2006

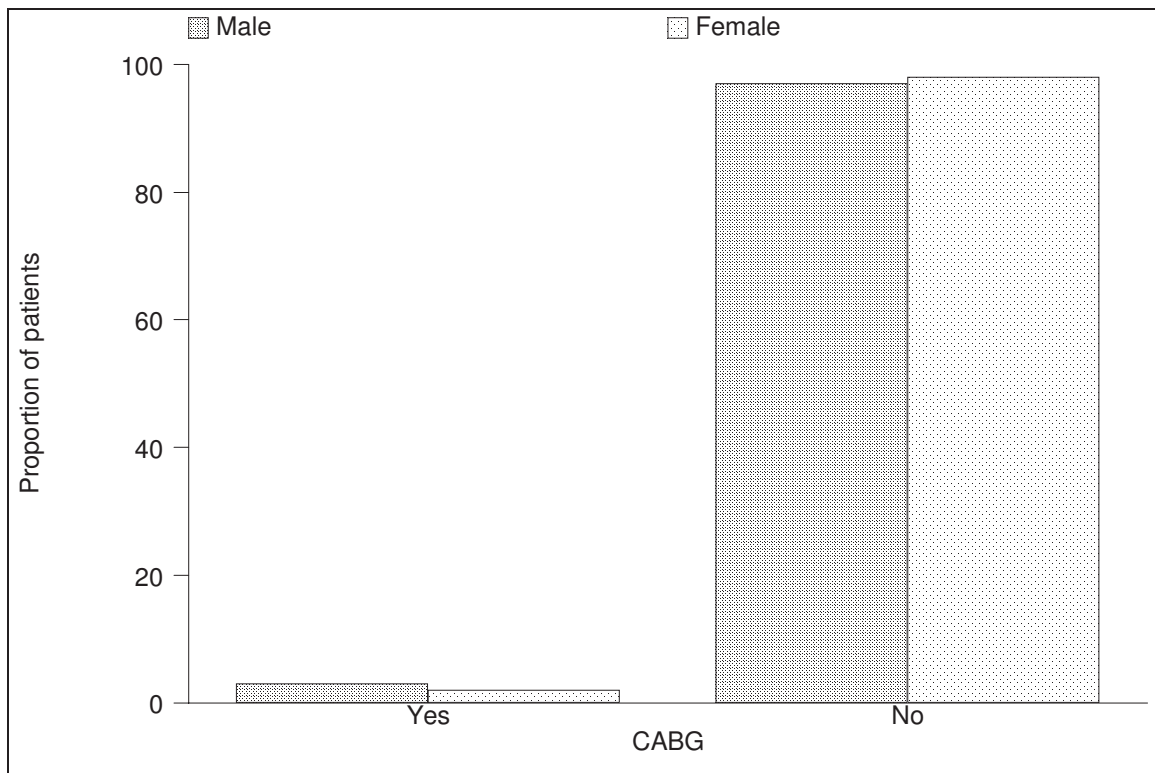


Table 4.3.3 Treatments for patients with NSTEMI/UA by ethnic group, Malaysia 2006

	Malay N=904	Chinese N=485	Indian N=513	Others* N=75
Total admission days**				
• N	879	477	506	72
• Mean, SD	6 (4)	6 (4)	6 (4)	5 (3)
• Median, (min, max)	5 (1,26)	4 (1,30)	5 (1,30)	4 (2,18)
Number of days on CCU				
• N	278	156	115	41
• Mean, SD	4 (3)	3 (2)	4 (3)	3 (2)
• Median, (min, max)	3 (1,24)	2 (1,17)	3 (1,14)	3 (1,9)
Number of days on ICU/CICU				
• N	59	43	17	18
• Mean, SD	4 (3)	3 (3)	4 (3)	3 (3)
• Median, (min, max)	3 (1,23)	3 (1,11)	3 (1,12)	2 (1,10)
Cardiac catheterization, no. %				
• Yes	156 (17)	83 (17)	103 (20)	15 (20)
• No	731 (81)	392 (81)	402 (78)	60 (80)
• No-Transferred to another centre	17 (2)	10 (2)	8 (2)	0 (0)
Percutaneous coronary intervention, no. %				
• Yes	100 (11)	62 (13)	70 (14)	10 (13)
• No	804 (89)	423 (87)	443 (86)	65 (87)
CABG, no. %				
• Yes	29 (3)	16 (3)	11 (2)	1 (1)
• No	875 (97)	469 (97)	502 (98)	74 (99)
Pre-admission aspirin use				
• Yes	401 (44)	183 (38)	233 (45)	20 (27)
• No	335 (37)	194 (40)	155 (30)	41 (55)
• Unknown	168 (19)	108 (22)	125 (24)	14 (19)

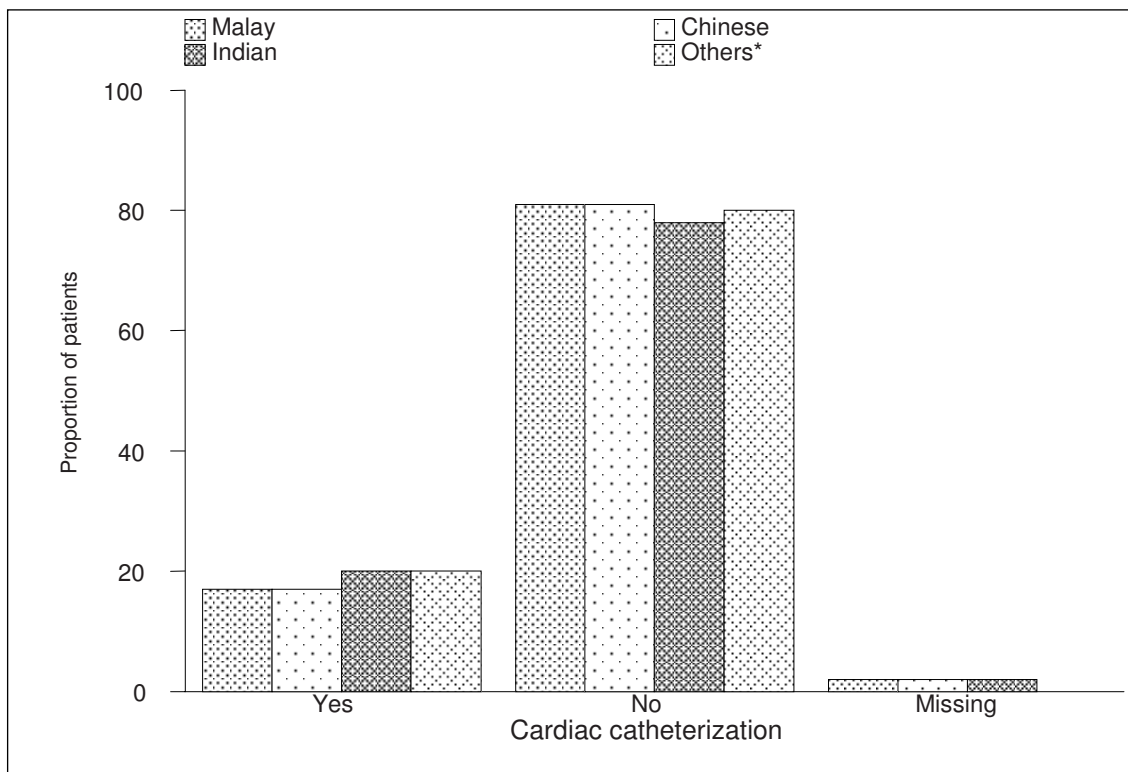
	Malay N=904	Chinese N=485	Indian N=513	Others* N=75
Pharmacological therapy given during admission, no. %				
• ASA	814 (90)	440 (91)	462 (90)	67 (89)
• ADP antagonist	461 (51)	299 (62)	335 (65)	46 (61)
• GP receptor inhibitor	27 (3)	20 (4)	15 (3)	4 (5)
• Unfractionated heparin	284 (31)	47 (10)	64 (12)	5 (7)
• LMWH	495 (55)	369 (76)	390 (76)	50 (67)
• Beta blocker	575 (64)	353 (73)	350 (68)	46 (61)
• ACE inhibitor	510 (56)	259 (53)	303 (59)	35 (47)
• Angiotensin II receptor blocker	82 (9)	49 (10)	63 (12)	7 (9)
• Statin	798 (88)	443 (91)	480 (94)	70 (93)
• Other lipid lowering agent	55 (6)	40 (8)	34 (7)	4 (5)
• Diuretics	330 (37)	172 (35)	181 (35)	22 (29)
• Calcium antagonist	190 (21)	104 (21)	135 (26)	19 (25)
• Oral hypoglycaemic agent	227 (25)	140 (29)	218 (42)	15 (20)
• Insulin	195 (22)	107 (22)	188 (37)	13 (17)
• Anti-arrhythmic agent	60 (7)	34 (7)	22 (4)	5 (7)

* Others includes Orang asli, Kadazan, Melanau, Murut, Bajau, Bidayuh, Iban, other Malaysian and foreigner

** Total admission days is derived as Outcome date – Admission date + 1

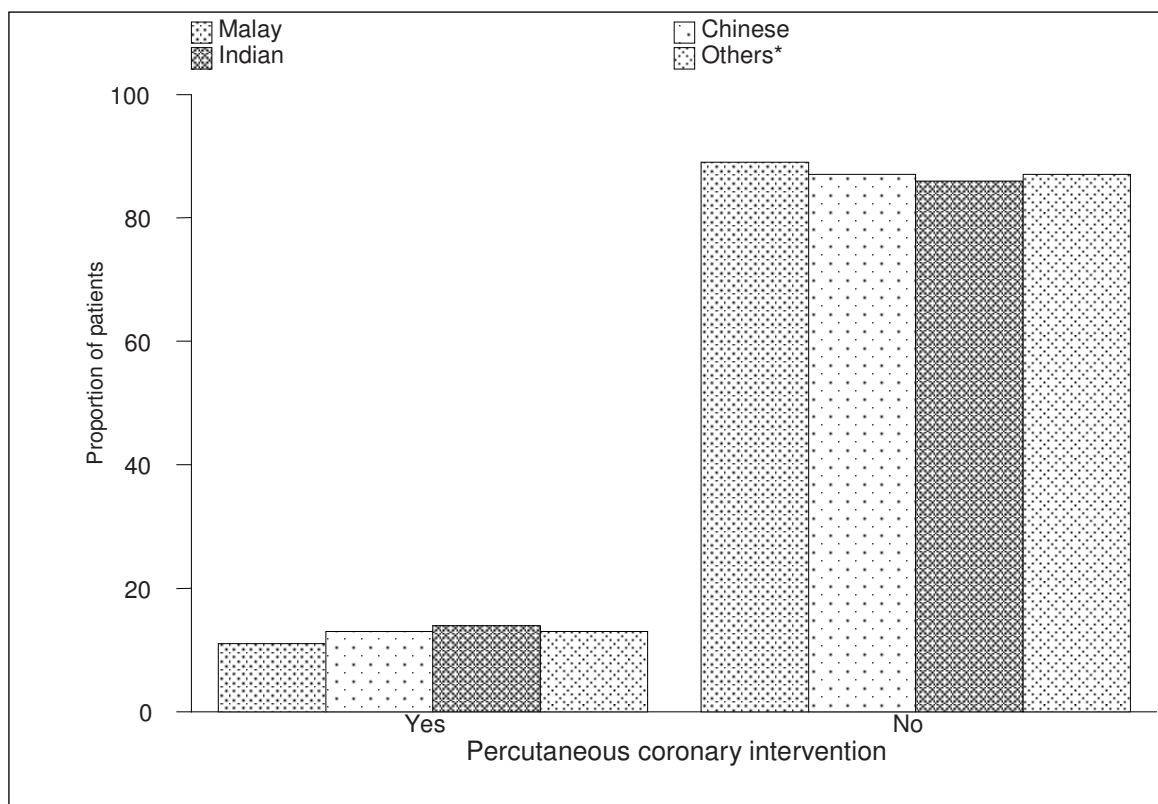
Note: Percentage is to the nearest decimal point.

Figure 4.3.3a Cardiac catheterization for patients with NSTEMI/UA by ethnic group, Malaysia 2006



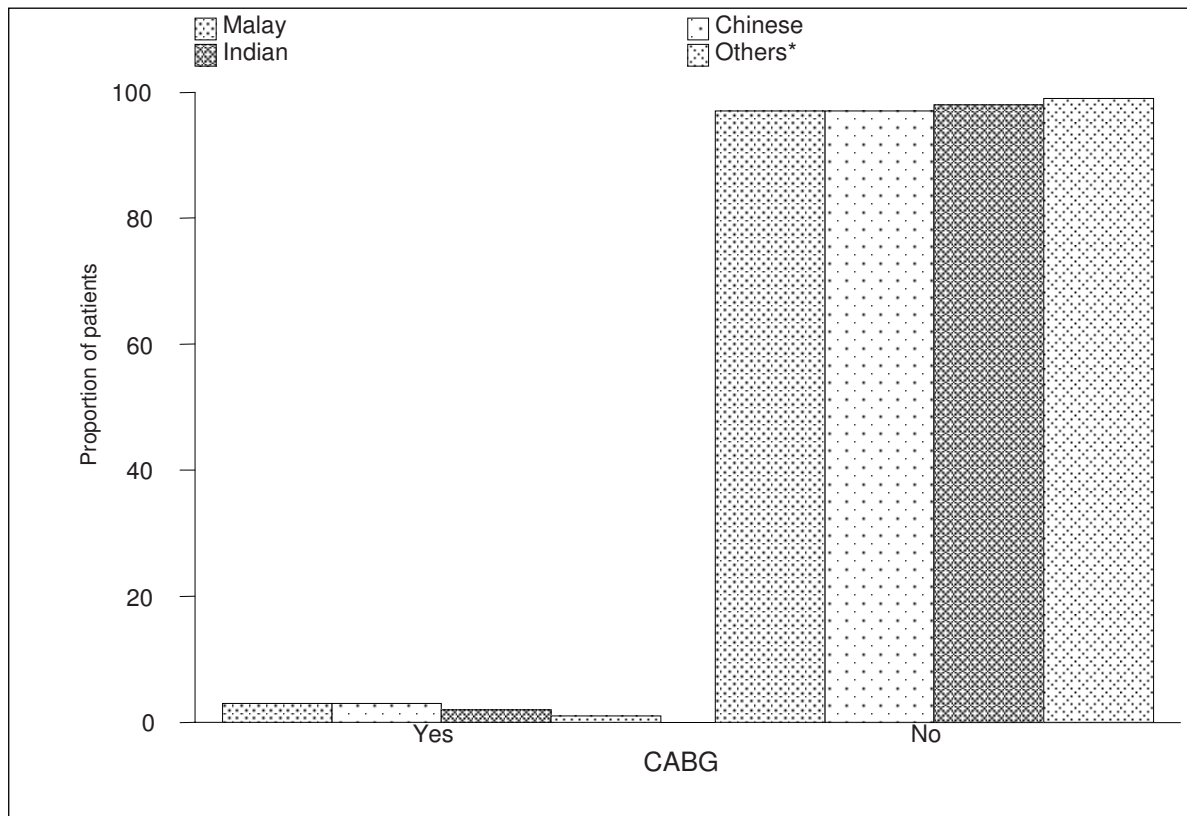
* Others includes Orang asli, Kadazan, Melanau, Murut, Bajau, Bidayuh, Iban, other Malaysian and foreigner

Figure 4.3.3b Percutaneous coronary intervention for patients with NSTEMI/UA by ethnic group, Malaysia 2006



* Others includes Orang asli, Kadazan, Melanau, Murut, Bajau, Bidayuh, Iban, other Malaysian and foreigner

Figure 4.3.3c CABG for patients with NSTEMI/UA by ethnic group, Malaysia 2006



* Others includes Orang asli, Kadazan, Melanau, Murut, Bajau, Bidayuh, Iban, other Malaysian and foreigner

CHAPTER 5

OUTCOMES

Chong Wei Peng

Wan Azman Wan Ahmad

In-hospital and 30-day outcome (Table 5.1 & 5.3)

The in-hospital mortality for the entire cohort of patients was 7% (229 deaths) while 30-day mortality was 8% (288 deaths) (Table 5.1). Based on the ACS stratum, the in-hospital death rates for patients with STEMI, NSTEMI and UA were 9%, 7% and 3% respectively. At 30 days, the death rates were 11%, 8% and 4% respectively (Table 5.3). As nearly a quarter of the patients were lost to follow-up, the 30-day mortalities were likely to be underestimated.

Similar patterns of mortality – highest in STEMI, followed by NSTEMI and lowest in UA, were observed in other prospective surveys of ACS^{2,3,5}.

The mortality rates in the current survey were similar to other registries in the late 90s. In GRACE study⁵, the in-hospital death rates for STEMI, NSTEMI and UA were 7%, 6% and 3% respectively. The National Registry of Myocardial Infarction (NRM) 3 reported a 9% in-hospital death rate for STEMI⁴. But more recent data from the western countries showed a significant reduction in ACS mortality. In-hospital death rates for STEMI and NSTEMI ACS were 4.6% and 2.2% respectively in GRACE study² in year 2005, and 5.3% and 2.5% respectively in the second Euro Heart Survey for ACS³ in 2004.

Outcome by pre-specified variables (Table 5.2.1 to 5.2.5)

The highest in-hospital and 30-day mortality, 10% and 13% respectively were observed in the elderly age group (Table 5.2.1). The mortalities in the young and middle-age groups were similar. Female patients experienced higher in-hospital and 30-day mortality (8% and 10% respectively) compared to the male patients (6% and 8%) (Table 5.2.2). Patients with pre-morbid diabetes mellitus also had a higher in-hospital and 30-day mortality (7% and 10% respectively) than those without diabetes mellitus (5% and 6%) (Table 5.2.3). The mortality in patients with pre-morbid hypertension was similar to those without hypertension (6% in-hospital and 8% 30-day mortality vs. 7% and 8% respectively) (Table 5.2.4). Interestingly, pre-morbid dyslipidaemia was associated with lower in-hospital and 30-day mortality rates (5% and 6% respectively) compared to those without dyslipidaemia (7% and 9%) (Table 5.2.5).

Outcome of STEMI by treatment (Table 5.4.1 to 5.4.4)

In STEMI, the use of fibrinolysis was associated with lower in-hospital and 30-day mortality rates (7% and 9% respectively vs. 13% and 16%) (Table 5.4.1). On the other hand, mortality of patients who had PCI was similar to those who did not have PCI (Table 5.4.2). PCI was performed in 21% of patients as primary PCI, rescue PCI or PCI for post-infarct angina. Therefore these data do not compare primary PCI with fibrinolysis in STEMI. Only 10 patients had CABG during the admission for STEMI, and all 10 were alive upon discharge and at 30 days (Table 5.4.3).

Outcome of NSTEMI/UA by treatment (Table 5.5.1 to 5.5.3)

In NSTEMI and UA, only 12.3% of the patients in this survey had in-hospital PCI, compared to 37.1% reported in the second Euro Heart Survey on ACS³, and 28% (in NSTEMI) and 18% (in UA) in GRACE¹. The in-hospital and 30-day mortalities of patients who had PCI were slightly lower compared to those medically treated (4% vs. 5% and 5% vs. 7% respectively) (Table 5.5.1). In contrast to STEMI, more NSTEMI/UA patients (n=57) underwent CABG during hospitalization for the index event. Both in-hospital and 30-day mortalities were higher in this group of patients (14% for both) compared to those who did not have CABG (5% and 6% respectively) (Table 5.5.2).

Prognostic factors (Table 5.6.1 to 5.6.4)

The following were associated with an increased risk of in-hospital death in patients with STEMI: higher Killip class, higher TIMI risk score, former or current cigarette smoking, family history of premature cardiovascular disease, dyslipidaemia, hypertension and diabetes mellitus. Older age was associated with increased risk of in-hospital death. Prognostic factors for an increased death in 30 days among STEMI patients were almost similar with in-hospital death with the exception of dyslipidemia. In NSTEMI/UA, the following predicts higher in-hospital mortality: higher Killip class, former or current cigarette smoking, diabetes mellitus and heart failure. Older age is again associated with increased risk of death. For 30-day mortality, higher Killip class, cigarette smoking and diabetes mellitus were poor prognostic factors.

Summary Points:

- Total in hospital mortality for patients with ACS was 7% while 30-day mortality was 8%.
- The mortality was higher in STEMI followed by NSTEMI and lowest in UA. Our mortality rates were similar to other Western registries in the late 90s.
- In STEMI, the use of fibrinolysis was associated with lower in-hospital and 30-day mortality rates. In contrast there was no difference in outcome between those who underwent PCI on the same admission and those who did not.
- For STEMI and UA the in-hospital and 30-day mortalities of patients who had PCI were slightly lower compared to medically treated patients.
- Important prognostic factors for STEMI were higher Killip class, higher TIMI risk score and the presence of conventional risk factors. Higher Killip class was also an important prognostic factor for NSTEMI/UA.

References:

1. Fox KAA, Goodman SG, Klein W et al., GRACE investigators. Management of acute coronary syndromes. Variations in practice and outcome. Findings from the Global Registry of Acute Coronary Events (GRACE). *Eur Heart J.* 2002; 23:1177-1189.
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4. Rogers WJ, Canto JG, Lambrew CT et al. Temporal trends in the treatment of over 1.5 million patients with myocardial infarction in the US from 1990 through 1999: the National Registry of Myocardial Infarction 1, 2 and 3. *J Am Coll Cardiol* 2000; 36: 2056-63.
5. Steg PG, Goldberg RJ, Gore JM et al., GRACE Investigators. Baseline characteristics, management practices, and in-hospital outcomes of patients hospitalized with acute coronary syndromes in the Global Registry of Acute Coronary Events (GRACE). *Am J Cardiol.* 2002; 90(4):358-63.

Table 5.1 Overall outcomes for patients with ACS, Malaysia 2006

Outcome	Overall outcome			
	In-hospital		30-day*	
	No.	%	No.	%
• Discharged / Alive	3186	93	2302	67
• Died	229	7	288	8
• Lost to follow-up	NA	NA	832	24
• Missing	7	0	0	0

*Including patients who died in-hospital

Note: Percentage is to the nearest decimal point.

Figure 5.1.1 In-hospital outcomes for patients with ACS, Malaysia 2006

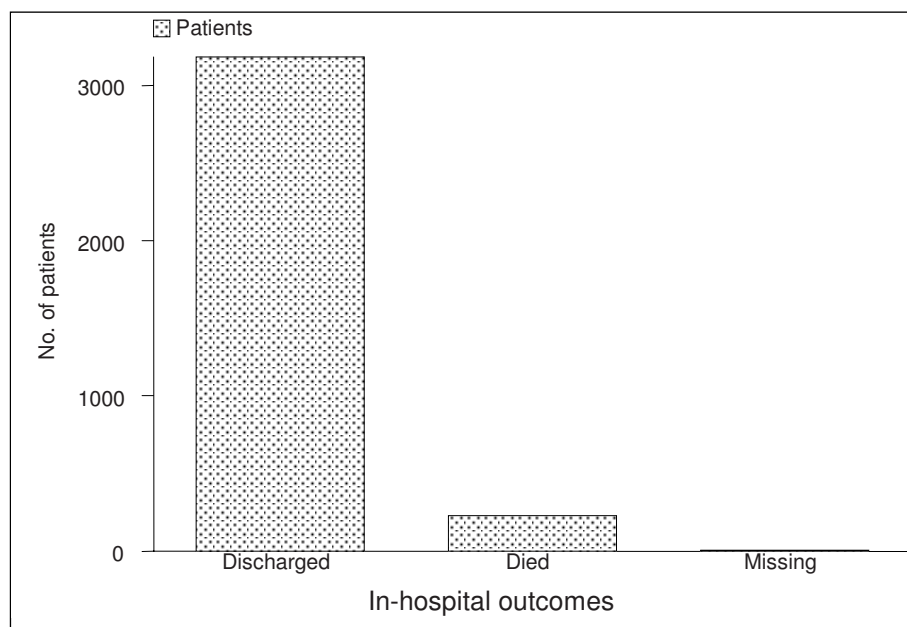


Figure 5.1.2 30-day outcomes for patients with ACS, Malaysia 2006

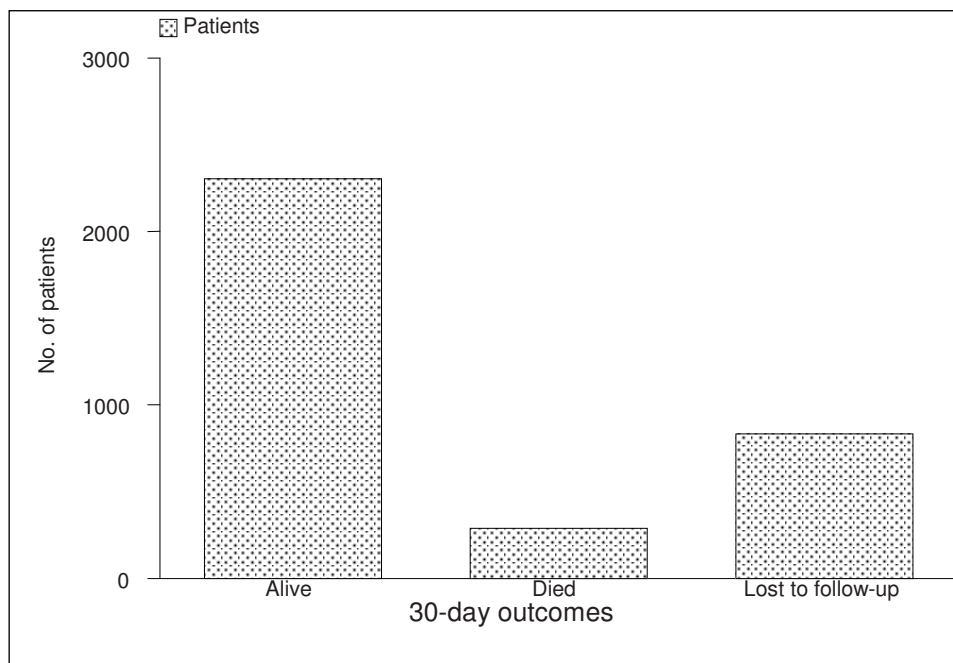


Table 5.2.1 Overall outcomes for patients with ACS by age group (years), Malaysia 2006

Outcome	In-hospital						30-day*					
	Young		Middle-age		elderly		Young		Middle-age		elderly	
	No.	%	No.	%	No.	%	No.	%	No.	%	No.	%
• Discharged / Alive	162	98	1610	96	1414	89	117	70	1139	68	1046	66
• Died	4	2	63	4	162	10	6	4	82	5	200	13
• Lost to follow-up	NA	NA	NA	NA	NA	NA	43	26	454	27	335	21
• Missing	0	0	2	0	5	0	0	0	0	0	0	0

*Including patients who died in-hospital.

Notes:

1. Young is defined as age from 20 to less than 40 years, middle-age is defined as age between 40 to less than 60 years and elderly is defined as 60 years and above.
2. Percentage is to the nearest decimal point.

Figure 5.2.1a In-hospital outcomes for patients with ACS by age group (years), Malaysia 2006

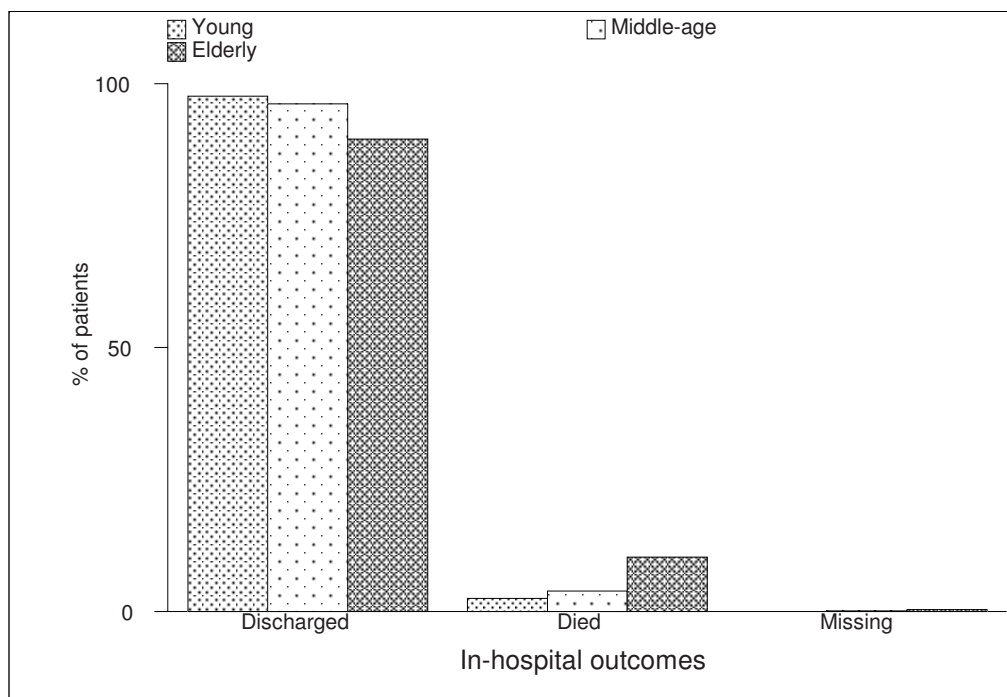


Figure 5.2.1b 30-day outcomes for patients with ACS by age group (years), Malaysia 2006

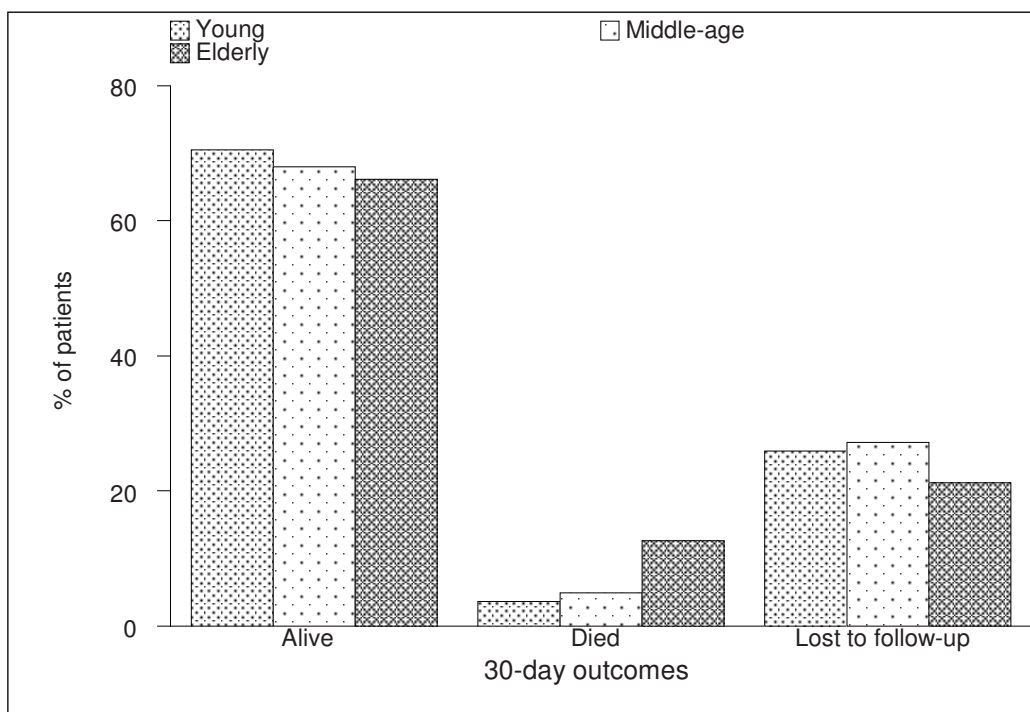


Table 5.2.2 Overall outcomes for patients with ACS by gender, Malaysia 2006

Outcome	In-hospital				30-day*			
	Male		Female		Male		Female	
	No.	%	No.	%	No.	%	No.	%
• Discharged / Alive	2401	93	785	92	1752	68	550	64
• Died	164	6	65	8	203	8	85	10
• Lost to follow-up	NA	NA	NA	NA	614	24	218	26
• Missing	4	0	3	0	0	0	0	0

*Including patients who died in-hospital.

Note: Percentage is to the nearest decimal point.

Figure 5.2.2a In-hospital outcomes for patients with ACS by gender, Malaysia 2006

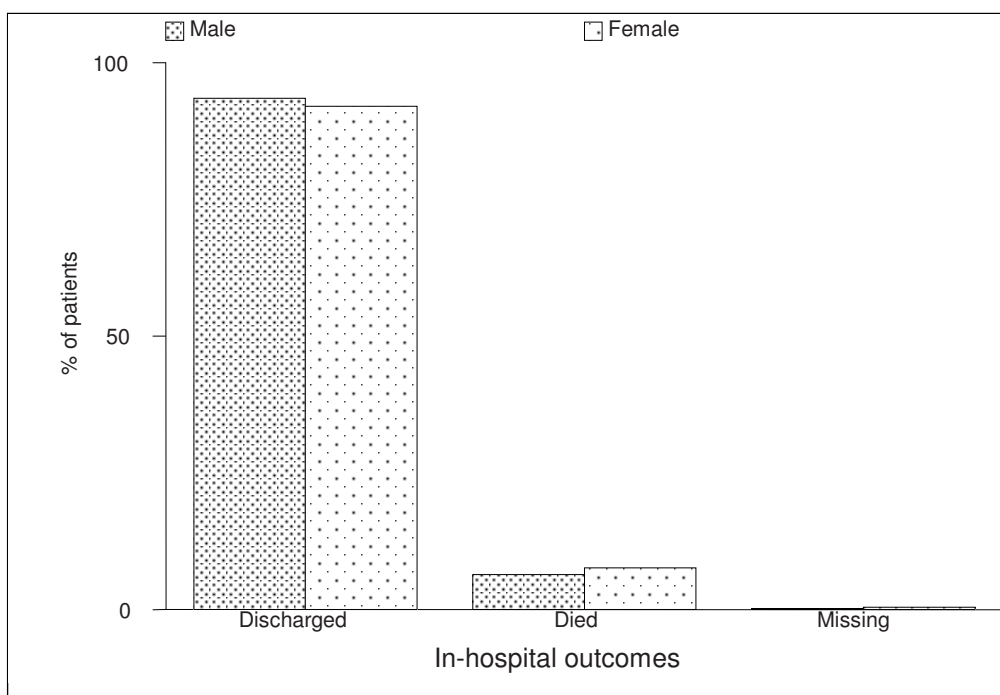


Figure 5.2.2b 30-day outcomes for patients with ACS by gender, Malaysia 2006

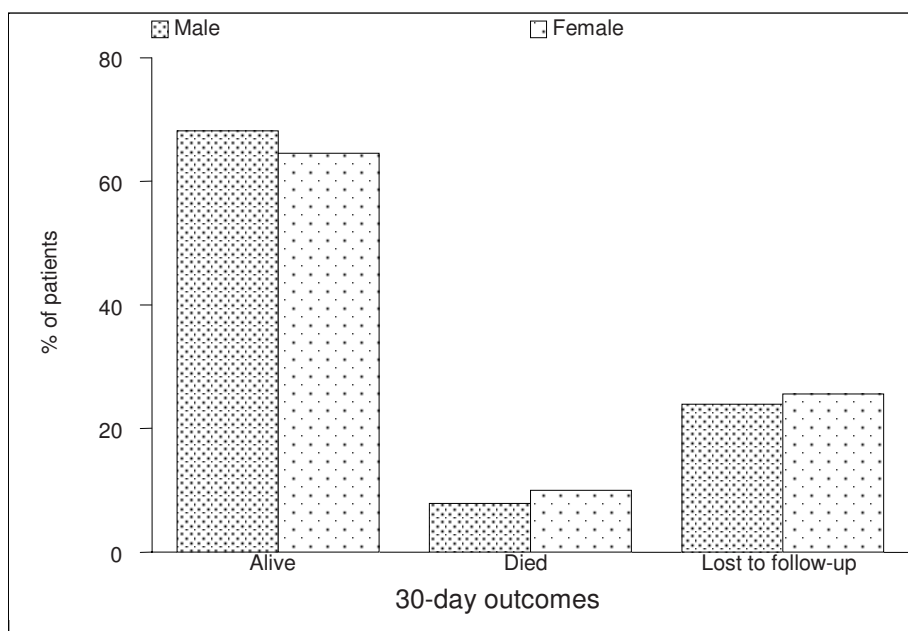


Table 5.2.3 Overall outcomes for patients with ACS by pre-morbid diabetes, Malaysia 2006

Outcome	In-hospital						30-day*					
	Diabetic		Non-diabetic		Not known		Diabetic		Non-diabetic		Not known	
	No.	%	No.	%	No.	%	No.	%	No.	%	No.	%
• Discharged / Alive	1388	93	1160	95	638	91	1010	67	817	67	475	68
• Died	108	7	64	5	57	8	143	10	78	6	67	10
• Lost to follow-up	NA	NA	NA	NA	NA	NA	344	23	331	27	157	22
• Missing	1	0	2	0	4	1	0	0	0	0	0	0

*Including patients who died in-hospital.

Note: Percentage is to the nearest decimal point.

Figure 5.2.3a In-hospital outcomes for patients with ACS by pre-morbid diabetes, Malaysia 2006

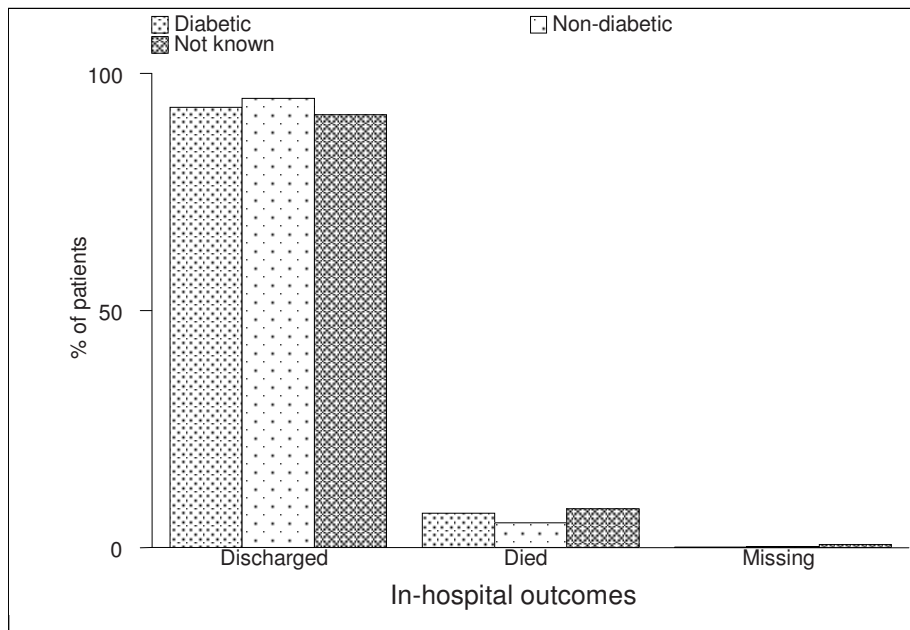


Figure 5.2.3b 30-day outcomes for patients with ACS by pre-morbid diabetes, Malaysia 2006

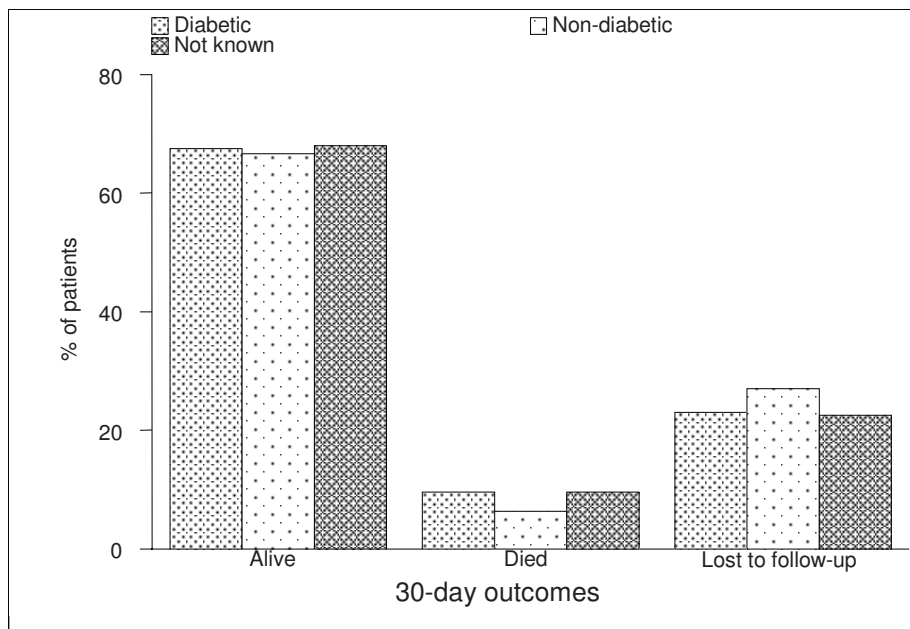


Table 5.2.4 Overall outcomes for patients with ACS by pre-morbid hypertension, Malaysia 2006

Outcome	In-hospital						30-day*					
	Hypertensive		Non-hypertensive		Not known		Hypertensive		Non-hypertensive		Not known	
	No.	%	No.	%	No.	%	No.	%	No.	%	No.	%
• Discharged / Alive	1947	93	733	93	506	92	1425	68	510	65	367	66
• Died	134	6	52	7	43	8	171	8	63	8	54	10
• Lost to follow-up	NA	NA	NA	NA	NA	NA	488	23	213	27	131	24
• Missing	3	0	1	0	3	1	0	0	0	0	0	0

*Including patients who died in-hospital.

Note: Percentage is to the nearest decimal point.

Figure 5.2.4a In-hospital outcomes for patients with ACS by pre-morbid hypertension, Malaysia 2006

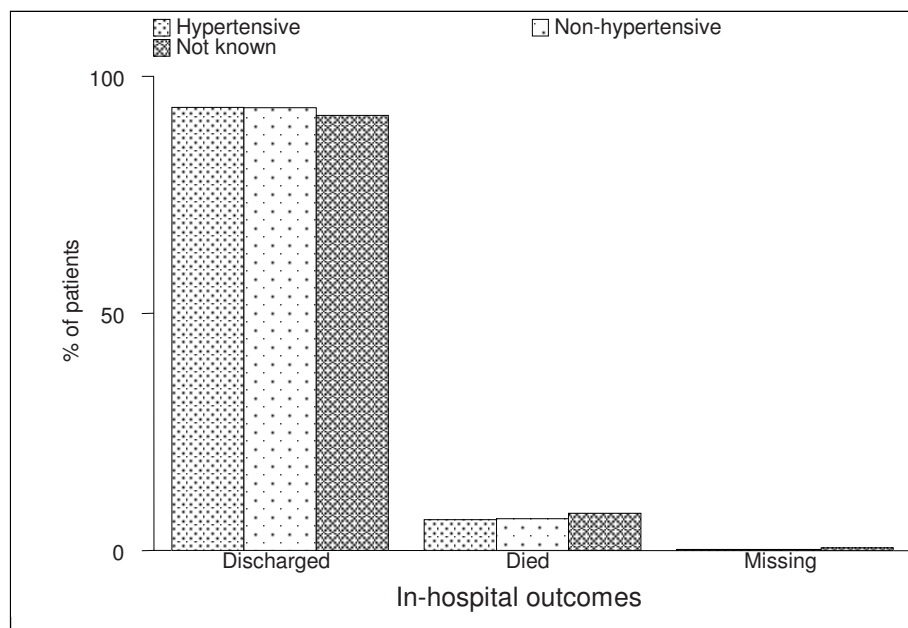


Figure 5.2.4b 30-day outcomes for patients with ACS by pre-morbid hypertension, Malaysia 2006

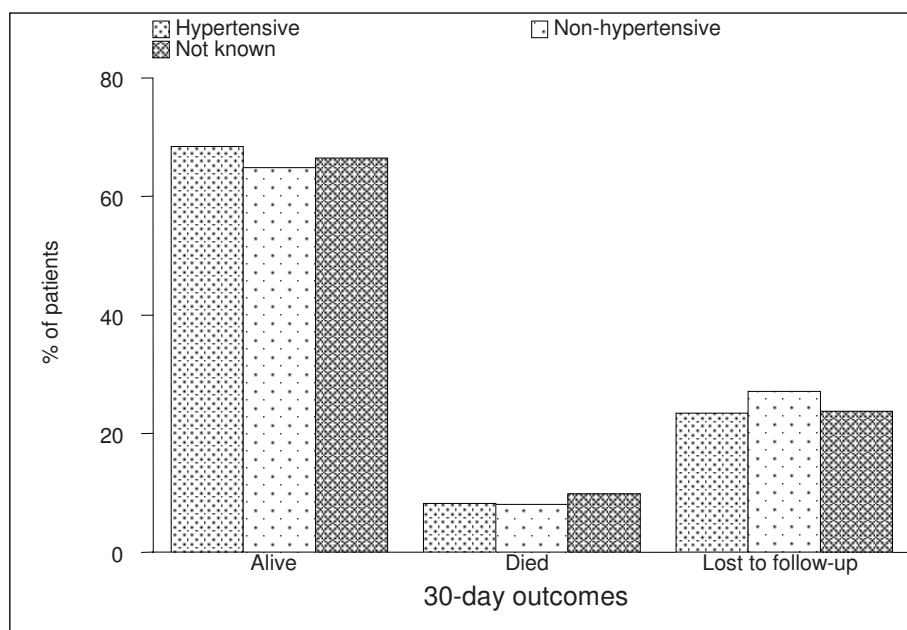


Table 5.2.5 Overall outcomes for patients with ACS by pre-morbid dyslipidaemia, Malaysia 2006

Outcome	In-hospital						30-day*					
	Dyslipidaemia						Dyslipidaemia					
	Yes		No		Not known		Yes		No		Not known	
	No.	%	No.	%	No.	%	No.	%	No.	%	No.	%
• Discharged / Alive	1075	95	838	93	1273	92	843	75	546	61	913	66
• Died	55	5	63	7	111	8	70	6	82	9	136	10
• Lost to follow-up	NA	NA	NA	NA	NA	NA	218	19	274	30	340	24
• Missing	1	0	1	0	5	0	0	0	0	0	0	0

*Including patients who died in-hospital.

Note: Percentage is to the nearest decimal point.

Figure 5.2.5a In-hospital outcomes for patients with ACS by pre-morbid dyslipidaemia, Malaysia 2006

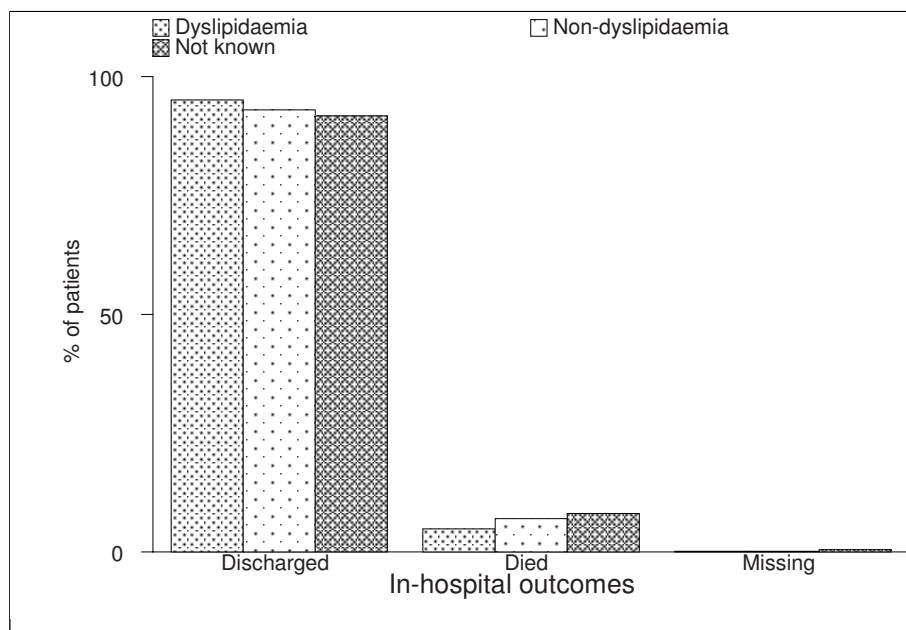


Figure 5.2.5b 30-day outcomes for patients with ACS by pre-morbid dyslipidaemia, Malaysia 2006

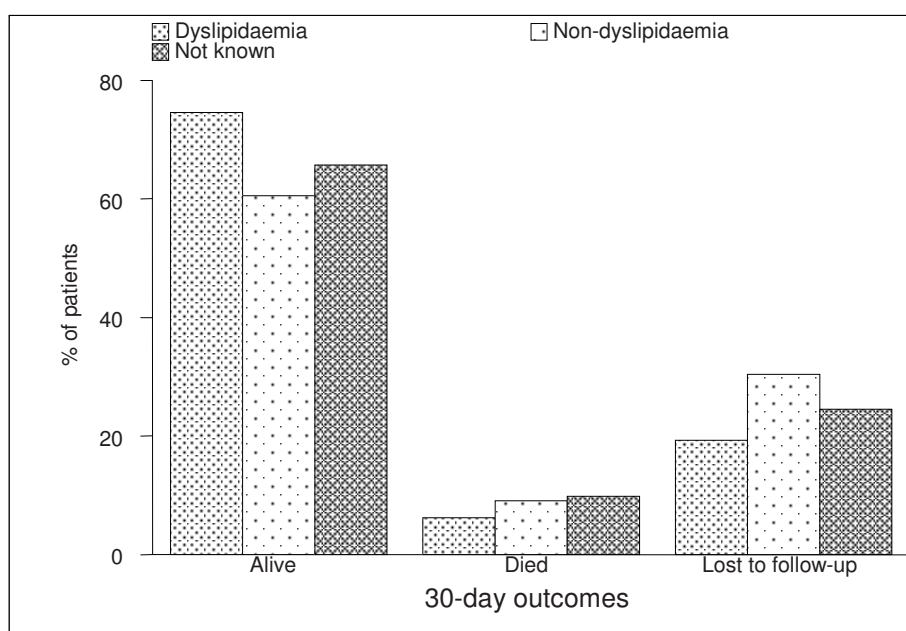


Table 5.3 Overall outcomes for patients with ACS by ACS stratum, Malaysia 2006

Outcome	In-hospital						30-day*					
	STEMI		NSTEMI		UA		STEMI		NSTEMI		UA	
	No.	%	No.	%	No.	%	No.	%	No.	%	No.	%
• Discharged / Alive	1312	91	1056	93	818	97	939	65	796	70	567	67
• Died	129	9	75	7	25	3	158	11	92	8	38	4
• Lost to follow-up	NA	NA	NA	NA	NA	NA	348	24	244	22	240	28
• Missing	4	0	1	0	2	0	0	0	0	0	0	0

*Including patients who died in-hospital.

Note: Percentage is to the nearest decimal point.

Figure 5.3.1 In-hospital outcomes for patients with ACS by ACS stratum, Malaysia 2006

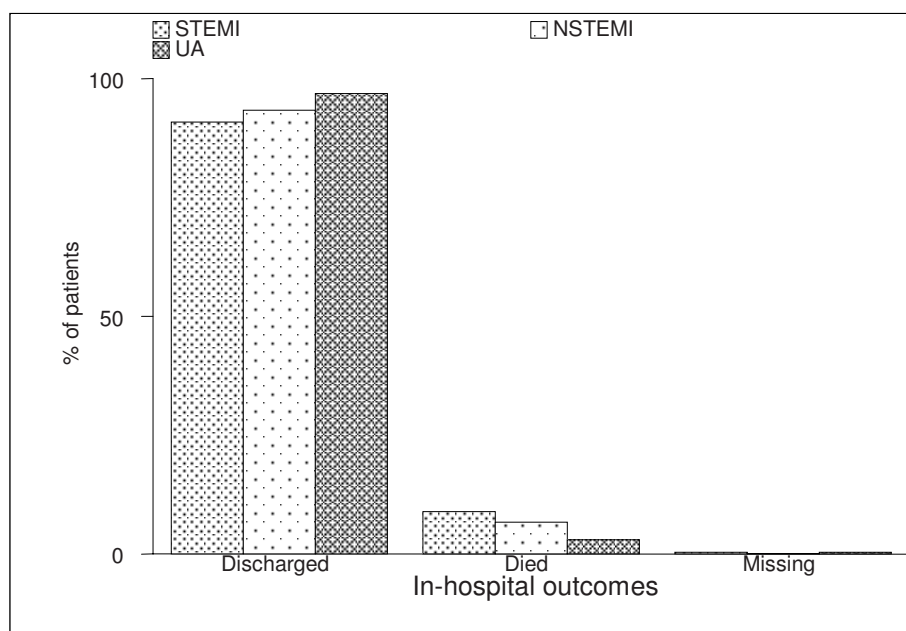


Figure 5.3.2 30-day outcomes for patients with ACS by ACS stratum, Malaysia 2006

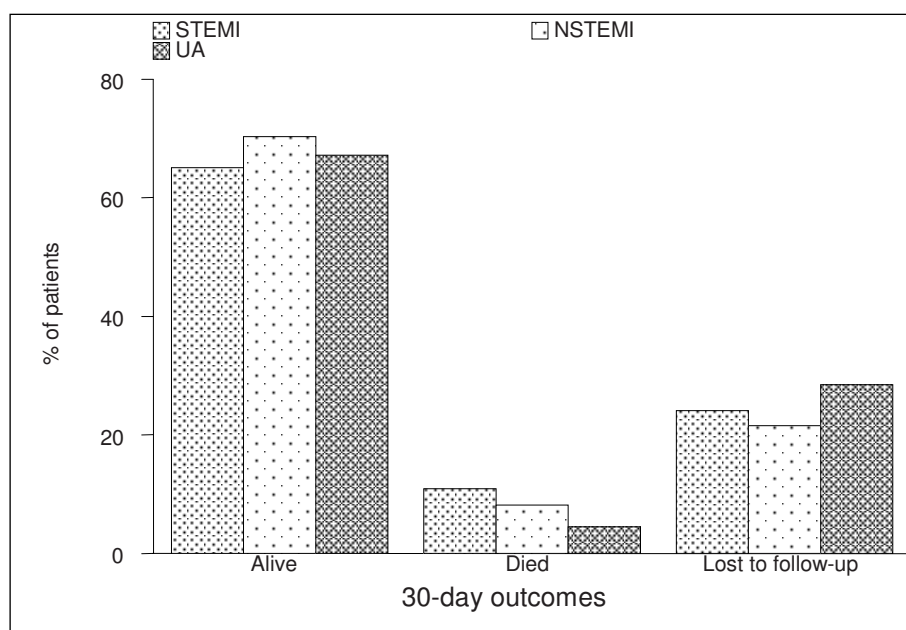


Table 5.4.1 Overall outcomes for patients with STEMI by fibrinolytic therapy, Malaysia 2006

Outcome	In-hospital				30-day*			
	Fibrinolytic therapy				Fibrinolytic therapy			
	Yes		No		Yes		No	
	No.	%	No.	%	No.	%	No.	%
• Discharged / Alive	940	92	372	87	686	67	253	59
• Died	74	7	55	13	90	9	68	16
• Lost to follow-up	NA	NA	NA	NA	242	24	106	25
• Missing	4	0	0	0	0	0	0	0

*Including patients who died in-hospital.

Note: Percentage is to the nearest decimal point.

Figure 5.4.1a In-hospital outcomes for patients with STEMI by fibrinolytic therapy, Malaysia 2006

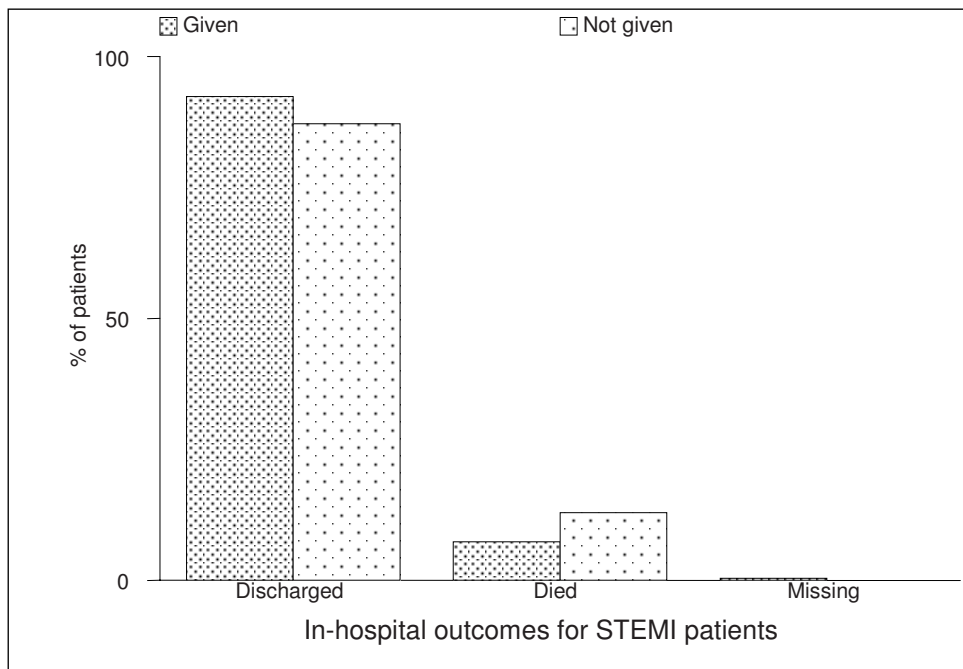


Figure 5.4.1b 30-day outcomes for patients with STEMI by fibrinolytic therapy, Malaysia 2006

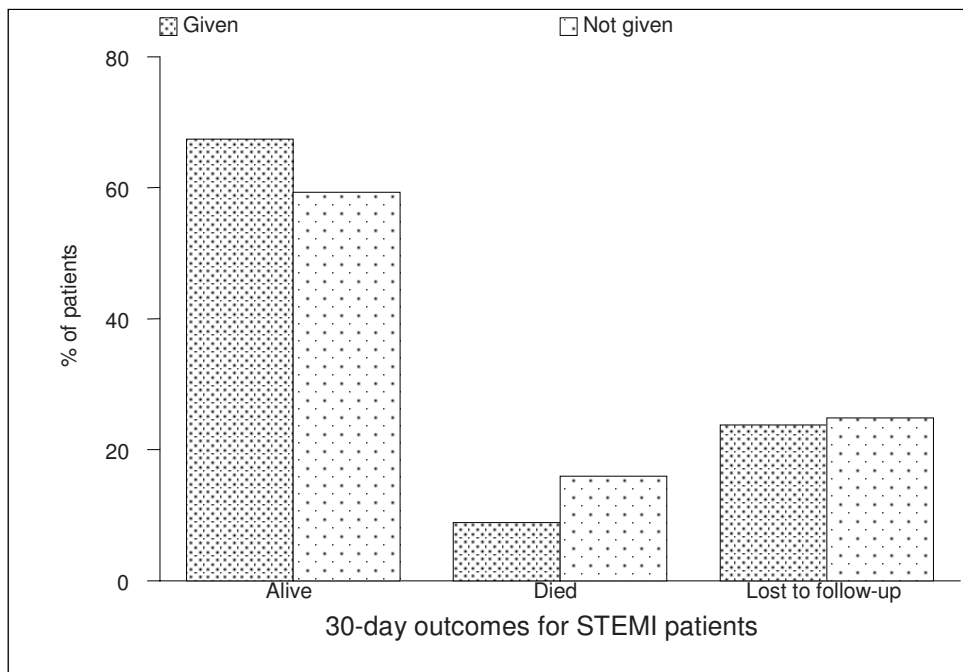


Table 5.4.2 Overall outcomes for patients with STEMI by percutaneous coronary intervention at admission, Malaysia 2006

Outcome	In-hospital				30-day*			
	Percutaneous coronary intervention				Percutaneous coronary intervention			
	Yes		No		Yes		No	
	No.	%	No.	%	No.	%	No.	%
• Discharged / Alive	283	92	1029	91	242	79	697	61
• Died	25	8	104	9	34	11	124	11
• Lost to follow-up	NA	NA	NA	NA	32	10	316	28
• Missing	0	0	4	0	0	0	0	0

*Including patients who died in-hospital.

Notes:

1. Percentage is to the nearest decimal point.

2. Percutaneous Coronary Intervention includes primary, rescue and facilitated intervention

Figure 5.4.2a In-hospital outcomes for patients with STEMI by percutaneous coronary intervention at admission, Malaysia 2006

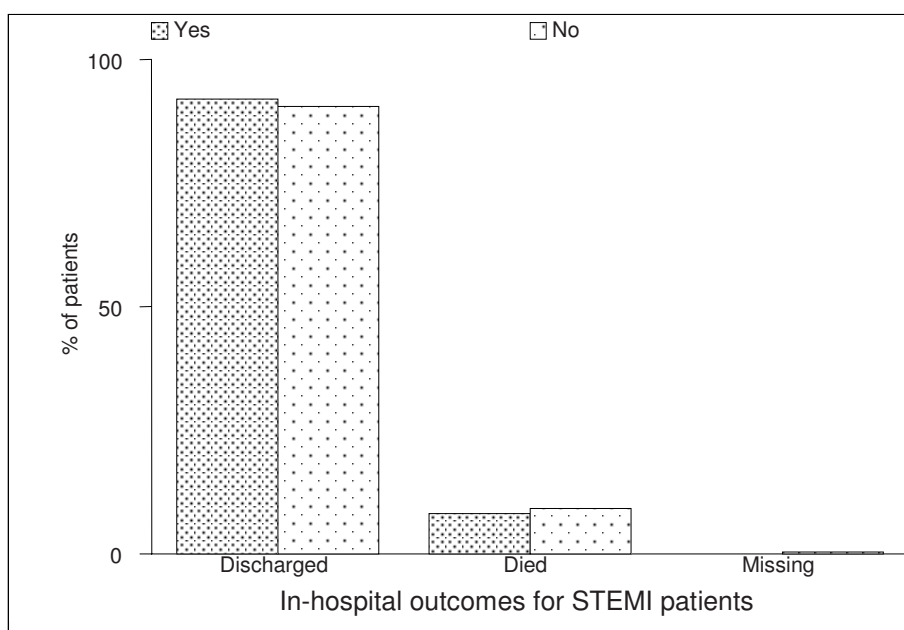


Figure 5.4.2b 30-day outcomes for patients with STEMI by percutaneous coronary intervention at admission, Malaysia 2006

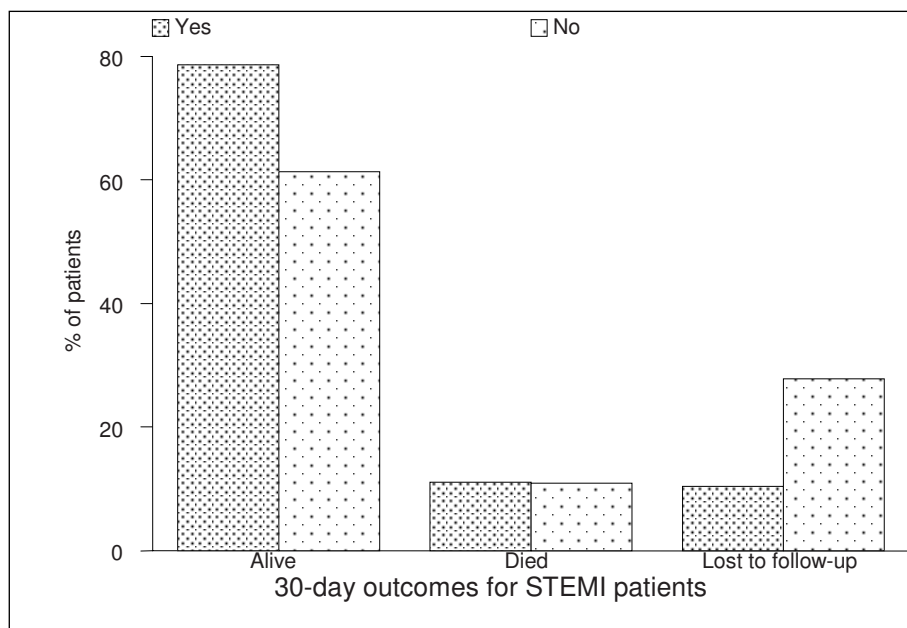


Table 5.4.3 Overall outcomes for patients with STEMI by CABG at admission, Malaysia 2006

Outcome	In-hospital				30-day*			
	CABG				CABG			
	Yes		No		Yes		No	
	No.	%	No.	%	No.	%	No.	%
• Discharged / Alive	10	100	1302	91	10	100	929	65
• Died	0	0	129	9	0	0	158	11
• Lost to follow-up	NA	NA	NA	NA	0	0	348	24
• Missing	0	0	4	0	0	0	0	0

*Including patients who died in-hospital.

Note: Percentage is to the nearest decimal point.

Figure 5.4.3a In-hospital outcomes for patients with STEMI by CABG at admission, Malaysia 2006

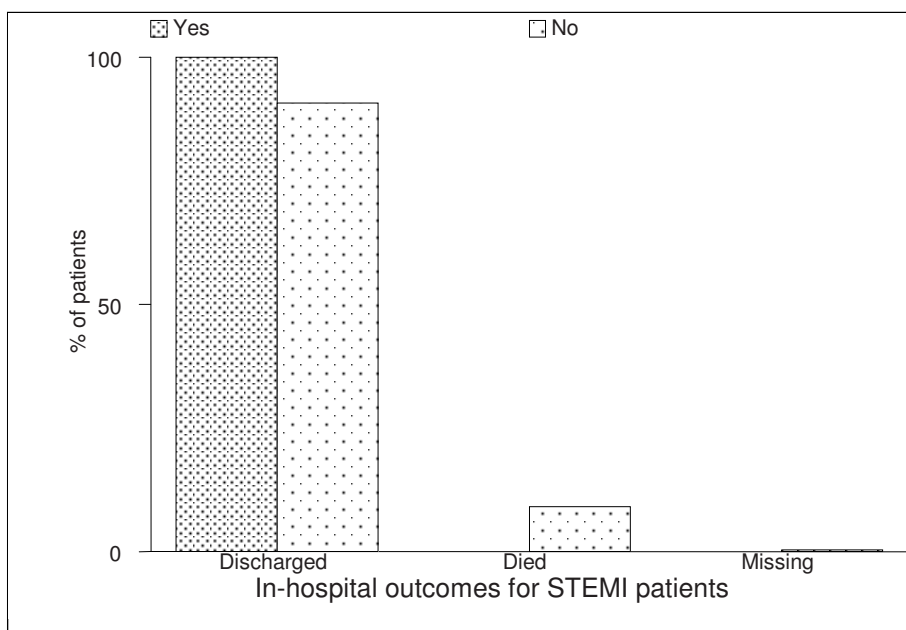


Figure 5.4.3b 30-day outcomes for patients with STEMI by CABG at admission, Malaysia 2006

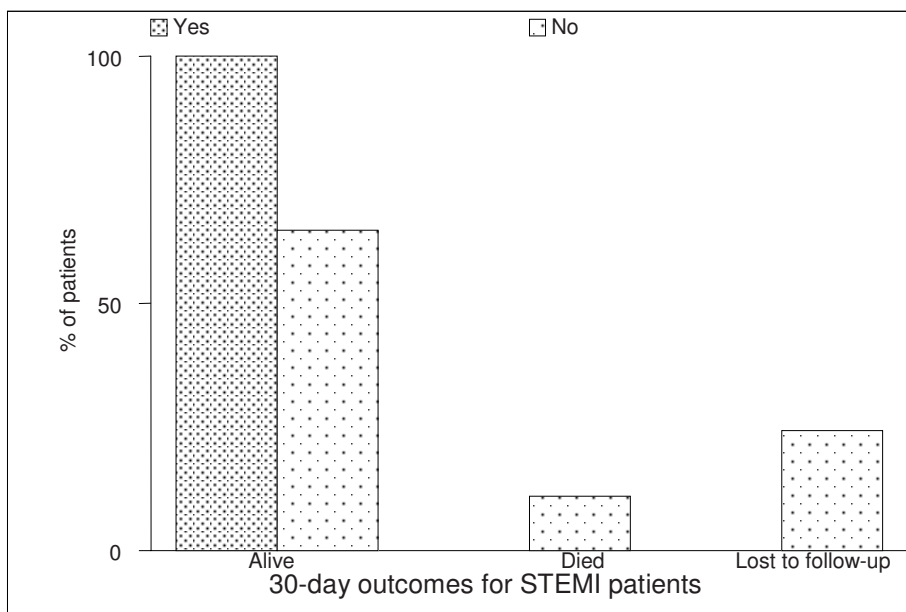


Table 5.4.4 Overall outcomes for patients with STEMI by pre-admission aspirin use, Malaysia 2006

Outcome	In-hospital						30-day*					
	Pre-admission aspirin use						Pre-admission aspirin use					
	Yes		No		Unknown		Yes		No		Unknown	
	No.	%	No.	%	No.	%	No.	%	No.	%	No.	%
• Discharged / Alive	200	88	885	92	227	90	128	56	660	68	151	60
• Died	27	12	76	8	26	10	33	15	95	10	30	12
• Lost to follow-up	NA	NA	NA	NA	NA	NA	66	29	210	22	72	28
• Missing	0	0	4	0	0	0	0	0	0	0	0	0

*Including patients who died in-hospital.

Note: Percentage is to the nearest decimal point.

Figure 5.4.4a In-hospital outcomes for patients with STEMI by pre-admission aspirin use, Malaysia 2006

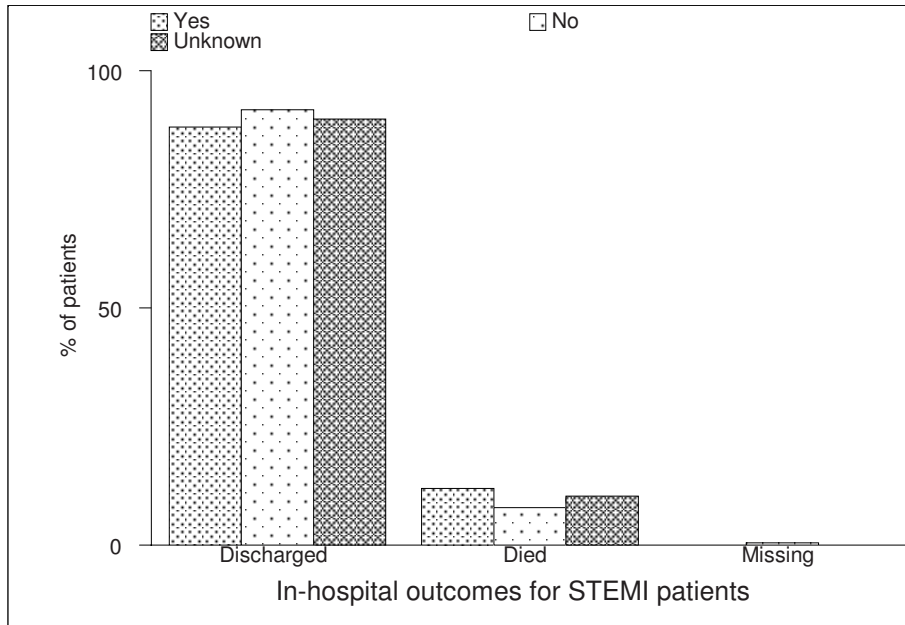


Figure 5.4.4b 30-day outcomes for patients with STEMI by pre-admission aspirin use, Malaysia 2006

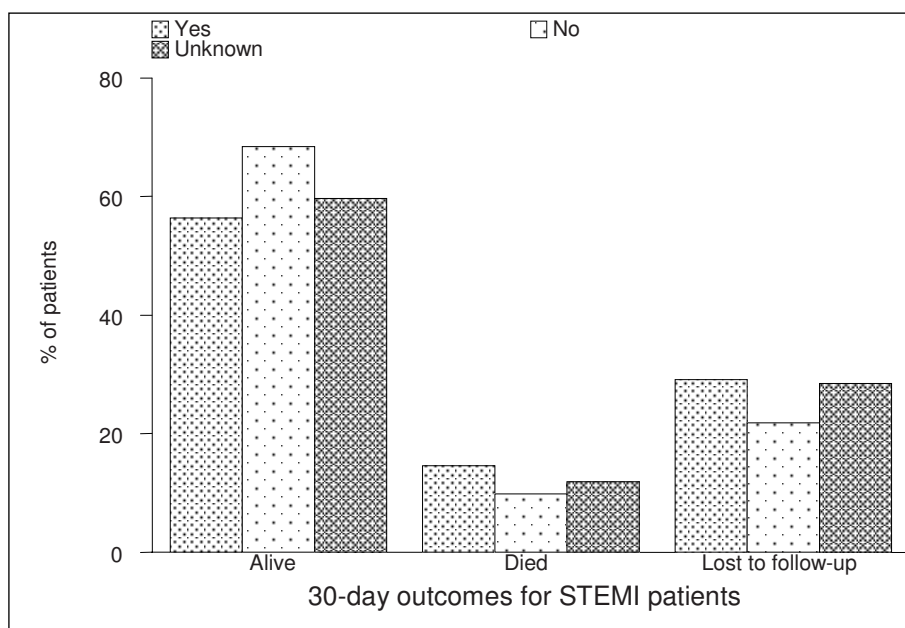


Table 5.5.1 Overall outcomes for patients with NSTEMI/UA by percutaneous coronary intervention, Malaysia 2006

Outcome	In-hospital				30-day*			
	Percutaneous coronary intervention				Percutaneous coronary intervention			
	Yes		No		Yes		No	
	No.	%	No.	%	No.	%	No.	%
• Discharged / Alive	233	96	1641	95	207	86	1156	67
• Died	9	4	91	5	12	5	118	7
• Lost to follow-up	NA	NA	NA	NA	23	10	461	27
• Missing	0	0	3	0	0	0	0	0

*Including patients who died in-hospital.

Note: Percentage is to the nearest decimal point.

Figure 5.5.1a In-hospital outcomes for patients with NSTEMI/UA by percutaneous coronary intervention, Malaysia 2006

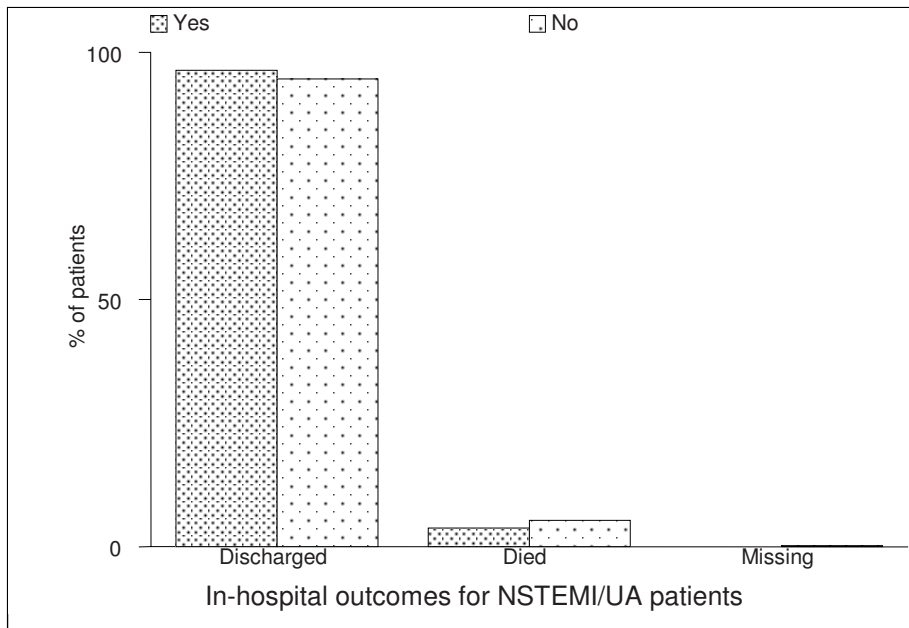


Figure 5.5.1b 30-day outcomes for patients with NSTEMI/UA by percutaneous coronary intervention, Malaysia 2006

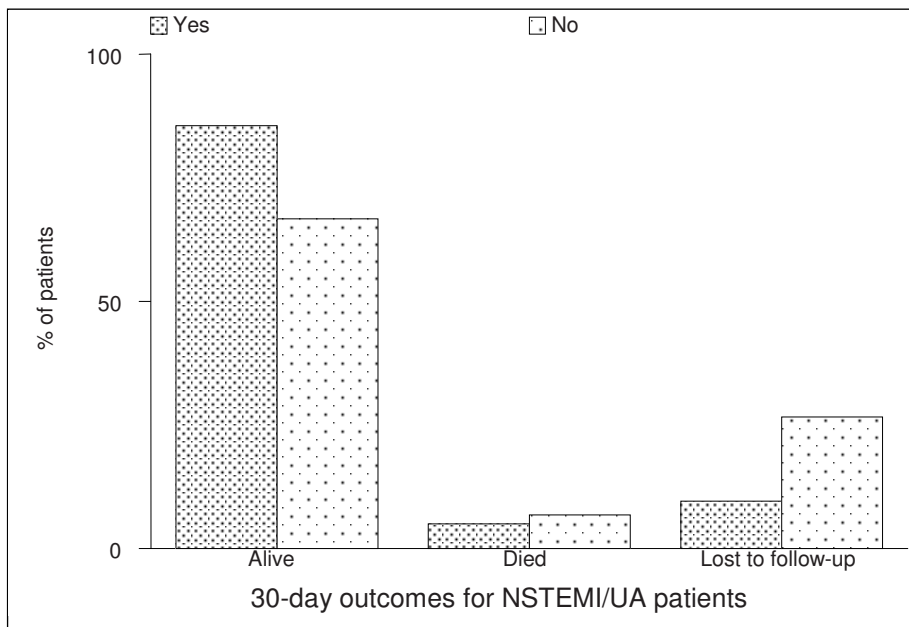


Table 5.5.2 Overall outcomes for patients with NSTEMI/UA by CABG, Malaysia 2006

Outcome	In-hospital				30-day*			
	CABG				CABG			
	Yes		No		Yes		No	
	No.	%	No.	%	No.	%	No.	%
• Discharged / Alive	49	86	1825	95	47	82	1316	69
• Died	8	14	92	5	8	14	122	6
• Lost to follow-up	NA	NA	NA	NA	2	4	482	25
• Missing	0	0	3	0	0	0	0	0

*Including patients who died in-hospital.

Note: Percentage is to the nearest decimal point.

Figure 5.5.2a In-hospital outcomes for patients with NSTEMI/UA by CABG, Malaysia 2006

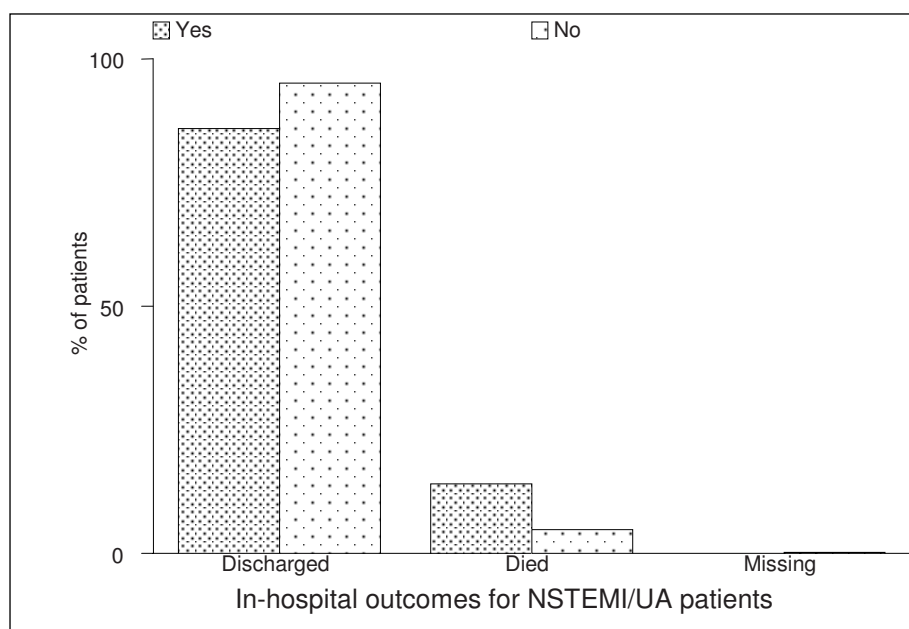


Figure 5.5.2b 30-day outcomes for patients with NSTEMI/UA by CABG, Malaysia 2006

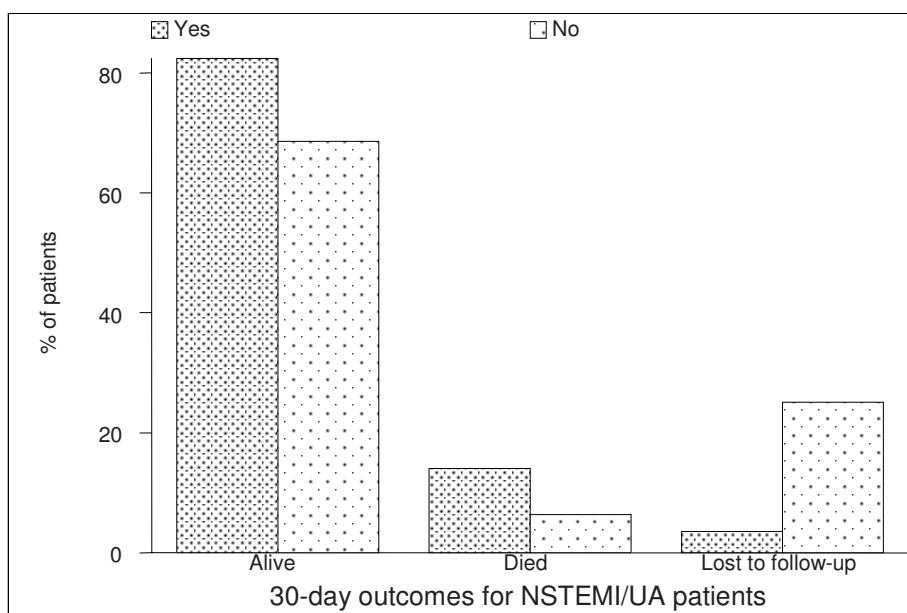


Table 5.5.3 Overall outcomes for patients with NSTEMI by pre-admission aspirin use, Malaysia 2006

Outcome	In-hospital						30-day*					
	Pre-admission aspirin use						Pre-admission aspirin use					
	Yes		No		Unknown		Yes		No		Unknown	
	No.	%	No.	%	No.	%	No.	%	No.	%	No.	%
• Discharged / Alive	796	95	688	95	390	94	582	70	501	69	280	67
• Died	41	5	34	5	25	6	56	7	44	6	30	7
• Lost to follow-up	NA	NA	NA	NA	NA	NA	199	24	180	25	105	25
• Missing	0	0	3	0	0	0	0	0	0	0	0	0

*Including patients who died in-hospital.

Note: Percentage is to the nearest decimal point.

Figure 5.5.3a In-hospital outcomes for patients with NSTEMI/UA by pre-admission aspirin use, Malaysia 2006

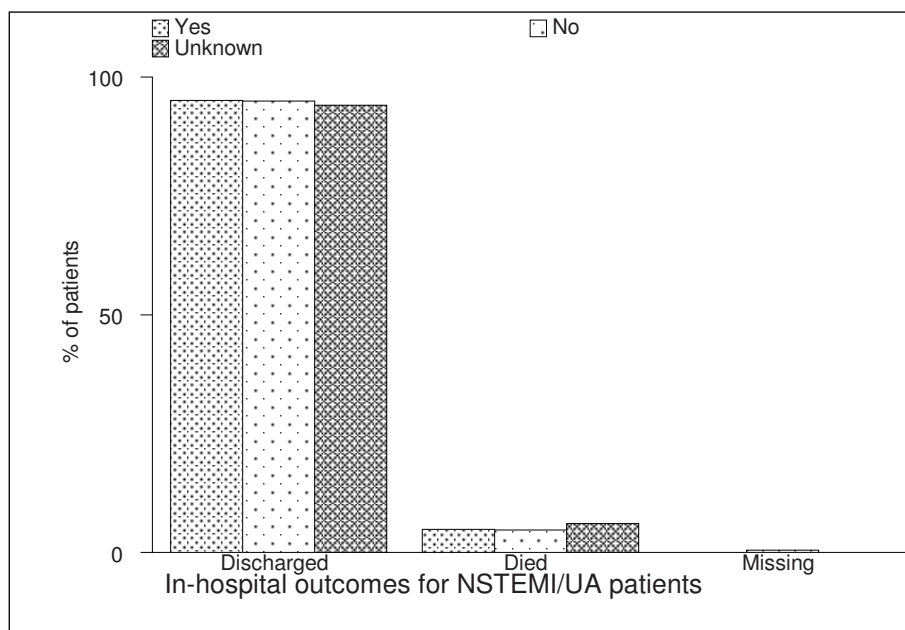


Figure 5.5.3b 30-day outcomes for patients with NSTEMI/UA by pre-admission aspirin use, Malaysia 2006

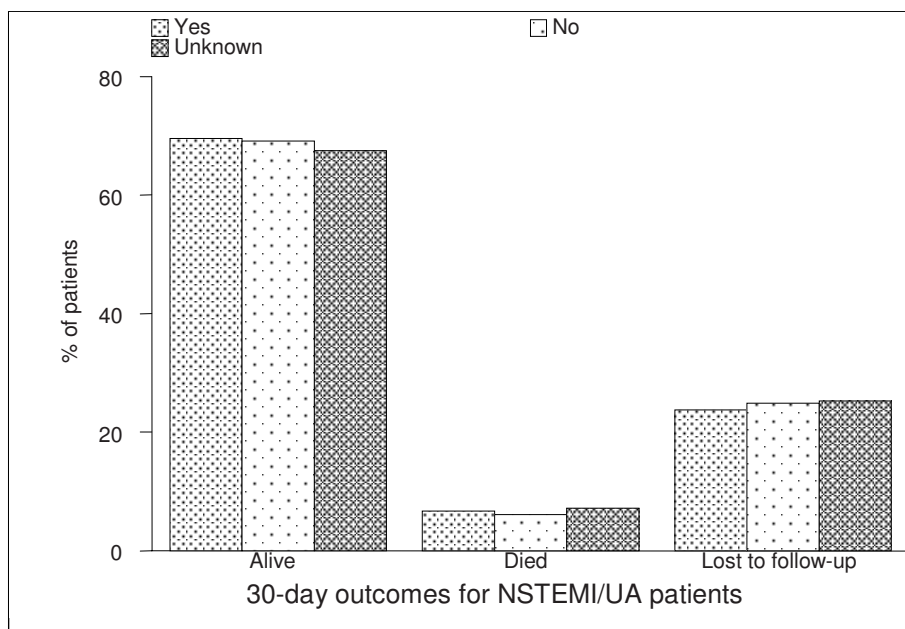


Table 5.6.1 Prognostic factors for death in hospital among STEMI patients, Malaysia 2006

Factors	N	Odd ratio	95% CI	P-value
Age group, years				
• 20 - <40 (ref)	113	1.00	-	-
• 40 - < 60	801	1.02	(0.29, 3.67)	0.97
• > =60	527	1.93	(0.53, 0.70)	0.32
Gender				
• Male (ref)	1226	1.00	-	-
• Female	215	1.23	(0.64, 2.39)	0.53
Ethnic group*				
• Malay	777	0.82	(0.32, 2.13)	0.69
• Chinese	300	0.81	(0.29, 2.35)	0.68
• Indian	286	0.72	(0.26, 2.01)	0.53
• Others (ref)	78	1.00	-	-
Killip classification code				
• I (ref)	891	1.00	-	-
• II	288	2.02	(1.17, 3.50)	0.01
• III	62	3.37	(1.54, 7.37)	0.002
• IV	66	8.50	(4.18, 17.27)	<0.001
• Not stated/ inadequately described	134	0.77	(0.30, 2.01)	0.59
Percutaneous coronary intervention				
• Yes	308	0.94	(0.39, 2.26)	0.88
• No (ref)	1133	1.00	-	-
Cardiac catheterization				
• Yes	298	0.84	(0.34, 2.09)	0.71
• No (ref)	1143	1.00	-	-
TIMI risk score				
• 0-2 (ref)	641	1.00	-	-
• 3-4	375	1.09	(0.55, 2.18)	0.80
• 5-7	337	2.03	(1.09, 3.78)	0.03
• >7	88	6.78	(3.22, 14.28)	<0.001
Fibrinolytic therapy				
• Given	1014	0.67	(0.43, 1.05)	0.08
• Not given (ref)	427	1.00	-	-

Factors	N	Odd ratio	95% CI	P-value
Smoking				
• Never (ref)	417	1.00	-	-
• Former (quit >30 days)	270	4.43	(1.89, 10.40)	0.001
• Current (any tobacco use within last 30 days)	721	3.37	(1.46, 7.82)	0.01
• Unknown	33	2.37	(0.80, 7.02)	0.12
Family history of premature cardiovascular disease				
• Yes	168	3.28	(1.23, 8.76)	0.02
• No (ref)	742	1.00	-	-
• Unknown	531	0.75	(0.40, 1.38)	0.35
Dyslipidaemia				
• Yes	278	2.56	(1.06, 6.15)	0.04
• No (ref)	458	1.00	-	-
• Unknown	705	1.20	(0.67, 2.18)	0.53
Hypertension				
• Yes	680	5.15	(2.24, 11.84)	<0.001
• No (ref)	433	1.00	-	-
• Unknown	328	1.00	(0.43, 2.32)	0.99
Diabetes				
• Yes	525	6.16	(2.82, 13.45)	<0.001
• No (ref)	538	1.00	-	-
• Unknown	378	1.57	(0.73, 3.39)	0.25
Heart failure				
• Yes	48	0.99	(0.38, 2.57)	0.99
• No (ref)	1008	1.00	-	-
• Unknown	385	1.30	(0.58, 2.92)	0.53
Coronary artery disease**				
• Yes	779	1.06	(0.48, 2.36)	0.88
• No (ref)	356	1.00	-	-
• Unknown	306	1.15	(0.46, 2.90)	0.77

*Others include Orang asli, Kadazan, Melanau, Murut, Bajau, Bidayuh, Iban, other Malaysian and foreigner.

**Coronary artery disease is defined as "Yes" on any of the following co-morbidities: 1) History of myocardial infarction, 2) Documented CAD >50% stenosis, 3) Chronic angina (onset more than 2 weeks ago), 4) New onset angina (less than 2 weeks).

Table 5.6.2 Prognostic factors for death in hospital among NSTEMI/UA patients, Malaysia 2006

Factors	N	Odd ratio	95% CI	P-value
Age group, years				
• 20 - <40 (ref)	53	1.00	-	-
• 40 - < 60	871	1.37	(0.16, 11.95)	0.78
• > =60	1050	3.80	(0.45, 32.15)	0.22
Gender				
• Male (ref)	1339	1.00	-	-
• Female	635	0.98	(0.56, 1.71)	0.93
Ethnic group*				
• Malay	901	1.56	(0.44, 5.51)	0.49
• Chinese	485	0.91	(0.24, 3.39)	0.89
• Indian	513	0.88	(0.23, 3.36)	0.85
• Others (ref)	75	1.00	-	-
Killip classification code				
• I (ref)	1247	1.00	-	-
• II	304	2.27	(1.27, 4.04)	0.01
• III	91	4.56	(2.16, 9.60)	<0.001
• IV	34	11.74	(4.81, 28.63)	<0.001
• Not stated/ inadequately described	298	0.78	(0.36, 1.67)	0.52
Percutaneous coronary intervention				
• Yes	242	0.64	(0.26, 1.63)	0.36
• No (ref)	1732	1.00	-	-
Cardiac catheterization				
• Yes	357	1.84	(0.88, 3.85)	0.10
• No (ref)	1617	1.00	-	-
TIMI risk score				
• 0-2 (ref)	1137	1.00	-	-
• 3-4	689	0.72	(0.43, 1.19)	0.20
• 5-7	148	1.94	(0.88, 4.27)	0.10

Factors	N	Odd ratio	95% CI	P-value
Smoking				
• Never (ref)	951	1.00	-	-
• Former (quit >30 days)	533	3.30	(1.52, 7.16)	0.002
• Current (any tobacco use within last 30 days)	414	2.46	(1.03, 5.89)	0.04
• Unknown	76	2.79	(1.09, 7.12)	0.03
Family history of premature cardiovascular disease				
• Yes	236	1.11	(0.37, 3.33)	0.86
• No (ref)	939	1.00	-	-
• Unknown	799	1.30	(0.74, 2.30)	0.36
Dyslipidaemia				
• Yes	852	1.22	(0.53, 2.79)	0.64
• No (ref)	443	1.00	-	-
• Unknown	679	0.64	(0.32, 1.26)	0.19
Hypertension				
• Yes	1401	1.57	(0.70, 3.53)	0.28
• No (ref)	352	1.00	-	-
• Unknown	221	0.37	(0.14, 1.01)	0.05
Diabetes				
• Yes	971	3.03	(1.40, 6.55)	0.01
• No (ref)	686	1.00	-	-
• Unknown	317	2.39	(1.02, 5.60)	0.05
Heart failure				
• Yes	236	2.15	(1.18, 3.91)	0.01
• No (ref)	1278	1.00	-	-
• Unknown	460	1.68	(0.81, 3.52)	0.17
Coronary artery disease**				
• Yes	1419	1.72	(0.82, 3.61)	0.15
• No (ref)	331	1.00	-	-
• Unknown	224	1.79	(0.66, 4.84)	0.25

*Others include Orang asli, Kadazan, Melanau, Murut, Bajau, Bidayuh, Iban, other Malaysian and foreigner.

**Coronary artery disease is defined as "Yes" on any of the following co-morbidities: 1) History of myocardial infarction, 2) Documented CAD >50% stenosis, 3) Chronic angina (onset more than 2 weeks ago), 4) New onset angina (less than 2 weeks).

Table 5.6.3 Prognostic factors for death in 30 days among STEMI patients, Malaysia 2006

Factors	N	Odd ratio	95% CI	P-value
Age group, years				
• 20 - <40 (ref)	88	1.00	-	-
• 40 - < 60	581	1.18	(0.39, 3.59)	0.78
• > =60	428	1.88	(0.61, 5.84)	0.28
Gender				
• Male (ref)	927	1.00	-	-
• Female	170	1.45	(0.78, 2.70)	0.24
Ethnic group*				
• Malay	588	1.35	(0.51, 3.58)	0.54
• Chinese	246	0.93	(0.33, 2.60)	0.88
• Indian	209	1.12	(0.39, 3.18)	0.83
• Others (ref)	54	1.00	-	-
Killip classification code				
• I (ref)	707	1.00	-	-
• II	223	1.40	(0.84, 2.34)	0.20
• III	49	2.64	(1.20, 5.79)	0.02
• IV	53	6.64	(3.15, 14.00)	<0.001
• Not stated/ inadequately described	65	0.78	(0.30, 2.06)	0.63
Percutaneous coronary intervention				
• Yes	276	0.66	(0.31, 1.42)	0.29
• No (ref)	821	1.00	-	-
Cardiac catheterization				
• Yes	270	1.09	(0.50, 2.38)	0.82
• No (ref)	827	1.00	-	-
Fibrinolytic therapy				
• Given	776	0.54	(0.35, 0.84)	0.01
• Not given (ref)	321	1.00	-	-
TIMI risk score				
• 0-2 (ref)	492	1.00	-	-
• 3-4	270	1.61	(0.90, 2.89)	0.11
• 5-7	263	2.26	(1.26, 4.04)	0.01
• >7	72	7.70	(3.62, 16.40)	<0.001
Smoking				
• Never (ref)	314	1.00	-	-
• Former (quit >30 days)	200	5.19	(2.33, 11.56)	<0.001
• Current (any tobacco use within last 30 days)	560	3.18	(1.45, 6.97)	0.01
• Unknown	23	1.74	(0.55, 5.44)	0.34

Factors	N	Odd ratio	95% CI	P-value
Family history of premature cardiovascular disease				
• Yes	127	4.23	(1.76, 10.20)	0.02
• No (ref)	583	1.00	-	-
• Unknown	387	0.96	(0.53, 1.74)	0.89
Dyslipidaemia				
• Yes	225	1.87	(0.81, 4.29)	0.14
• No (ref)	331	1.00	-	-
• Unknown	541	1.11	(0.64, 1.93)	0.71
Hypertension				
• Yes	534	4.92	(2.24, 10.79)	<0.001
• No (ref)	308	1.00	-	-
• Unknown	255	1.11	(0.49, 2.47)	0.81
Diabetes				
• Yes	390	5.62	(2.70, 11.71)	<0.001
• No (ref)	399	1.00	-	-
• Unknown	308	0.95	(0.46, 2.00)	0.90
Heart failure				
• Yes	37	2.12	(0.84, 5.37)	0.11
• No (ref)	780	1.00	-	-
• Unknown	280	1.57	(0.71, 3.46)	0.27
Coronary artery disease**				
• Yes	599	0.86	(0.40, 1.87)	0.71
• No (ref)	258	1.00	-	-
• Unknown	240	1.28	(0.53, 3.10)	0.59

*Others include Orang asli, Kadazan, Melanau, Murut, Bajau, Bidayuh, Iban, other Malaysian and foreigner.

**Coronary artery disease is defined as "Yes" on any of the following co-morbidities: 1) History of myocardial infarction, 2) Documented CAD >50% stenosis, 3) Chronic angina (onset more than 2 weeks ago), 4) New onset angina (less than 2 weeks).

Table 5.6.4 Prognostic factors for death in 30 days among NSTEMI/UA patients, Malaysia 2006

Factors	N	Odd ratio	95% CI	P-value
Age group, years				
• 20 - <40 (ref)	35	1.00	-	-
• 40 - < 60	639	0.65	(0.13, 3.20)	0.60
• > =60	819	1.97	(0.41, 9.52)	0.40
Gender				
• Male (ref)	1028	1.00	-	-
• Female	465	0.97	(0.59, 1.58)	0.89
Ethnic group*				
• Malay	631	1.13	(0.41, 3.12)	0.82
• Chinese	403	0.62	(0.21, 1.79)	0.37
• Indian	400	0.75	(0.26, 2.18)	0.59
• Others (ref)	59	1.00	-	-
Killip classification code				
• I (ref)	967	1.00	-	-
• II	266	1.88	(1.13, 3.11)	0.02
• III	82	3.20	(1.60, 6.40)	0.001
• IV	28	6.96	(2.82, 17.21)	<0.001
• Not stated/ inadequately described	150	1.07	(0.55, 2.06)	0.84
Percutaneous coronary intervention				
• Yes	219	0.64	(0.29, 1.43)	0.28
• No (ref)	1274	1.00	-	-
Cardiac catheterization				
• Yes	334	1.43	(0.75, 2.70)	0.28
• No (ref)	1159	1.00	-	-
TIMI risk score				
• 0-2 (ref)	806	1.00	-	-
• 3-4	554	0.76	(0.48, 1.18)	0.22
• 5-7	133	1.53	(0.76, 3.05)	0.23
Smoking				
• Never (ref)	714	1.00	-	-
• Former (quit >30 days)	428	2.43	(1.25, 4.72)	0.01
• Current (any tobacco use within last 30 days)	298	1.84	(0.85, 3.926)	0.12
• Unknown	53	1.63	(0.65, 4.12)	0.30

Factors	N	Odd ratio	95% CI	P-value
Family history of premature cardiovascular disease				
• Yes	176	1.29	(0.51, 3.31)	0.59
• No (ref)	699	1.00	-	-
• Unknown	618	1.28	(0.77, 2.12)	0.34
Dyslipidaemia				
• Yes	688	0.95	(0.47, 1.93)	0.90
• No (ref)	297	1.00	-	-
• Unknown	508	0.57	(0.31, 1.05)	0.07
Hypertension				
• Yes	1062	1.38	(0.67, 2.86)	0.38
• No (ref)	265	1.00	-	-
• Unknown	166	0.51	(0.20, 1.26)	0.14
Diabetes				
• Yes	763	3.12	(1.58, 6.16)	0.001
• No (ref)	496	1.00	-	-
• Unknown	234	1.70	(0.78, 3.71)	0.18
Heart failure				
• Yes	195	1.70	(0.98, 2.96)	0.06
• No (ref)	955	1.00	-	-
• Unknown	343	1.95	(1.02, 3.71)	0.04
Coronary artery disease**				
• Yes	1089	1.78	(0.92, 3.46)	0.09
• No (ref)	247	1.00	-	-
• Unknown	157	1.64	(0.67, 4.01)	0.28

*Others include Orang asli, Kadazan, Melanau, Murut, Bajau, Bidayuh, Iban, other Malaysian and foreigner.

**Coronary artery disease is defined as "Yes" on any of the following co-morbidities: 1) History of myocardial infarction, 2) Documented CAD >50% stenosis, 3) Chronic angina (onset more than 2 weeks ago), 4) New onset angina (less than 2 weeks).

APPENDIX A: DATA MANAGEMENT

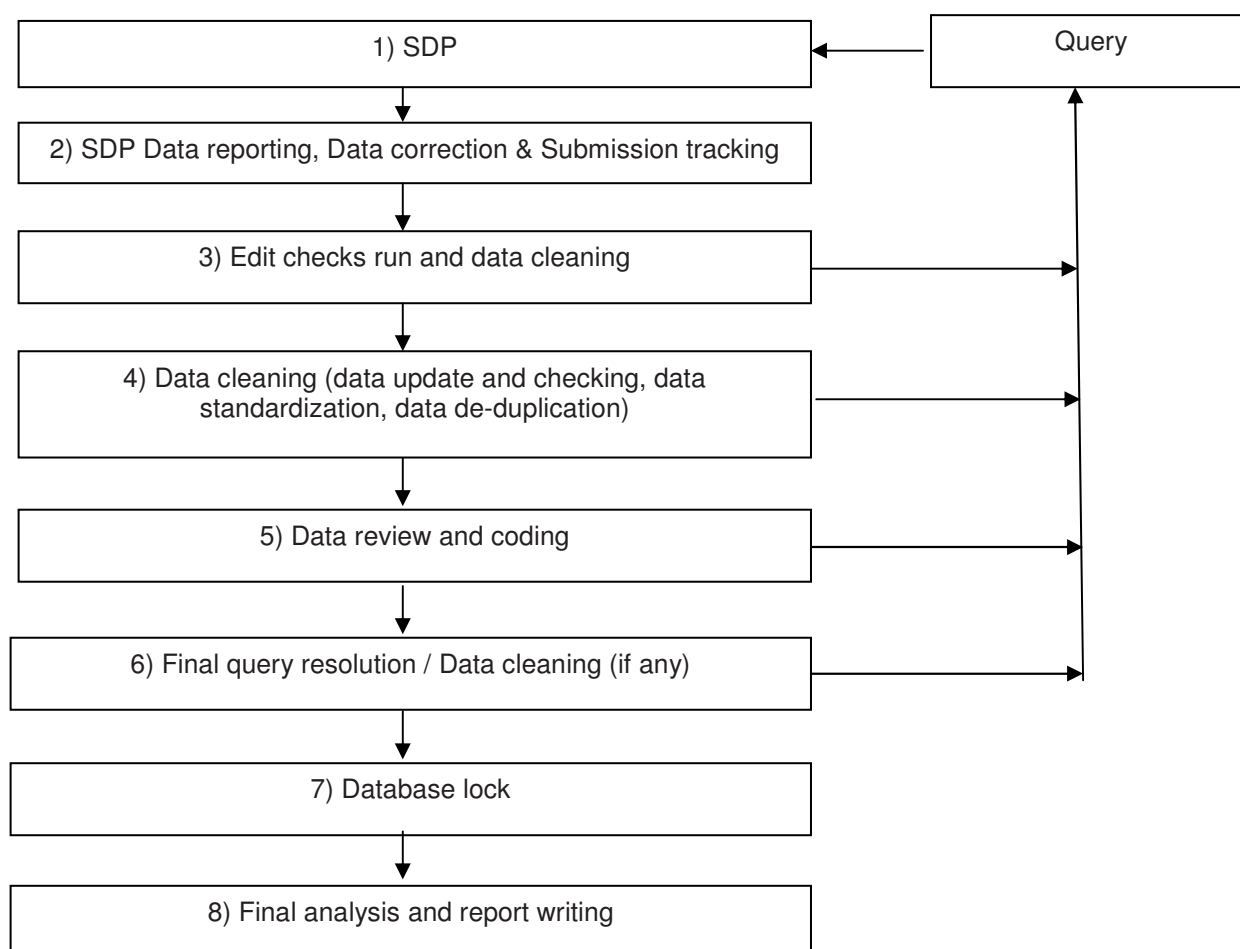
The National Cardiovascular Disease Database (NCVD) Registry maintains two different databases for cardiovascular disease, i.e. for Acute Coronary Syndrome and Percutaneous Coronary Intervention. Data is stored in SQL Server due to the high volume of data accumulated throughout the years.

Data sources

SDPs or Source Data Providers of NCVD-ACS registry comprise of hospitals with cardiologists and physicians who participated in the registry throughout Malaysia.

Data Flow Process

This section describes the data management flow process of the National Cardiovascular Disease Database Registry.



SDP Data reporting, Data correction and Submission tracking

Data reporting by SDP is done via Web Applications e-Case Report Forms.

There are a number of data security features that are designed into NCVD web application (eCRF) such as web owner authentication, 2-level user authentication (user name and password authentication and a Short Messaging System (SMS) of authorization code to mobile phone authentication), access control, data encryption, session management to automatically log off the application, audit trail and data backup and disaster recovery plan.

SDP submits NCVD-ACS Notification form on ad hoc basis whenever there is a case. SDP also submits follow-up data at 30 days, 1 year and other ad hoc follow-ups post notification date. An alert page containing all the overdue submissions for follow up at 30-day and 1-year post notification date is available to users to ease submissions tracking.

Prior to registering a patient record, a verification process is done by using the search function to identify if the patient exists in the entire registry. The application will still detect a duplicate record if the same IC number is keyed in should the step of searching patient be left out. This step is done to avoid duplicate of records. For patients that exist in the database, SDP merely needs to add a new ACS or PCI notification with basic patient particulars pre-filled based on existing patient information in the database. ACS and PCI share the same patient list.

There are a few in-built functionalities at the data entry page that serve to improve data quality. One such function is the auto calculation which is to reduce error of human calculation. There is also a function for inconsistency check that disables certain fields if these fields are answered in a certain manner. When value entered is out of range, user is prompted for the correct value.

A real-time data query page is also available via the web application to enable user to check which of the non-compulsory data is missing, out of range and inconsistent. A link is provided on the data query page for the user to click on to resolve the query for the particular patient.

Real-time reports are also provided in the web application. The aggregated data reports are presented as tables and graphs. The aggregated data reports are typically presented in two manners, the first as centre's own data aggregated data report and second as registry's overall aggregated data report. This way, the centre is able to be compared against the overall registry's average.

Data download function is also available in the web application to allow users to download their own centre's data for all the forms entered for their own further analysis. The data are downloadable as Text - tab delimited (.txt) format, Microsoft excel workbook (.xls) and Comma separated value (.csv) format.

Edit checks run and Data cleaning

Edit check was performed periodically by the registry manager to identify missing compulsory data, out of range values, inconsistency data, invalid values and error with de-duplication. Data cleaning is then performed based on the results of edit checks. Data update and data checking of the dataset are performed when there is a query of certain fields whenever necessary. It could be due to request by user, correction of data based on checking from data query in eCRF or after receiving results for preliminary data analysis. During data standardization, missing data are handled based on derivation from existing data. Data de-duplication is also performed to identify duplicate records in the database that might have been missed by SDP.

Final query resolution / Data cleaning / Database lock

A final edit check run was performed to ensure that data is clean. All queries were resolved before database is locked to ensure data quality and integrity. Final dataset is subsequently locked and exported to the statistician for analysis.

Data analysis

Please refer to section on Statistical Methods section for further details.

Data release policy

One of the primary objectives of the Registry is to make data available to the cardiovascular healthcare providers, policy makers and researchers. The Registry would appreciate that users acknowledge the Registry for the use of the data. Any request for data that requires a computer run must be made in writing (by e-mail, fax, or registered mail) accompanied with a Data Release Application Form and signed Data Release Agreement Form. These requests need prior approval by the Advisory Board before data can be released.

Registry ICT Infrastructure and Data centre

The operations of the NCVD are supported by an extensive ICT infrastructure to ensure operational efficiency and effectiveness.

NCVD subscribes to co-location service with a high availability and highly secured Internet Data Centre at Cyberjaya in order to provide NCVD with quality assured Internet Hosting services and state-of-the-art physical and logical security features without having to invest in costly internal data centre setup. Physical security features implemented includes state of the art security features such as anti-static raised flooring, fire protection with smoke and heat alarm warning system, biometric security access, video camera surveillance system, uninterrupted power supply, environmental control, etc.

Other managed security services include patch management of the servers, antivirus signature monitoring and update, firewall traffic monitoring and intrusion detection, security incidence response, daily, weekly and monthly basis data backup service, at least once yearly data recovery simulation to verify that backup works, half-yearly network security scan and penetration test, security policy maintenance, maintenance and monitoring of audit trail of user access, etc. Managed system services are also provided and these include usage and performance report, operating system maintenance and monitoring, bandwidth monitoring and systems health monitoring.

APPENDIX B: STATISTICAL METHODS

The statistical methods described were used to summarize the data collected from the National Cardiovascular Database (NCVD). In this report, two sources of data have been used for analysis. They were the centre survey data and the NCVD ACS registry data.

Provision of acute coronary care services in Malaysia

Chapter 1 of this report was based on the centre survey data rather than individual patient data reported to the database. This was to provide up-to-date information on patient and centre census in the country and thus overcome the inevitable time lag between processing individual patient data and subsequent reporting of results. The survey was conducted from November 15th 2007 to March 5th 2008. 73 out of 273 hospitals that approached through telephone survey were confirmed with availability of CCU services. Only 69 centres managed to return the survey form completely. Standard error estimates were not reported because no sample was taken. Results on distribution for Malaysia as a whole and also by state were expressed in per million-population since states obviously vary in their population sizes. State population data were based on the last census projection obtained from the Department of Statistics in Malaysia¹. Missing data on ASC services, admissions, utilization of acute coronary or cardiac services and cardiac care are estimated based on the sampling weight of the total beds in each hospital.

The analyses for the rest of this report were generated based on the NCVD ACS registry data, using the following analysis set:

The data without missing on initial diagnosis, final diagnosis is neither stable angina nor non-cardiac, and age at least 20 that were collected until 31st December 2006 by NCVD-ACS were analyzed. The data was stratified to reflect differences in

- Demography: race, gender, age
- Medical factors: premorbid or past medical history
- Initial diagnoses: ACS stratums
- Therapy: fibrinolytic given, aspirin use

Methods for handling missing data and outliers

Missing age was imputed using the hotdeck method. The outliers were set to missing (see table below)

Fields	Acceptable range
Number of distinct episodes of angina	≤20
Heart rate	25 – 200 beats/min (should not be 0)
Systolic BP	60 – 230 mmHg (should not be 0)
Diastolic BP	10 – 120 mmHg
Height	130cm – 210cm
Weight	30kg - 200kg
Waist circumference	≥ 36cm
Hip circumference	60 - 200cm
Peak CK-MB	<1000 Unit/L (should not be 0)
Peak CK	<10 000 Unit/L (should not be 0)
Peak Troponin – TnT	No range
Peak Troponin – Tnl	No range
TC	3 - 20 mmol/L
LDL	1 - 15 mmol/L
HDL-C	0.5 - 5 mmol/L
Tg	1 -15 mmol/L
Fasting Blood Glucose	3 - 30 mmol/L
Left Ventricular Ejection Fraction	5% - 80%
Onset to door	Should not be 0
Door to needle time (mins)	1 min -24 hours (or equivalent minutes) (should not be 0)
Door to balloon time (mins)	1 min-24 hours (or equivalent minutes) Apply only for patients with STEMI and planned for primary angioplasty (should not be 0)

Patient Characteristics

The information on patient characteristics was summarized in chapter 2 of the report. These tables included patients' age, gender, ethnic group, coronary risk factors, anthropometric measurements, co-morbidity, and also the distribution of patients by source data providers (SDP). For summarizing continuous data, the mean, standard deviation, median, minimum and maximum were reported. On the other hand, both the frequency count and percentage were reported for discrete data. Invariably, there were situations where there was missing data. For the purpose of analysis, subjects with missing age had their values imputed by using a hotdeck imputation method. For discrete data, analysis was confined to available data and no imputation was done.

Cardiac Presentation

Chapter 3 of the report basically was to summarize the patient characteristics, vital sign measurements, and laboratory parameters by ACS stratum such as STEMI, NSTEMI and UA, age groups namely young, middle-age and elderly, gender as well as the pre-morbid conditions such as diabetes, hypertension, and dyslipidaemia. For continuous data, the mean, standard deviation, median, minimum and maximum were reported. On the other hand, frequency count and percentage were reported for discrete data. Only the missing age was imputed for the purpose of analysis.

Treatment

The treatments that were provided to the patients were mainly summarized in chapter 4 of the report. This information was cross tabulated by ACS stratum, age group, gender as well as the main ethnic group in Malaysia. No imputation was done for this chapter.

Clinical Outcomes

Chapter 5 of the report summarized the overall in-hospital as well as 30-day outcomes for patients with ACS. Cross tabulations of outcomes by gender, pre-morbid conditions such as diabetes, hypertension, dyslipidaemia, and ACS stratum were included in this chapter. Tabulation of outcomes by fibrinolytic therapy was only presented for STEMI patients. Other tabulations such as outcomes by percutaneous coronary intervention at admission, CABG at admission, and also the pre-admission aspirin use were presented separately for patients with STEMI and NSTEMI/UA. Prognostic factors for in-hospital death as well as death in 30 days were summarized separately for STEMI and NSTEMI/UA patients. No imputation was done for this chapter.

Reference

1. Department of Statistics. Yearbook of Statistics.2002. Malaysia

APPENDIX C: PARTICIPATING CENTRE DIRECTORY

Ipoh Hospital

c/o Medical Department,
Jalan Hospital,
30450 Ipoh,
Perak.

Tel: 05-253 3333
Fax: 05-253 1541
Investigator:
Dr Khor Chiew Gek

Study coordinator:
Sr Zainab Yeop Ahmad

Malacca Hospital

c/o Medical Department,
Jalan Mufti Haji Khalil,
75400 Melaka,

Tel: 06-282 2344
Fax: 06-284 1590
Investigator:
Dr Sharifah Omar

Study coordinator:
S/N Faridah Sharif

Penang Hospital

c/o Cardiology Department,
Jalan Pahang,
50586 Kuala Lumpur.

Tel: 03-2615 6150
Fax: 03-2692 5920
Investigators:
Dato' Dr Omar Ismail
Dr Liew Chee Tat

Study coordinators:
S/N Chong Hooi Joo
S/N Norhafiza Abdul Aziz

Kuala Lumpur Hospital

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50586 Kuala Lumpur.

Tel: 03-2615 6150
Fax: 03-2692 5920
Investigator:
Dato' Dr Jeyaindran Sinnadurai

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S/N Thavamalar Rajanayagam
S/N Sarah Md Yusof

National Heart Institute

c/o Cardiology Department,
145, Jalan Tun Razak,
50400 Kuala Lumpur.

Tel: 03-2617 8200
Fax: 03-2698 2824
Investigator:
Dato' Dr Azhari Rosman

Study coordinators:
Nadiyah Sulaiman
Intan Safarinaz Sabian

Queen Elizabeth Hospital

c/o Medical Department,
Beg Berkunci 2029,
88586 Kota Kinabalu,
Sabah.

Tel: 088-218 166
Fax: 088-318 605
Investigator:
Dr Phanindtranath Mahadasa

Study coordinators:
Sr Wong Kath Koau
Sr Lawasa Mojimbal

Raja Perempuan Zainab II Hospital

c/o Medical Department,
Jalan Hospital,
15000 Kota Bharu
Kelantan.

Tel: 09-745 2000
Fax: 09-747 9532
Investigator:
Dr Monniaty Mohamed

Study coordinators:
S/N Zuryanawati Ahmad Sekeri
S/N Wan Ruzita Wan Hassan

Sarawak General Hospital

c/o Medical Department,
Jalan Tun Ahmad Zaidi Adruce,
93586 Kuching,
Sarawak.

Tel: 082-276 666
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Investigators:
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Dr Kalwinder S Khaira

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S/N Mary Kempo Tahak

Sultanah Aminah Hospital

c/o Cardiology Department,
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80100 Johor Bahru,
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Dr Lu Hou Tee

Study coordinators:
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S/N Delailah Ithnin

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Jalan Tun Ahmad Zaidi Adruce,
93586 Kuching,
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Tel: 082-276 666
Fax: 082-278 308
Investigators:
Prof. Dr Sim Kui-Hian
Dr Ang Choon Kiat

Study coordinators:
Sr Ngu Ching Huong
S/N Cenderella Nuah

Seberang Jaya Hospital

c/o Medical Department,
13700 Perai,
Seberang Jaya,
Pulau Pinang.

Tel: 04-382 7333
Fax: 04-397 0754
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Study coordinator:
Sr Hasmah Salleh

Sultanah Bahiyah Hospital

c/o Medical Department,
KM 6, Jalan Langgar,
05460 Alor Setar,
Kedah.

Tel: 04-740 6203
Fax: 04-740 6154
Investigator:
Dr Hasmannizar Abd Manap

Study coordinator:
S/N Zarina Abdul Hamid

Sultanah Nur Zahirah Hospital

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Jalan Sultan Mahmud,
20400 Kuala Terengganu,
Terengganu.

Tel: 09-621 2121
Fax: 09-622 1820
Investigator:
Dr Wan Mohd Razin Wan Hassan

Study coordinator:
Sr Nor Asiah Mohamad

Tengku Ampuan Rahimah Hospital

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Jalan Langat,
41200 Klang,
Selangor.

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Fax: 03-3372 9089
Investigator:
Datin Paduka Dr Santha Kumari

Study coordinator:
S/N Salbiah

Tuanku Ja'afar Hospital

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Jalan Rasah,
70300 Seremban,
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Dr Cham Yee Ling

Study coordinator:
S/N Narizan Zulkifli

Tengku Ampuan Afzan Hospital

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Jalan Tanah Putih,
25100 Kuantan,
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Dr Harris Ngow Abdullah

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S/N Asanah Asbi

Tuanku Fauziah Hospital

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01000 Kangar,
Perlis.

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Investigators:
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Dr Khor Boon Tat

Study coordinators:
Sr Zauriah Abu Bakar
S/N Rusmira Ramli

University Malaya Medical Centre

c/o Medical Department,
Lembah Pantai,
59100 Kuala Lumpur.

Tel: 03-7949 2821
Fax: 03-7949 2611
Investigators:
Prof. Dr Wan Azman Wan Ahmad
Dr Chong Wei Peng

Study coordinators:
Ms Manhaiyun Alwi
Ms Suzanna Hani Hussein

APPENDIX D: CCU SURVEY PARTICIPATION

We extend our appreciation to the following centres that have contributed in the CCU survey for year 2006, conducted from November 2007 to March 2008:

Ampang Hospital

Jalan Mewah Utara, Pandan Mewah,
68000 Ampang,
Selangor

Ampang Puteri Specialist Hospital

1 Jalan Mamanda 9,
Taman Dato' Ahmad Razali,
68000 Ampang,
Selangor

Assunta Hospital

Lot 68 Jalan Templer,
46990 Petaling Jaya,
Selangor

Bintulu Hospital

Jalan Nyabau,
97000 Bintulu,
Sarawak

Bukit Mertajam Hospital

Jalan Kulim,
14000 Bukit Mertajam,
Seberang Perai,
Pulau Pinang

Damansara Specialist Hospital

119, Jalan SS20/10 Damansara Utama,
47400 Petaling Jaya,
Selangor

Gleneagles Intan Medical Centre

282-286 Jalan Ampang,
50450 Kuala Lumpur

Gleneagles Medical Centre

1 Jalan Pangkor ,
10050 Georgetown,
Pulau Pinang

Kuala Lumpur Hospital

Jalan Pahang,
50586 Kuala Lumpur

Kuantan Clinical Diagnostic Centre

A 37, Jalan Dato' Lim Hoe Lek,
25000 Kuantan
Pahang

Ipoh Hospital

Jalan Hospital,
30990 Ipoh,
Perak

Ipoh Specialist Hospital

26 Jalan Raja Dihilir,
30350 Ipoh,
Perak

Island Hospital

308 Macalister Road,
10450 Georgetown,
Pulau Pinang

Johor Specialist Hospital

39-B, Jalan Abdul Samad,
80100 Johor Bahru,
Johor

Kajang Hospital

Jalan Semenyih,
43000 Kajang,
Selangor

Keningau Hospital

KM 5, Jalan Apin-Apin, Keningau,
Peti Surat 11,
89007 Keningau,
Sabah

Kepala Batas Hospital

Jalan Bertam,
13200 Kepala Batas,
Pulau Pinang

Koh Cardiology and Medical Clinic Sdn. Bhd.

Perak Community Specialist Hospital,
No. 277, Jalan Permaisuri Bainun,
30250 Ipoh,
Perak

Normah Medical Specialist Centre

Jalan Tun Abdul Rahman, Petra Jaya,
93050, Kuching
Sarawak

Pantai Ayer Keroh Hospital Sdn. Bhd.

No.2418-1, KM 8 Lebuah Ayer Keroh,
75450 Melaka

Kulim Hospital

Jalan Mahang,
09000 Kulim,
Kedah

Lam Wah Ee Hospital

141, Jalan Tan Sri Teh Ewe Lim,
11600 Georgetown,
Pulau Pinang

Loh Guan Lye Specialist Centre

19 & 21, Jalan Logan,
10400 Georgetown,
Pulau Pinang

Mahkota Medical Centre

No. 3 Mahkota Melaka, Jalan Merdeka,
75000 Melaka

Malacca Hospital

Jalan Mufti Haji Khalil,
75400 Bandar Melaka,
Melaka

Metro Specialist Hospital

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Kedah

National Heart Institute

145 Jalan Tun Razak
50400 Kuala Lumpur

Pusrawi Hospital Sdn. Bhd.

Lot 149, Jalan Tun Razak,
50400 Kuala Lumpur

Puteri Specialist Hospital

33 Jalan Tun Abdul Razak (Susur 5),
80350 Johor Bahru,
Johor

Putra Specialist Hospital (Malacca) Sdn. Bhd.

169, Jalan Bendahara,
75100 Melaka

Putrajaya Hospital

Pusat Pentadbiran Kerajaan Persekutuan,
Presint 7,
62250 Putrajaya

Queen Elizabeth Hospital

Karung Berkunci No 2029,
88586 Kota Kinabalu,
Sabah

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55100 Kuala Lumpur

Pantai Medical Centre

8 Jalan Bukit Pantai,
59100 Kuala Lumpur

Pekan Hospital

26600 Pekan,
Pahang

Penang Adventist Hospital

465, Jalan Burma
10350 Pulau Pinang

Penang Hospital

Jalan Resideni,
10900 Georgetown,
Pulau Pinang

Perdana Specialist Hospital

Suite 19, Lot PT37 & 600,
Seksyen 14, Jalan Bayam,
15200 Kota Bharu,
Kelantan

Port Dickson Hospital

KM 11 Jalan Pantai, Sirusa,
71050 Port Dickson,
Negeri Sembilan

Selangor Medical Centre

Lot 1, Jalan Singa 20/1 Section 20,
40300 Shah Alam
Selangor

Selayang Hospital

Lebuhraya Selayang-Kepong,
68100 Batu Caves,
Selangor

Sentosa Medical Centre

36 Jalan Chemur,
Damai Complex,
50400 Kuala Lumpur

Serdang Hospital

Jalan Puchong,
43000 Kajang,
Selangor

Seremban Specialist Hospital

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Sabah Medical Centre

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88300 Kota Kinabalu,
Sabah

Sarawak General Hospital

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Sarawak

Seberang Jaya Hospital

Jalan Tun Hussein Onn,
13700 Seberang Perai,
Pulau Pinang

Subang Jaya Medical Centre

1 Jalan SS12/1A,
47500 Subang Jaya,
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Sultan Abdul Halim Hospital

Jalan Lencongan Timur,
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Sultan Haji Ahmad Shah Hospital

28000 Temerloh,
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Sultan Ismail Hospital

Jalan Persiaran Mutiara Emas Utama,
Taman Mount. Austin,
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Sultanah Aminah Hospital

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Sultanah Bahiyah Hospital

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Sibu Hospital

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Slim River Hospital

35800 Slim River,
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Sri Kota Specialist Medical Centre

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Taiping Hospital

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Tawakal Specialist Centre

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Tuanku Ampuan Najihah Hospital

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APPENDIX E: NOTE OF APPRECIATION

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APPENDIX F: DATA DEFINITIONS

ACS stratum	Risk stratum of the patient presenting with clinical features consistent with an ACS (chest pain or overwhelming shortness of breath), defined by accompanying clinical, ECG and biochemical features
Bleeding complication (TIMI criteria)	<p>The person's episode of bleeding as described by the thrombolysis in myocardial infarction (TIMI) criteria:</p> <p>Major; Overt clinical bleeding (or documented intracranial or retroperitoneal hemorrhage) associated with a drop in hemoglobin of greater than 5 g/dl (0.5 g/l) or in hematocrit of greater than 15% (absolute)</p> <p>Minor; Overt clinical bleeding associated with a fall in hemoglobin of 3 to less than or equal to 5g/dl (0.5 g/l) or in hematocrit of 9% to less than or equal to 15% (absolute)</p> <p>None; No bleeding event that meets the major or minor definition</p>
Blood pressure (diastolic) at presentation	The person's measured diastolic blood pressure (at presentation)
Blood pressure (systolic) at presentation	The person's measured systolic blood pressure (at presentation)
Cerebrovascular disease	Indicates if the patient has a history of stroke and/or transient ischaemic attack (TIA) or documented evidence of cerebrovascular disease (CT scan, MRI) prior to this hospital admission
Chronic angina (onset more than 2 weeks ago)	Indicates if the patient has an angina for more than 2 weeks prior to this hospital admission
Chronic lung disease	Indicates if the patient has a history of chronic lung disease including chronic obstructive pulmonary disease (COPD), chronic pulmonary fibrosis, cystic fibrosis or bronchiectasis, or is receiving treatments for these conditions, prior to this hospital admission. Previous acute pneumonia and ventilation for acute respiratory distress are excluded
Chronic renal disease	Indicates if the patient has documented underlying moderate to severe impairment of renal function prior to this hospital admission
'Culprit' artery	The vessel considered as responsible for the Acute Coronary Syndrome
Current smoker	Patient regularly smokes tobacco product / products one or more times per day or has smoked in the 30 days prior to this admission
Date of onset ACS symptoms	Defines the date of onset of index event resulting in admission.
Date patient presented	Defines the date that patient presented to the reporting centre.

Diabetes mellitus	Indicates if the patient has a history of diabetes mellitus diagnosed prior to this hospital admission or currently receiving treatment for diabetes. Defines TIMI UA/NSTEMI score. Defines TIMI STEMI score
Documented CAD > 50% stenosis	Indicates if the patient has angiographically-proven coronary disease involving at least 1 vessel with greater than 50% stenosis, or have undergone percutaneous angioplasty (PCI) or coronary artery bypass graft (CABG) prior to this hospital admission. Defines TIMI UA/NSTEMI score
Door to balloon time (mins)	The duration between time patients presented to the reporting centre to time of first angioplasty balloon inflation/stenting by the same centre. Applicable only to patients with STEMI undergoing urgent PCI.
Door to needle time	The duration between time patients presented to the reporting centre to time intravenous fibrinolytic therapy was administered or initiated by that same centre. Applicable only to STEMI patients receiving thrombolysis at the reporting centre
Dyslipidaemia	Indicates if the patient has a history of dyslipidaemia diagnosed prior to this hospital admission or currently receiving treatment for dyslipidaemia. Defines TIMI UA/NSTEMI score
ECG abnormalities location	Describes the area in which the main abnormalities are located on the standard 12-lead ECG
ECG abnormalities type	Describes the type of abnormalities seen on the ECG
Facilitated PCI	PCI may be performed as part of planned revascularization strategy in conjunction with pharmacological thrombolysis
Family history of premature cardiovascular disease	Indicates if the patient has a 1st degree family member (parents or siblings) who suffered a myocardial infarction and/or stroke before the age of 55 years. Defines TIMI UA/NSTEMI score
Fasting blood glucose	A method for learning how much glucose there is in a blood sample taken after an overnight fast (in mmol/L)
Fibrinolytic drug used	Identifies the fibrinolytic drugs used. Applicable only to patients presenting with STEMI. This data may be entered by the reporting centre for patients who received thrombolysis prior to transfer
Fibrinolytic therapy status	Identifies the person's fibrinolytic therapy status. Applicable only to patients presenting with STEMI. This data may be entered by the reporting centre for patients who received thrombolysis prior to transfer. Also indicates whether and where thrombolysis was given. Only thrombolysis instituted by the participating centre will be calculated for 'Door-to-Needle' time
Final diagnosis at discharge	Indicates one of the following the final diagnosis after all procedures and investigations had been performed <ul style="list-style-type: none"> ▪ Q-wave MI, or ▪ Non Q-wave MI, or ▪ Unstable angina, or ▪ Stable angina, or ▪ Non cardiac

Former smoker	Patient has stopped smoking tobacco products more than 30 days before this admission
HDL-C	The person's latest measured high-density lipoprotein cholesterol (HDL-C) latest level before event (in mmol/L)
Heart failure	Indicates if the patient has a history of heart failure or documented evidence (echocardiography, MRI, nuclear imaging, ventriculography) of left ventricular systolic dysfunction prior to this hospital admission
Heart rate at presentation	The heart rate recorded in beats per minute (at presentation)
Height (cm)	Measures the patient's height in cm. Measurements may be taken at any time prior to discharge. However, measurements taken after prolonged hospitalization (>2 weeks) or following surgery or prolonged intensive unit stay may not be accurate
Hip circumference (cm)	Measures the patient's hip circumference at presentation. Measurements may be taken at any time prior to discharge. However measurements taken after prolonged hospitalization (>2 weeks) or following surgery or prolonged intensive unit stay may not be accurate
Hypertension	Indicates if the patient has a history of hypertension diagnosed prior to this hospital admission or currently receiving treatment for hypertension. Defines TIMI UA/NSTEMI score. Defines TIMI STEMI score
Intravenous Fibrinolytic therapy (date)	The date intravenous fibrinolytic therapy was administered or initiated. Applicable only to patients presenting with STEMI. This data may be entered by the reporting centre for patients who received thrombolysis prior to transfer
Intravenous Fibrinolytic therapy (time)	The time intravenous fibrinolytic therapy was administered or initiated. Applicable only to patients presenting with STEMI. This data may be entered by the reporting centre for patients who received thrombolysis prior to transfer
Killip classification	Identifies the Killip class, as a measure of haemodynamics compromise, of the person at the time of presentation Class I includes individuals with no clinical signs of heart failure Class II includes individuals with rales in the lungs, an S3 gallop, and elevated jugular venous pressure Class III describes individuals with frank pulmonary edema Class IV describes individuals in cardiogenic shock
LDL-C	The person's latest measured low-density lipoprotein cholesterol (LDL-C) latest level before event (in mmol/L)
Left Main Stem Involvement	Left main disease (>50%) is counted as TWO-vessel disease (LAD and Circumflex). This field to be entered if there is at least 50% stenosis in the left main stem
Lipid Profile (fasting)	The person's measured lipid profile values
LVEF (%)	Ejection fraction is the percentage of the blood that fills the left ventricle during diastole that is then pumped into the body during systole. It measures the blood-pumping efficiency of the left ventricle

Myocardial infarction history	Indicates if the patient has a history of myocardial infarction prior to this hospital admission. Defines TIMI UA/NSTEMI score
New onset angina (Less than 2 weeks)	Indicates if the patient has an angina in the past 2 weeks prior to this hospital admission
No of days in CCU / ICU / CICU	Total number of days spent in a cardiac care unit (CCU) at the reporting centre only, either consecutively or intermittently
NSTEMI	Patients with chest pains characteristic of unstable angina AND showing evidence of biochemical myocardial necrosis. For patients who have received thrombolysis prior to admission to reporting centre, NSTEMI may also be diagnosed if after early reperfusion there may be rapid resolution of existing ST elevation associated with CK rise <2xULN or small Trop rise only
Number of diseased vessels	The number of major coronary vessels systems (LAD system, Circumflex system, and / or Right System) with > 50% narrowing in any angiographic view, or significant lesion defined by IVUS or pressure wire assessment
Number of distinct episodes of angina in past 24 hours	Number of distinct episodes of anginal pain that occurred in the 24 hours before hospital presentation
Peak CK	The person's maximum measured CK level over 48 hours from the time of presentation (in Unit/L)
Peak CK-MB	The person's maximum measured CK-MB isoenzyme over 48 hours from the time of presentation (in Unit/L)
Peak Troponin T / I	The person's maximum measured troponin (T / I / or both) over 48 hours from the time of presentation (in microgram/L) or state as positive or negative
Peripheral vascular disease	Indicates if the patient has a history and/or documented evidence and/or have undergone treatment for peripheral vascular disease (including aortic aneurysm; peripheral artery disease, intermittent claudication and/or previous peripheral artery stenting or bypass; renal artery stenosis and/or previous renal artery stenting)
Primary PCI	PCI intended as the primary mode of coronary revascularization
Rescue PCI	After initial thrombolysis, PCI may be performed when there is on-going cardiac ischemia or perceived failure of thrombolytic drug to achieve adequate reperfusion
Smoking status	Indicates if the patient has a history confirming any form of tobacco use in the past. This includes use of cigarettes / cigars / pipes/ tobacco chewing
STEMI	History consistent with diagnosis plus ST elevation in contiguous leads or new LBBB
Time of onset ACS symptoms	Defines the time of onset of index event resulting in admission using the 24-hour clock
Time patient presented	Defines the time that patient presented to the reporting centre

TIMI flow classification	<p>Angiographic criteria of severity of coronary flow impediment prior to PCI as defined by the TIMI score.</p> <ul style="list-style-type: none"> 0 No perfusion; I Penetration without perfusion; II Partial perfusion. Contrast opacifies the entire coronary bed distal to the stenosis. However the rate of entry and/or clearance is slower in the coronary bed distal to the obstruction than in comparable areas not perfused by the vessel; III Complete perfusion. Filling and clearance of contrast equally rapid in the coronary bed distal to stenosis as in other coronary beds
TIMI Risk score for STEMI	<p>Thrombolysis in myocardial infarction (TIMI) risk score for STEMI is based on the following criteria: (Max 14 points)</p> <ul style="list-style-type: none"> a) Age \geq 75 - 3 point, Age 65 to 74 - 2 points b) Diabetes OR Hypertension OR Chronic angina (onset more than 2 weeks ago) OR New onset angina (Less than 2 weeks) - 1 point c) Systolic BP $<$100 mmHg - 3 points d) Heart Rate $>$ 100 beat per minute - 2 points e) Killip II-IV - 2 points f) Weight $<$ 67 kg - 1 point g) Anterior Leads: V1 to V4, Bundle Branch block (BBB) - 1 point h) Time to Treatment $>$4 - 1 point
TIMI Risk score for UA/NSTEMI	<p>Thrombolysis in myocardial infarction (TIMI) risk score for UA/NSTEMI is based on the following criteria: 1 point for each of the following criteria</p> <ul style="list-style-type: none"> a) Age \geq 65 b) At least 3 risk factors (Past medical history: dyslipidaemia, HPT, diabetes, premature cardiovascular disease family history status) c) Known CAD (stenosis \geq 50%) (Past medical history: Myocardial infarction history, Documented CAD $>$50% stenosis) d) ST (ECG) deviation \geq 0.5mm (ECG Abnormalities Type: ST-segment elevation \geq 1mm (0.1 mV) in \geq 2 contiguous leads, ST-segment elevation \geq 2mm (0.2 mV) in \geq 2 contiguous leads, ST-segment depression \geq 0.5mm (0.05 mV) in \geq 2 contiguous leads (includes reciprocal changes) e) Recent (\leq 24 hrs) Severe angina (\geq 2 angina in last 24 hrs) f) Use of anti-platelet agent (ASA) in last 7 days (Used at least one of ASA, ADP Antagonist) g) Elevated cardiac enzymes/markers: CK-MB (Peak CKMB value $>$ CKMB Reference Upper limit) and Troponin (Peak Troponin value $>$ Troponin Reference Upper limit)
Total cholesterol	The person's latest measured total cholesterol latest level before event (in mmol/L)
Total days of admission	Total number of days spent at reporting centre from the day of admission till the day of outcome, because of discharge, transfer or patient death

Triglycerides	The person's latest measured triglycerides latest level before event (in mmol/L)
Unstable angina	Angina (or other distinct chest pain patterns) without evidence of biochemical myocardial necrosis BUT with any 1 of the following: (1) Angina occurring at rest and prolonged >20mins; (2) New-onset angina of at least CCS III severity; (3) Recent acceleration of angina by at least 1 CCS class.
Waist circumference (cm)	Measures the patient's waist circumference at presentation. Measurements may be taken at any time prior to discharge. However measurements taken after prolonged hospitalization (>2 weeks) or following surgery or prolonged intensive unit stay may not be accurate
Weight (kg)	Measures the patient's weight in kg. Measurements may be taken at any time prior to discharge. However, measurements taken after prolonged hospitalization (>2 weeks) or following surgery or prolonged intensive unit stay may not be accurate