

# Department of Cardiothoracic Surgery Cardiac Surgery Biennial Report 2010 - 2011



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Message From Chief Executive by Dr P.Y. Leung, Chief Executive, Hospital Authority

Patient safety is always of utmost priority to the Hospital Authority (HA) and it must never be compromised. Yet, we could not be able to fully understand our performance without the support from data.

I am glad to see the release of this biennial Cardiothoracic Surgery Report, the first comprehensive review aiming at a better understanding of the performance in the Queen Mary Hospital. The report provides a systemic approach and structured analysis for us to plan and determine the direction for improving the quality of care for cardiac patients.

I would like to take this opportunity to express my appreciation to our dedicated professionals for their unrelenting efforts in ensuring patient safety and building a culture of continuous improvement. This will enable us to move toward our vision of being trusted by the community. I am confident that we will see greater achievements together in the years to come, and more substantial contributions to patient care, to the HA and to the community.



### Foreword

#### by Professor Chung-Mau LO, Chair Professor and Head of Department of Surgery, The University of Hong Kong

This is the first report on cardiac surgery at Queen Mary Hospital for the year 2010 to 2011 based on data generated by the Dendrite Clinical System. The report clearly shows that Hong Kong has a world-class cardiac surgery service.

Despite a more complex and diverse case mix, the data indicates that the outcomes of cardiac surgery at Queen Mary Hospital are comparable if not superior to those of the United Kingdom as published by the Society of Cardiothoracic Surgeons in Great Britain & Ireland. Of the 867 adult cardiac surgical procedures performed over the two-year period, the most common operation was isolated coronary artery bypass surgery, followed by isolated valve surgery. There were more valvular and re-do valvular operations, aortic operations and transplantation compared to centers in the United Kingdom.

Our crude mortality rate for coronary artery bypass surgery, isolated or combined with other kinds of cardiac surgery, was on a par with that in the United Kingdom and that reported by the European Association for Cardio-Thoracic Surgery. For isolated coronary artery bypass surgery, the ratio of observed mortality to predicted mortality was 0.44 only. The results of valve surgery were even more impressive as our crude mortality of different kinds of valve surgery, isolated or combined, were consistently lower than that in the United Kingdom and Europe. The predicted mortality rate for all single valve surgery was 6.9% and the observed rate was 1.6%, with an observed-to-predicted-mortality ratio of 0.23. For all kinds of valve surgery, the observed mortality rates were lower than the predicted mortality rates.

It is heartening to see that cardiac surgery at Queen Mary Hospital is another clinical practice par excellence. This outstanding achievement could not have been possible without the skill and hard work of a dedicated and committed team of surgeons, nurses, anaesthesiologists, and allied health, of which we are proud.



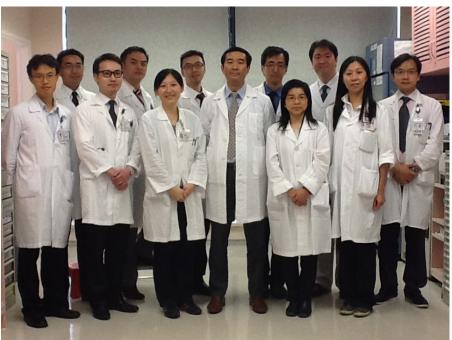
## Introduction

by Dr Timmy Wing Kuk Au, Chief of Service, Department of Cardiothoracic Surgery, Queen Mary Hospital

he Department of Cardiothoracic Surgery is very proud to publish our first biennial adult cardiac surgical report for the year 2010 and 2011. Cardiac surgery has been performed in our department for nearly 50 years but systematic clinical data collection and outcome audit were not carried out at the Grantham Hospital (our predecessor) until the late 1990s. Risk stratification using EuroSCORE protocol was then our main program in the adult cardiac surgery outcome audit at that time. This prospective surgical audit in adult cardiac surgery was fully supported by our previous Chief of Service Dr Chiu Shiu-Wah and Dr Cheng Lik-Cheung. Reports had been published in peer review journals and our data presented in various local and international meetings with good feedback. With the support of the Hong Kong West Cluster, we moved on to adopt the Dendrite Clinical System for our data management in late 2009. Currently, we have a dedicated team of 2 cardiothoracic surgeons and one nursing officer who are responsible for data collection and follow ups. We believe that this report could act as our continuous quality improvement program as well as unveil our practice of adult cardiac surgery to our colleagues in the Hospital Authority and to the public both locally and overseas. Finally, this report would never



Introduction



Cardiothoracic Surgery team photo

have been accomplished without the untiring efforts, skill and passionate countenance of our staffs and their devotion towards excellence in patient care.

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## **Database overview**

6. DE



## The overall workload at QMH, Hong Kong for the calender years 2010 & 2011

#### **Procedure groupings**

- All data comparisons are made to the data from United Kingdom published by the Society for Cardiothoracic Surgeon in Great Britain & Ireland in their Sixth National Adult Cardiac Surgical Database Report 2008, & relate to the most up-to-date data in that document, from the financial year ending 2008.
- We have performed 867 adult cardiac surgical procedures (defined as open heart surgery for patients over the age of 18 years).
- Percentage of our coronary artery bypass graft surgery (CABG) was lower than the United Kingdom data (33% in Queen Mary Hospital *versus* 58% in United Kingdom).
- On the other hand, percentage of our valves surgery was higher than the United Kingdom data (Isolated Valves + Valves & Others is 40% in Queen Mary Hospital *versus* 23% in United Kingdom).
- Adult congenital heart surgery comprised a greater proportion of our workload than reported for the United Kingdom in their Sixth National Adult Cardiac Surgical Database Report 2008.
- Aortic surgery and heart transplantation were also relatively more common in Queen Mary Hospital compared to the data reported by the United Kingdom surgeons.

Overall workload at Queen Mary Hospital, Hong Kong; calendar years 2010 & 2011

		Data		
		Count	Proportion	Proportion in the UK
	Isolated CABG	286	33.0%	58.3%
бu	Isolated valve	240	27.7%	18.9%
grouping	CABG & valve	56	6.5%	11.5%
gro	CABG & other	21	2.4%	2.0%
ure	Valve & other	107	12.3%	4.5%
Procedure	CABG, valve & other	17	2.0%	1.6%
Pro	Other	140	16.1%	3.2%
	All	867		

For details on the other procedures, please see the data on page 16.



#### Procedure detail

- The procedure groupings that contain other in their description (CABG & other; CABG, valve & other; Valve & other; Other) describe operations that involve a cardiac surgical procedure other than coronary artery bypass and valve surgery.
- These other procedures include : atrial septal defect (ASD) repair, ventricular septal defect repair (VSD), aortic surgery, radiofrequency ablation surgery (MAZE) and thoracic resection.
- 3 or more grafts were performed in 257 isolated coronary artery surgery, which was a higher rate than that reported in the United Kingdom data (89% in Queen Mary Hospital *versus* 74% in United Kingdom).
- More detail information on CABG and valve surgery at Queen Mary Hospital can be found in dedicated sections later in this report.

			Procedure groupings							
			Isolated CABG	Isolated valve	CABG & valve	CABG & other	Valve & other	CABG, valve & other	Other	All
	>	1 graft	2	0	25	9	0	12	0	48
	CABG surgery	2 grafts	25	0	15	5	0	3	0	48
_	ins 5	3 grafts	185	0	12	5	0	2	0	204
etai	ABC	4 grafts	69	0	3	2	0	0	0	74
Procedure detail	0	≥5 grafts	3	0	0	0	0	0	0	3
edu	>	Aortic alone	0	64	28	0	19	7	0	118
Proc	surgery	Mitral alone	0	65	14	0	23	0	0	102
	a sur	Aortic & mitral	0	18	1	0	5	2	0	26
	Valve	Mitral & tricuspid	0	35	9	0	22	3	0	69
	>	Other valve combinations	0	52	4	0	29	3	0	88

Procedure detail



#### Other procedure detail

- It is important to remember that the patients may have more than one of the other procedures. For example, there are 3 patients who had both surgery on the aorta **and** congenital surgery.
- The group 'other procedures not listed above' included all those patients for whom there was an other procedure of some kind recorded, but who did not fall into any of the categories listed above; examples of these kinds of procedures would be: myxomas, HOCM myomectomy surgery, lung resections, atrial reduction plasty, concomitant peripheral vascular procedures and TEVAR with cardiopulmonary bypass.

		Data	
		Count	Proportion
	No other procedures	582	67.1%
	All operations with an other component	285	32.9%
	Surgery on the aorta	97	11.2%
S	Cardiac transplant	21	2.4%
Other procedures	Adult congenital surgery	40	4.6%
ocec	Pulmonary embolectomy	4	0.5%
r pr	Pericardiectomy	2	0.2%
the	Epicardial pacemaker	2	0.2%
0	LV aneurysmectomy	2	0.2%
	Radio-frequency ablation	51	5.9%
	Other procedure not listed above	51	5.9%
	All	867	

Other procedures performed



#### Previous cardiac surgery

- All comparisons to data from the United Kingdom come from results published by the Society for Cardiothoracic Surgeon in Great Britain & Ireland in their Sixth National Adult Cardiac Surgical Database Report 2008, & relate to the most up-to-date data in that document, from the financial year ending 2008.
- Isolated CABG with previous cardiac surgery was 2.4% compared to 1.6% in United Kingdom.
- Patients with coronary artery diseases with previous cardiac surgery who then require further coronary intervention may now more frequently undergo PCI rather than surgery and the situation is similar in United Kingdom.
- Isolated valve surgery with previous cardiac surgery was 30% in Queen Mary Hospital while all valves & other surgery with previous cardiac surgery was 26% compared to 9% for isolated AVR with previous cardiac surgery and 2% for isolated MVR with previous cardiac surgery from United Kingdom data.
- Overall 16% of our adult cardiac surgery patients had previous cardiac surgery done.

		Previous cardiac surgery			
		No	Yes	Proportion prior surgery	
	Isolated CABG	279	7	2.4%	
ng	Isolated valve	169	71	29.6%	
grouping	CABG & valve	53	3	5.4%	
-	CABG & other	20	1	4.8%	
Procedure	Valve & other	79	28	26.2%	
ocec	CABG, valve & other	15	2	11.8%	
Pro	Other	112	28	20.0%	
	All	727	140		

Previous surgery



#### Mortality

- In-hospital mortality is used as our primary outcome rather than 30-day mortality.
- Our isolated CABG and CABG combined surgery crude mortality were on par with the United Kingdom data and EACTS database.
- Our isolated valve surgery and valves combined surgery crude mortality was lower than the United Kingdom data and EACTS database.

In-hospital, post-operative mortality rates for each procedure group

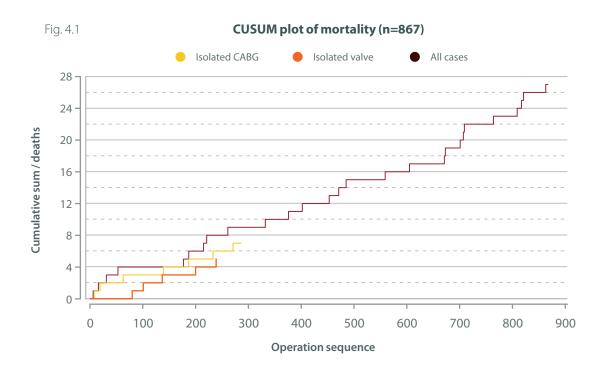
		Mortality			
		Alive	Died	Rate (95% Cl)	
	Isolated CABG	279	7	2.4% (1.1-5.2%)	
ng	Isolated valve	235	5	2.1% (0.8-5.1%)	
grouping	CABG & valve	54	2	3.6% (0.6-13.4%)	
	CABG & other	19	2	9.5% (1.7-31.8%)	
Procedure	Valve & other	104	3	2.8% (0.7-8.6%)	
ocec	CABG, valve & other	17	0	0.0% (0.0-16.2%)	
Pre	Other	132	8	5.7% (2.7-11.3%)	
	All	840	27		

International comparison of in-hospital, post-operative mortality rates for each procedure group

		Mortality data			
		Queen Mary Hospital, Hong Kong	United Kingdom NACSD 2008	EACTS database 2006-2008	
	Isolated CABG	2.4% (286; 1.1-5.2%)	1.5% (22,808; 1.3-1.6%)	2.2% (219,053; 2.2-2.3%)	
_	Isolated valve	2.1% (240; 0.8-5.1%)	3.5% (7,379; 3.1-4.0%)	3.4% (75,247; 3.3-3.5%)	
grouping	CABG & valve	3.6% (56; 0.6-13.4%)	6.1% (4,508; 5.4-6.8%)	6.2% (37,721; 6.0-6.5%)	
	CABG & other	9.5% (21; 1.7-31.8%)	7.8% (766; 6.1-10.0%)	7.0% (4,327; 6.3-7.8%)	
Procedure	Valve & other         2.8%           (107; 0.7-8.6%)		5.5% (1,780; 4.5-6.7%)	4.9% (12,883; 4.5-5.3%)	
	CABG, valve & other	alve & other 0.0% (17; 0.0-16.2%)		11.3% (3,097; 10.2-12.5%)	
	Other	5.7% (140; 2.7-11.3%)	7.9% (1,271; 6.5-9.5%)	7.7% (11,562; 7.2-8.2%)	



- The cumulative sum technique is a method of plotting an accumulation of events over time.
- The cumulative mortality plot provides visual representation of performance against the expected outcome rate of a particular risk scoring protocol.
- When observed CUSUM mortality plot compares with the predicted CUSUM mortality plot allows the detection of trends and corrective actions and it provides an excellent audit to surgeons and hospital administrators.
- There were no indications of odd results in the CUSUM plot for Queen Mary Hospital.





## Isolated CABG surgery

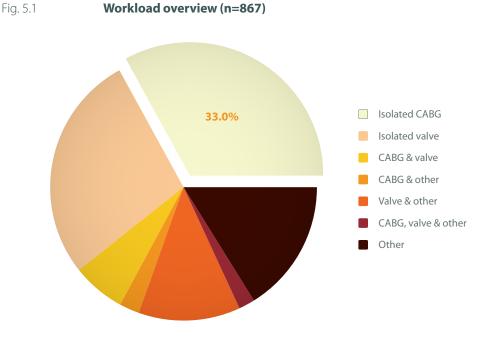


## **Isolated coronary surgery**

#### CABG in the context of overall workload

- Coronary surgery contributed 33% of workload in our department.
- In contrast to western countries that CABG usually attributes 70% of case load.
- Total 286 patients had isolated CABG in the year 2010 to 2011.

Workload overview (n=867)



#### **Pre-operative risk factors**

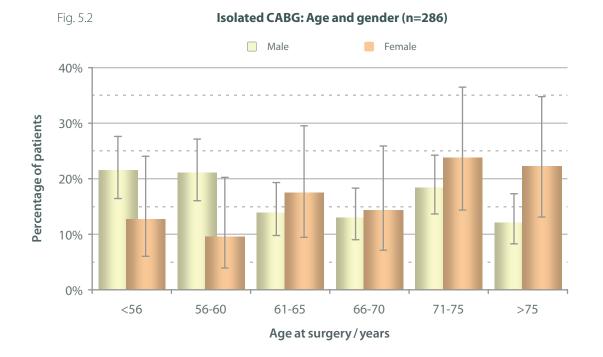
#### Age and gender

#### Age and gender distributions

- 286 patients (223 male and 63 female) had undergone isolated coronary surgery in year 2010 and 2011.
- Patients undergoing coronary artery bypass grafting was dominated by male (78%).
- Overall number was distributed similarly in different age groups. However, male patients tended to present at younger age.

#### Queen Mary Hospital, Hong Kong





Age, gender and mortality

- Crude mortality showed female patients had lower mortality rate, which is different from the *EuroSCORE* where female itself as a risk factor.
- When different age groups are taken into consideration, older men had higher mortality while this trend is not seen in women.

		Gender			
			Male		Female
Ś	<56	0.0%	(0.0-6.1%)	0.0%	(0.0-31.2%)
at surgery / years	56-60	2.1%	(0.1-12.7%)	0.0%	(0.0-39.3%)
ry/}	61-65	0.0%	(0.0-9.2%)	9.1%	(0.5-42.9%)
Irge	66-70	3.4%	(0.2-19.6%)	0.0%	(0.0-28.3%)
itsu	71-75	4.9%	(0.8-17.8%)	0.0%	(0.0-18.1%)
Age a	>75	7.4%	(1.3-25.8%)	0.0%	(0.0-19.3%)
A	All	2.7%	(1.1-6.0%)	1.6%	(0.1-9.7%)

Isolated CABG surgery: crude mortality, age and gender

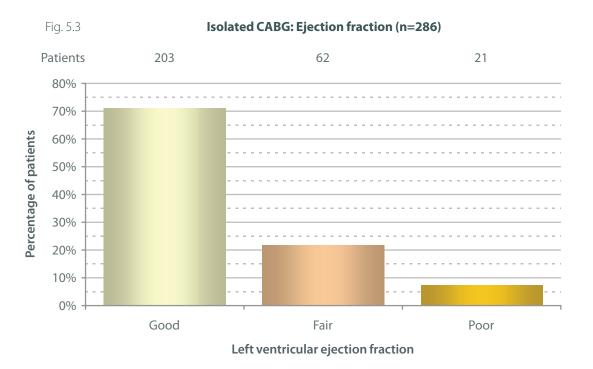
Isolated CABG surgery



#### Left ventricular ejection fraction

Left ventricular ejection fraction distributions

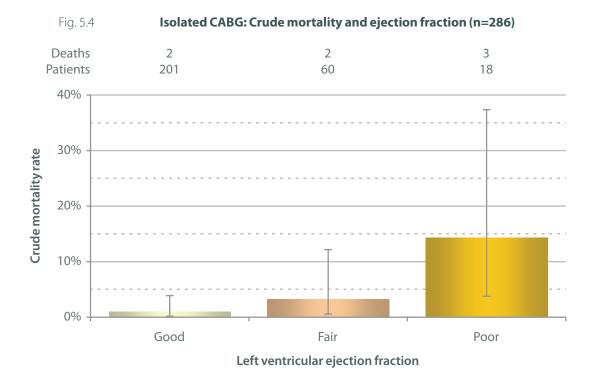
- Ventricular function is mainly assessed by echocardiogram and expressed as ejection fraction.
- Before operation, 21.7% of patients had impaired left ventricular function while 7.3% of them were considered as poor (ejection fraction <30%).</li>
- An intra-aortic balloon pump (IABP) would be inserted before operation if the patient's ejection fraction were poor, or had unstable angina or unstable haemodynamics. 37 of our patients (12.9%) fell into this category.





Left ventricular ejection fraction and mortality

- Poor left ventricular function is a well known risk factor in revascularization surgery.
- Mortality was correlated to left ventricular function in our patient population. There was only 1% mortality in good function group.

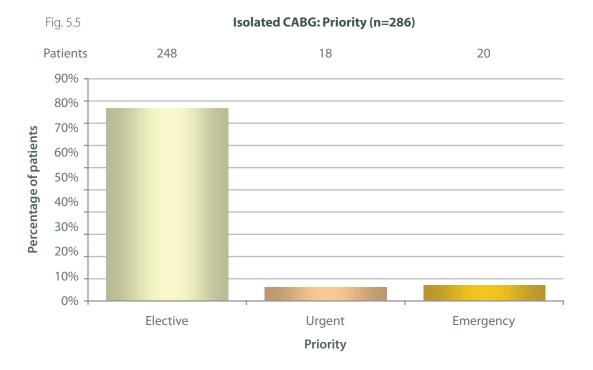




#### Priority

**Priority distributions** 

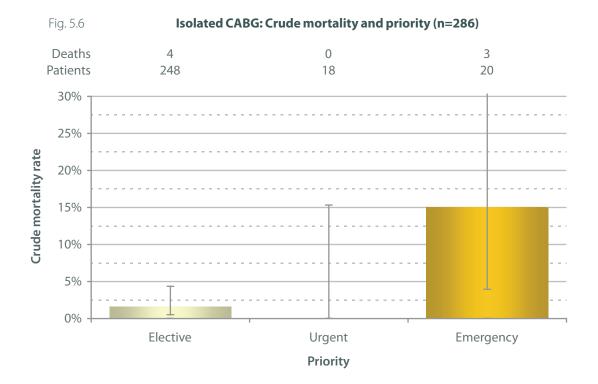
- As shown in the chart, most of the patients underwent CABG on elective basis (86.7%).
- Urgent CABG is defined by operation on next available working day while emergent CABG (including salvaging surgery) indicates patients need surgery the same day because of clinical situation. These represented 6.3% and 7% of all coronary operation respectively.
- According to the United Kingdom National Adult Cardiac Database Report 2008, emergent and salvage coronary surgery comprised 2.5% of all cases (560/22831).





**Priority and mortality** 

- Operations performed in urgent or emergency setting are always considered as high risk. This is because patients in these situations might have on-going ischemia of myocardium, frequent malignant arrhythmia or even unstable haemodynamics.
- Our data correlate with this idea. Mortality was 15% in emergency setting compared to 1.6% in those performed electively.
- According to the United Kingdom National Adult Cardiac Database Report 2008, emergent and salvaging coronary surgery contributed 2.5% of cases (56%22831). Mortality rate was 10.5% (5%560).





#### Mortality and other risk factors

The table below shows risk factors that are considered significant for coronary surgery. Being female, low body mass index, presence of left main disease, previous cardiac surgery, diabetic, hypertensive and presence of other arterial diseases would have higher mortality according to *EuroSCORE*.

Fisher's exact test showed that there were no significant differences in mortality rates for the different classes of these risk factors (p=0.520; p=0.630; p=0.067; p=0.839; p=0.207; p=0.568; p=0.491 respectively)

			Mortality		
			Alive	Died	Rate
	Gender	Male	217	6	2.7%
	Gender	Female	62	1	1.6%
	Dedu Mess Indeu	≥25 kg m <sup>-2</sup>	117	3	2.5%
	Body Mass Index	<25 kg m <sup>-2</sup>	160	4	2.4%
	Left main stem disease	No	138	1	0.7%
S		Yes	139	6	4.1%
Risk factors	Previous cardiac surgery	No	272	7	2.5%
skf		Yes	7	0	0.0%
Ri	Diabetes	No	144	2	1.4%
		Yes	135	5	3.6%
	I have a store size a	No	31	1	3.1%
	Hypertension	Yes	248	6	2.4%
	Extra-cardiac	No	254	6	2.3%
	arteriopathy	Yes	25	1	3.8%

Isolated CABG surgery: in-hospital, post-operative mortality rates for various risk factors



#### The grafting process

#### Arterial grafting at each site

- There were 1,067 distal anastomoses made in 286 patients. On average, each patient received 3.7 grafts in isolated CABG.
- Arterial grafts, especially the left internal mammary artery (LIMA), are considered as better conduit in CABG. Younger patients may benefit from a second arterial graft, usually radial artery or right internal mammary artery.
- We had approaching 90% arterial graft usage in isolated CABG (243 LIMA and 54 radial artery). While almost 20% of them had 2 or more arterial graft conduit.
- Usage of left internal mammary artery in United Kingdom was around 93% in 2008 and 18% of patients received 2 or more arterial grafts.

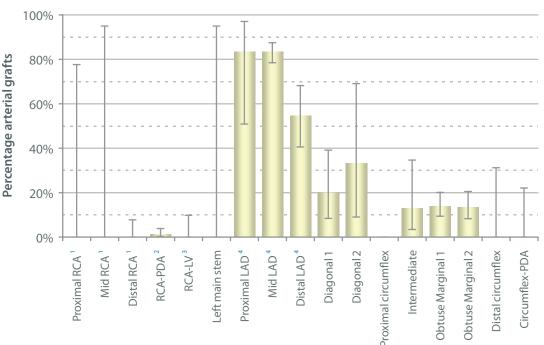


Fig. 5.7Isolated CABG: Arterial graft usage at each of the coronary artery sites treated<br/>(n=1,059 conduits)

#### Coronary artery site

- 1. RCA  $\rightarrow$  Right coronary artery.
- 2. PDA  $\rightarrow$  Posterior descending artery.
- 3. LV  $\rightarrow$  Left ventricle.
- 4. LAD  $\rightarrow$  Left anterior descending artery.



#### **Endoscopic harvest of conduits**

- Our department began using endoscopic vein harvesting techniques in 2005; we began endoscopic radial artery harvesting in 2007. This is now become the standard and preferred way of harvesting these conduits in our practice.
- 54 patients had a radial artery used as conduit. In all but one of these patients the radial artery was harvested using the endoscopic technique.
- For the 262 patients who had a vein graft 85.1% of them were harvested by endoscopic method.

**Isolated CABG surgery:** endoscopic conduit harvest for patients where the named conduit was used in the CABG

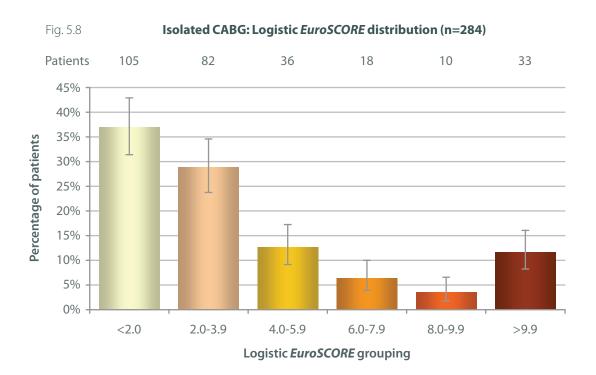
		Endoscopic harvest of the named conduit		
		No	Yes	Endoscopic harvest rate
onduit	Radial artery used	1	53	98.1%
Con	Vein used	39	223	85.1%



#### **Risk adjustment**

#### Logistic EuroSCORE distributions

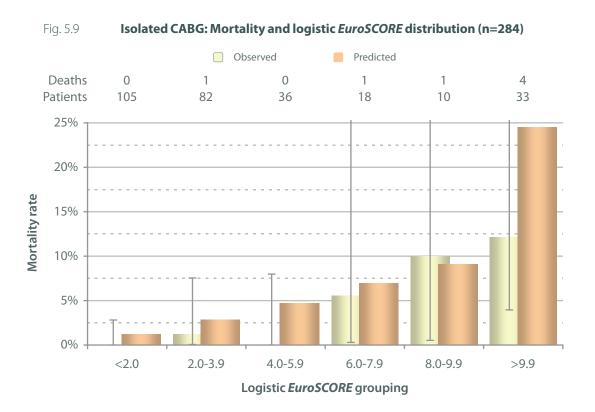
- Logistic *EuroSCORE* is a commonly used risk stratification and prediction method in cardiac surgery. The value equals to the predicted mortality risk for a particular patient.
- In the isolated CABG group, 37% of patients fell into low risk group (<2% mortality) and the number descended in higher risk groups.
- We performed more high risk coronary surgery. As 11.6% of patients were in the highest risk group (>9.9% mortality). When compare to UK data, 8.1% of patients fell into this group.





#### Logistic EuroSCORE and mortality

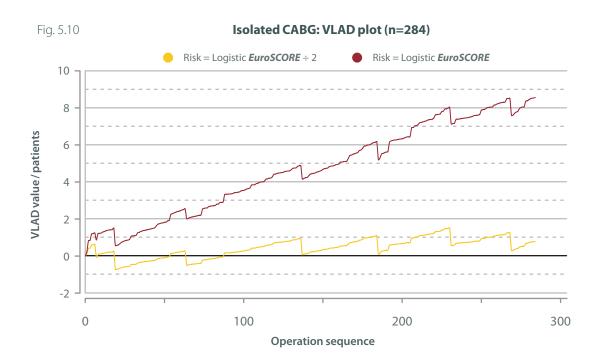
- Observed mortality rates were close to the predicted rates. Most of the mortality was associated with the higher-risk groups.
- Overall predicted mortality was 5.5% (15.6 patients). While observed mortality was 2.4% (7 patients). The observed *versus* predicted mortality ratio was 0.44 for isolated CABG.



- This Variable Life-Adjusted Display (VLAD) covers all risk-scored isolated CABG procedures performed during 2010 and 2011. The plot is risk adjusted and performance as predicted should run approximately around the horizontal zero line (the heavy black line).
- The upslope of the curve demonstrated a net gain of patients' life and the performance was better than predicted.
- At the end of the curve, almost 9 lives had been saved at Queen Mary Hospital.



• An *EuroSCORE* divided by 2 is also shown in the graph. This graph demonstrates performance very close to the predicted value.





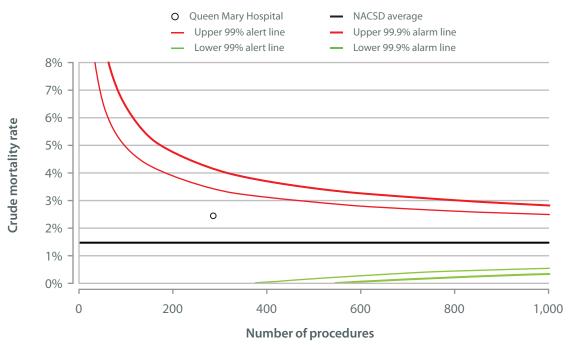
#### International benchmarking of outcomes

- Funnel plots are a graphical means of displaying outcomes compared to a given standard, with upper and lower control limits to define a range of acceptable results.
- The following pages show funnel plots for the outcomes:
  - Crude mortality
  - Re-operation for bleeding
  - Post-operative stroke
  - New haemofiltration / dialysis
- All four of the following charts compare the results at Queen Mary Hospital against the 2008 results from the United Kingdom National Adult Cardiac Surgical Database Report, with alert and alarm lines set at 99.0% and 99.9% respectively.
- The first plot shows that the crude mortality at Queen Mary Hospital rate (2.4%) fell well within the alert lines. Although it was slightly higher than the average from United Kingdom, the rate is not adjusted to take account of the patients' risk profile.
- The second chart places the Queen Mary Hospital bleeding rate in the context of United Kingdom results. Queen Mary Hospital's re-operation rate was 2.1%, which was between the alert lines of the funnel plot, and actually lower than the average reported from the United Kingdom database.
- The third and fourth charts represent the crude stroke rate and the proportion of patients that need haemofiltration / dialysis for acute renal failure; the rates for both of these outcomes at Queen Mary Hospital fell well within the funnel plot alert lines, and almost exactly on the United Kingdom average line.
- There is evidence that the results for these four outcomes at Queen Mary Hospital are in line with internationally-published results from the United Kingdom.



#### **In-hospital mortality**

#### Fig. 5.11 Isolated CABG: Crude mortality rate for QMH (n=286) compared to the data from the financial year 2008 in the United Kingdom NACSD



#### **Re-operation for bleeding**

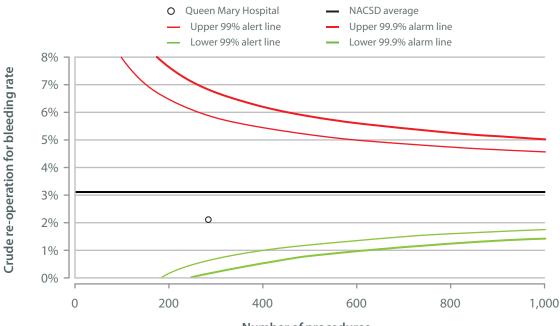
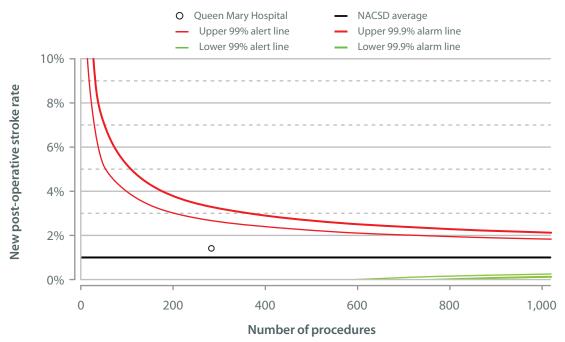


Fig. 5.12Isolated CABG: Crude re-operation for bleeding rate for QMH (n=284) compared<br/>to the data from the financial year 2008 in the United Kingdom NACSD

Number of procedures



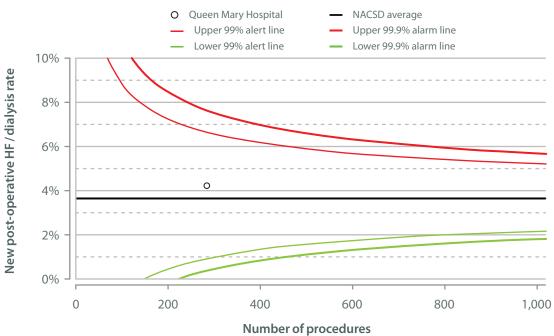
#### Post-operative stroke



### Fig. 5.13 Isolated CABG: Crude post-operative stroke rate for QMH (n=283) compared to the data from the financial year 2008 in the United Kingdom NACSD

New haemofiltration / dialysis





#### Queen Mary Hospital, Hong Kong





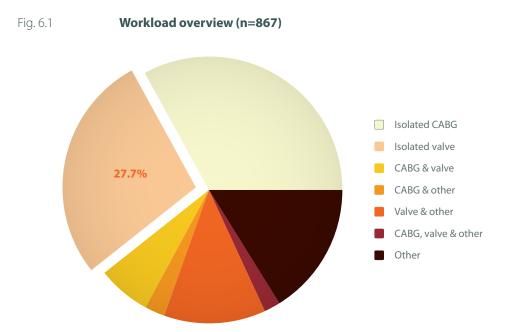
## Isolated valve surgery



# **Isolated single valve surgery**

## Isolated valve surgery in the context of overall workload

- During the year 2010 & 2011, there were 240 isolated valve operations performed at Queen Mary Hospital, contributing to 27.7% of the overall workload.
- Other than that, there were 180 (20.8%) valve operations performed in combination with the other procedures like CABG, radiofrequency atrial ablation, aortic surgery, septal defect repair, *etc*.
- Among the 240 isolated valve operations, there were 11 isolated tricuspid valve operations and 11 isolated pulmonary valve operations.





## **Priority**

- There were 151 isolated single valve operations performed during the year 2010 & 2011.
- 135 out of 151 (89.4%) isolated single valve operations were performed electively.
- The remaining urgent or emergency operations were carried out for infection, jammed mechanical valves, acute decompensated heart failure or patients with unstable haemodynamics.

Isolated single valve surgery: operative urgency distribution

		Valve treated			
		Aortic alone	Mitral alone	Other singles	
	Elective	56	57	22	
riority	Urgent	5	5	0	
Prio	Emergency	3	3	0	
<u> </u>	All	64	65	22	

## Previous cardiac surgery

- In all isolated single valve operations, 41 out of 151 (27.7%) had previous cardiac operations.
- In all isolated multiple valve operations, 30 out of 89 (33.7%) had previous cardiac surgeries.
- Among those redo cardiac operations, some of them are second or more redo operations.

Isolated single valve surgery: prior cardiac surgery

		Valve treated					
		Aortic alone	Mitral alone	Other singles	All		
Previous surgery	No previous cardiac surgery	58	51	1	110		
	Previous cardiac surgery	6	14	21	41		
	All	64	65	22	151		



## Haemodynamic pathology

- Over 50% isolated aortic valve operations were for patients with aortic stenosis.
- For patients who had isolated aortic valve replacement, 20 out of 64 (31.3%) received biological valves while remaining 44 (68.7%) received mechanical prostheses.
- In all patients who had isolated mitral valve surgery, 67.7% of them had mitral regurgitation.

Isolated single valve surgery: haemodynamic pathology distribution

		Valve treated			
		Aortic alone	Mitral alone	Other singles	
Haemodynamic pathology	Stenosis	33	14	0	
	Regurgitation	18	44	16	
	Mixed	12	5	1	



## Native valve pathology

- Calcific degenerative disease (46.9%) and bicuspid aortic valve disease (43.8%) represented the two most common pathology for aortic valve operations.
- While for the isolated mitral valve surgery, more than half were for degenerative changes (56.9%), and nearly a quarter were for chronic rheumatic changes (24.6%), and these 2 etiologies were the two major causes for mitral valve surgery in this locality.
- There were 2 cases of aortitis in this cohort.

		Valve treated				
		Aortic alone	Mitral alone	Other singles		
	Native valve not present	3	3	0		
	Congenital	28	1	9		
gy	Degenerative	17	37	1		
pathology	Active infective endocarditis	4	5	0		
bath	Previous infective endocarditis	0	8	0		
valve p	Rheumatic	6	16	7		
e val	Annuloaortic ectasia	4	0	0		
Native	Calcific degeneration	30	1	0		
N	Ischaemic	0	1	0		
	Functional regurgitation	0	0	4		
	Other	2	1	0		

Isolated single valve surgery: native valve pathology distribution



## **Mitral valve repair**

## Mitral valve repair in the context of all mitral valve surgery

- For isolated valve surgery, there were 136 operations involving mitral valve in year 2010 & 2011.
- Out of these 136 operations, 71 (52.2%) were operated for regurgitant valves. The rest of them were mainly for rheumatic mitral stenosis, which is still a common pathology in this locality.
- Mitral valve repair is always the procedure of choice for treatment of mitral regurgitation in Queen Mary Hospital. The goals of valve repair include preservation of leaflet mobility, restoration of leaflet coaptation and stabilization of repair with remodeling annuloplasty.
- Based on The Society for Cardiothoracic Surgery (SCTS) in Great Britain & Ireland 6<sup>th</sup> National Adult Cardiac Surgical Database Report, in 2008, 67% underwent mitral valve repair for degenerative mitral valve disease.
- Our proportion of mitral valve repair surgery was comparable to international benchmark. During the year 2010 to 2011, 58 out of 71 (81.7%) regurgitant mitral valves were successfully repaired.

		Haemodynamic pathology			
		Stenosis	Regurgitation	Mixed	
Valve treated	Replacement	43	13	18	
	Repair	0	58	0	
	All	43	71	18	

Mitral valve surgery: haemodynamic pathology and type of valve procedure



## Type of mitral vale repair

- All isolated mitral valve repair operations were complex repairs, involving two or more repair procedures.
- The reconstructive techniques are systematic that involve intraoperative valve inspection & analysis, meticulous application of repair techniques, implantation of remodeling annuloplasty and finally evaluation of repair by saline test and transesophageal echocardiogram.
- More than 90% of mitral valve repair operations had ring annuloplasty.
- Nearly 60% of operations had implanted one or more artificial chords.

Isolated valve surgery involving the mitral valve repair: type of valve repair

		Data		
		Count	Rate	
	Commisurotomy	3	5.8%	
	Annuloplasty (ring)	48	92.3%	
	Annuloplasty (suture)	1	1.9%	
air	Leaflet resection	19	36.5%	
repair	Leaflet extension	1	1.9%	
ve	Chordal transfer	3	5.8%	
l va	Chordal shortening	0	0.0%	
Type of mitral valve	Artificial chord	30	57.7%	
of m	Papillary muscle repositioning	0	0.0%	
be d	Decalcification/debridement	1	1.9%	
Ţ	Leaflet patch	3	5.8%	
	Sub-valvar release	5	9.6%	
	Re-suspension	2	3.8%	
	Other	0	0.0%	



## Tricuspid valve repair

## Tricuspid vale repair in the context of all triscupid valve surgery

- At tricuspid valve position, repair surgery is the preferred technique for correcting tricuspid regurgitation because of its better surgical outcome and lower mortality.
- Tricuspid valve surgery mirror the surgical techniques used in mitral valve repair surgery.
- Most of the tricuspid valve surgery (85.5%) were performed in the presence of other valves pathology.
- Majority of the tricuspid valve alone operations were redo operations for rheumatic etiology, few of them were due to infective endocarditis.
- In our cohort, 73 out of 76 (96%) tricuspid valve pathology were being repaired.

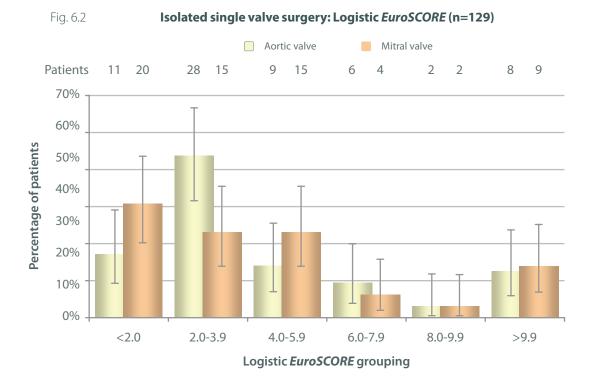
Isolated valve surgery involving the tricuspid valve: type of valve procedure

		Tricuspid valve procedure				
		Replacement	Repair	All		
Valve treated	Tricuspid alone	2	9	11		
	Tricuspid plus another valve	1	64	65		
	All that include tricuspid valve surgery	3	73	76		



## Logistic EuroSCORE

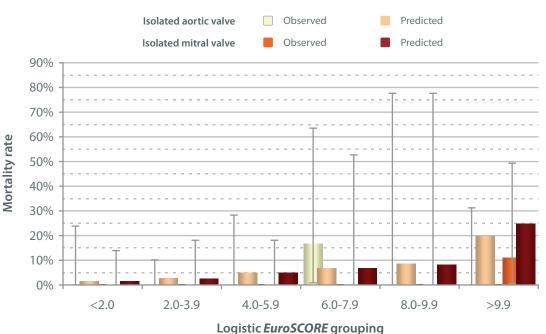
- Using the logistic *EuroSCORE*, most of our isolated single valve surgery had the predicted operation risks between 2-4%.
- While only around 15% of patients had their predicted risks greater than 9.9%.
- The mean ICU stay after isolated aortic valve surgery and mitral valve surgery were 2.4 days and 2.2 days respectively.
- The predicted risk for isolated aortic valve surgery alone in this cohort was <sup>3.5</sup>/<sub>4</sub>= 5.5%.
- The predicted risk for isolated mitral valve surgery alone in this cohort was <sup>4.</sup>/<sub>65</sub>= 6.3%.
- The predicted risk for all single valve (Aortic + Mitral) in this cohort was <sup>8.9</sup>/<sub>129</sub>= 6.9%.





## Mortality

- There were 2 deaths following single valve surgery:
  - a 75-year-old male patient who had an isolated AVR; EuroSCORE = 6.8
  - a 78-year-old NYHA 4 female patient who had an emergency MVR; *EuroSCORE* = 63.8
- The observed mortality for isolated aortic valve surgery was 1/64=1.6% and observed/predicted ratio was 0.29
- The observed mortality for isolated mitral valve surgery was <sup>1</sup>/<sub>65</sub> = 1.5% and observed/predicted ratio was 0.24
- The observed mortality for all single valve (aortic + mitral) was <sup>2</sup>/<sub>129</sub>=
   1.6% and the observed/ predicted ratio was 0.23
- The observed mortality was lower than the predicted mortality in all groups of valvular patients.



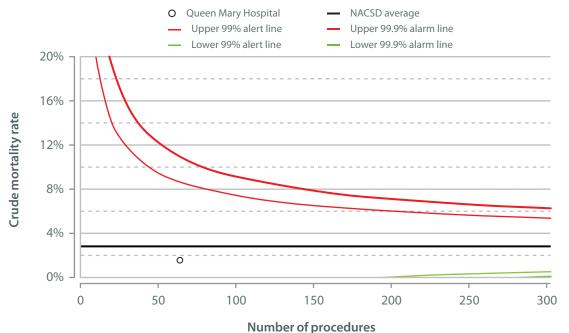
## Fig. 6.3Isolated single valve surgery: Mortality and logistic EuroSCORE (n=129)



## International benchmarking of mortality

- The graph here is a funnel plot of in-hospital crude mortality for isolated aortic valve surgery, with alert lines and United Kingdom National Adult Cardiac Surgical Database (2008) average line.
- The crude mortality in isolated aortic valve was 1.6%, which was lower than the average mortality in United Kingdom. And this number fell in the normal distribution.

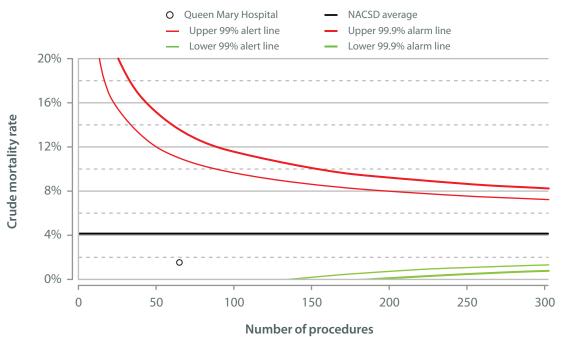






- The graph here is a funnel plot of in-hospital crude mortality for isolated mitral valve surgery, with alert lines and United Kingdom National Adult Cardiac Surgical Database (2008) average line.
- The crude mortality in isolated mitral valve surgery was 1.5%, which fell in the normal distribution and lower than the average from United Kingdom.

## Fig. 6.5Isolated mitral valve surgery: Crude mortality rate for QMH (n=65) compared to<br/>the data from the financial years 2004-2008 in the United Kingdom NACSD





# Isolated multiple valve surgery

## Valves treated

- There were 83 patients having multiple valve surgery contributing to 34.6% (<sup>83</sup>/<sub>240</sub>) of the overall isolated valve operations.
- Among these 83 patients, more than 20% (18/83) had triple valve surgery.

Isolated multiple valve surgery: valves treated

		Valve treated			
		Male	Female	All patients	
ted	Aortic & mitral	8	10	18	
	Aortic & tricuspid	4	6	10	
treated	Mitral & tricuspid	11	24	35	
vest	Pulmonary & tricuspid	1	1	2	
Valves	Aortic, mitral & tricuspid	6	12	18	
	All	30	53	83	

## Mortality

There were only 3 deaths in this group: all were female patients undergoing elective surgery:

- The first patient had a combined mitral valve repair & tricuspid valve repair; she was a 65-year-old patient, with a NYHA grade of 3 and a logistic *EuroSCORE* of 3.1; she had not had any cardiac surgery in the past.
- The second patient had a combined mitral valve replacement & tricuspid valve repair; she was a 46-year-old patient, with a NYHA grade of 3 and a logistic *EuroSCORE* of 5.5; she had had a cardiac surgical procedure in the past.
- The third patient had a combined aortic valve replacement, mitral valve replacement & tricuspid valve repair; she was a 59-year-old patient, with a NYHA grade of 3 and a logistic *EuroSCORE* of 14.5; she had had a cardiac surgical procedure in the past.

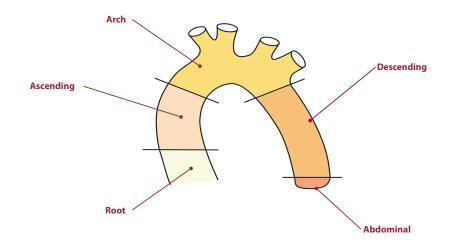


# **Other procedures**



## Surgery on the aorta

## **Segments treated**



Surgery on the aorta: number and details of segments treated

			Cardiac procedure group				
			CABG & Other	Valve & other	CABG, valve & other	Other <sup>1</sup>	All
		Root	2	7	3	5	17
		Ascending	2	7	3	40	52
q	1	Arch	0	0	0	2	2
Segments treated		Descending	0	1	0	2	3
s tre		Abdominal	0	0	0	0	0
lent		Root & ascending	1	6	0	5	12
egn	2	Ascending & arch	1	1	0	6	8
Š		Descending & abdominal	1	0	0	0	1
	3	Root, ascending & arch	0	1	0	1	2
		All	7	23	6	61	97

1. Includes surgery aorta  $\pm$  another procedure (other than CABG and valve surgery).



## **Surgical technique**

- In aortic surgery involving the ascending aorta and the aortic arch, the preferred cannulation site is through the axillary or subclavian artery, both in emergency and elective settings, to achieve antegrade flow to end-organs and avoid obliterations of the true lumen, which is more seen in retrograde arterial perfusion.
- In aortic aneurysm with no dissection, femoral artery and ascending is still used for arterial cannulation.
- Descending thoracic aorta has also been used as a cannulation site for cases where a left thoracotomy is required
- 60 patients required deep hypothermic circulatory arrest (DHCA)
- 98.3% (59 out of 60) of DHCA had antegrade cerebral perfusion via direct cannulation of the neck vessels at the aortic arch for cerebral protection.
- Cerebral oximetry with near infra-red spectrometry was used through out the operation to assess adequacy of cerebral oxygenation

Surgery on the aorta: cannulation

		Count				Count
al tion	Ascending aorta	31		_	Right atrial	5
	Axillary / subclavian	50		/enou inulat	RA / IVC 2-stage	71
Arterial	Femoral	21			Bi-caval	7
A	Other	1			Femoral	18
U	All	97			All	97

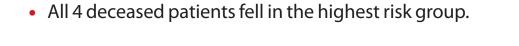
#### Surgery on the aorta: cerebral perfusion

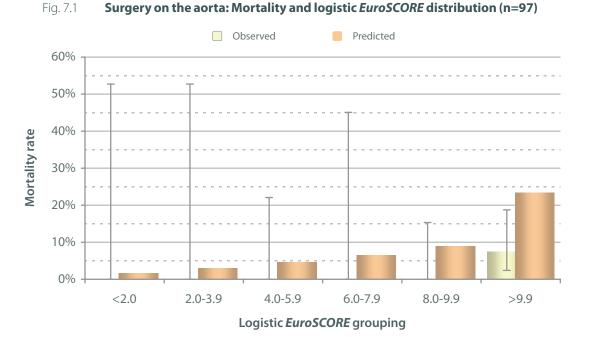
		Count
al on	None	30
<b>Cerebral</b> <b>Derfusion</b>	Antegrade	59
De De	Retrograde	1



## Mortality and morbidity

- Out of the 97 patients operated, there were 3 (3.1%) transient ischaemic attacks and 7 (7.2%) cerebral vascular accidents.
- 15 (15.5%) required reopen for haemostasis.
- 4 (4.1%) patients required renal replacement therapy.
- Predicted mortality of aortic surgery was 15.7% by using logistic *EuroSCORE*. The observed mortality was 4.1%. Predicted / observed mortality ratio was 0.26 in our centre.







# **Other cardiac procedures**

## **Adult congenital**

- In this report, we have defined surgery for adult congenital heart disease as patients >18 years-old who underwent surgery for congenital heart conditions.
- Any patients <18 years of age undergoing surgery for congenital heart conditions will be included in the paediatric population and therefore will not be reported in this section.
- In 2010 and 2011, 50 and 41 patients respectively underwent surgery for congenital heart conditions. Amongst these, 17 were repair of atrial septal defects (ASD), 15 were repair of ventricular septal defects (VSD), 25 were procedures on the right ventricular outflow tract with or without pulmonary valve replacement (PVR), 4 were heart transplantation, and the others include complex repair including surgery for Ebstein anomaly, aortic surgeries, *etc*.
- There was one mortality reported. This is a 70-year-old patient with secundum ASD, severe pulmonary hypertension and severe tricuspid regurgitation. He was unfortunately complicated by post-operative mediastinitis. Radical wound debridement and flap reconstruction was performed and prolonged antibiotics were given. However, the patient's condition deteriorated and finally succumbed from overwhelming sepsis.



## Queen Mary Hospital, Hong Kong First Cardiac Surgery Report 2012

Procedure detail

		Procedure groupings							
		Isolated CABG	Isolated valve	CABG & valve	CABG & other	Valve & other	CABG, valve & other	Other	All
	None recorded	284	235	56	9	19	6	59	668
	LV aneurysmectomy	0	0	0	1	0	0	1	2
	Acquired VSD	0	0	0	1	0	2	0	3
S	Atrial myxoma	0	0	0	0	0	0	5	5
dure	Pulmonary embolectomy	0	0	0	0	0	0	4	4
oced	Cardiac transplant	0	0	0	0	0	0	21	21
c pre	Pulmonary transplant	0	0	0	0	0	0	2	2
Other cardiac procedures	Epicardial pacemaker	0	0	0	0	1	0	1	2
r car	Pericardiectomy	0	0	0	1	1	0	0	2
the	ASD	0	2	0	0	16	0	14	32
0	Congenital surgery	0	2	0	1	17	1	19	40
	Atrial ablation	0	0	0	3	42	5	1	51
	Other procedures	2	2	0	5	22	3	19	53
	Patient denominator	286	240	56	21	107	17	140	867



## **Atrial ablation**

## **Patients and procedures**

- 51 atrial ablation were performed in the year 2010 to 2011. All of them were performed with other procedures.
- We adopted the modified Cox-Maze III procedure, using radiofrequency ablation for surgery for atrial fibrillation.
- From our own database in 2005 to 2009, amongst the 122 patients who underwent concomitant radiofrequency ablation for atrial fibrillation during open heart surgery, 77.5% patients were in sinus rhythm at 1 year follow-up. Positive predictors for success in radiofrequency ablation surgery were a left atrial size of < 55mm (p = 0.032) and a patient age < 60 years old (p = 0.004).</li>
- Nowadays, we would offer concomitant radiofrequency ablation during cardiac surgery in all suitable cardiac surgery candidates presenting to us with atrial fibrillation.

		Gender							
		Male	Female	All	Percentage				
Ś	<56	13	8	21	41.2%				
/ear	56-60	2	6	8	15.7%				
( / <b>V</b>	61-65	5	7	12	23.5%				
rge	66-70	4	2	6	11.8%				
it su	>70	2	2	4	7.9%				
Age at surgery / years	All	26	25	51					
A	Percentage	51.0%	49.0%						

Atrial ablation: age and gender

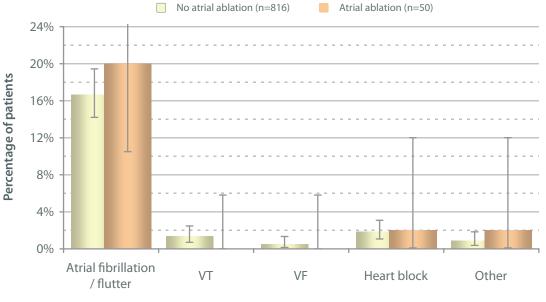


## Outcomes

- No deaths were recorded in the group of patients who had an atrial ablation procedure.
- The immediate post-operative success rate, defined as maintenance of sinus rhythm in the in-patient period, was 74.5% (<sup>38</sup>/<sub>51</sub>).
- Risk of heart block after atrial ablation was similar to patients without any arrhythmia surgery.

Atrial ablation: post-operative arrhythmia requiring intervention

		Procedure			
		No atrial ablation	Atrial ablation		
	None	651	38		
rative mia	Atrial fibrillation / flutter	136	10		
ъ е	VT	11	0		
t-op rhyt	VF	4	0		
Postari	Heart block	15	1		
	Other	7	1		



## Atrial ablation: Post-operative arrhythmias requiring intervention

Post-operative rhythm requiring intervention





# Appendices

CTSI

1-14



# **Appendices**

## **Database form**

	Adult Cardiac S	ospital, Hong Kong <b>Surgical Database</b> Version 1.1	3			
	Patient identification a	nd demographics				
Hospital number		Date of birth	dd / mm / yyyy			
Given name		Date / time of operation	dd / mm / yyyy hh:mm			
Family name		Gende	r O 1. Male O 2. Fem			
	Admission details & ca	rdiac history				
Date of referral	dd/mm/yyyy	Outpatient clini	c select from list			
Date of admission	dd/mm/yyyy					
Admission category	O 1. Health Authority	O 2.	Private			
Mode of admission	<ul><li>O 1. Elective</li><li>O 2. Planned inpatient</li></ul>	transfer O 3.	Emergency			
Angina status pre-surgery	<ul> <li>0. No angina</li> <li>1. No limitation of physical activity</li> <li>2. Slight limitation of ordinary activity</li> <li>3. Marked limitation of ordinary physical activity</li> <li>4. Symptoms at rest or minimal activity</li> </ul>					
Oyspnoea status pre-surgery	<ul> <li>1. No limitation of physical activity</li> <li>2. Slight limitation of ordinary activity</li> <li>3. Marked limitation of ordinary physical activity</li> <li>4. Symptoms at rest or minimal activity</li> </ul>					
Congestive cardiac failure	O0. NeverO1. In the pastOO2. Now					
Symptom status	O 1. Stable	O 2.	Unstable/recent deterioration			
Number of previous MIs	<ul><li>O. None</li><li>O. 1. One</li></ul>		Two or more Unknown			
Interval between surgery and last MI	<ul> <li>O. No previous MI</li> <li>1. MI &lt; 6 hours</li> <li>2. MI 6-24 hours</li> <li>3. MI 25-48 hours</li> </ul>	O 5.	MI 2-30 days MI 31-90 days MI > 90 days			
	Previous interventions					
Previous PCI	<ul> <li>O. No previous PCI</li> <li>1. PCI &lt; 24 hours before surgery</li> <li>2. PCI &gt; 24 hours before surgery; same admission</li> <li>3. PCI &gt; 24 hours before surgery; previous admission</li> </ul>					
Date of last PCI	dd/mm/yyyy					
Previous cardiac surgery	<ul> <li>O. No previous cardia</li> <li>1. CABG</li> <li>2. Valve</li> <li>3. Congenital cardiaa</li> <li>4. Other cardiaa</li> </ul>	□ 6. □ 7. c □ 8	Aortic - ascending / arch Aortic - descending / abdomina Other thoracic Carotid endarterectomy Other peripheral vascular			
Date of last cardiac operation	dd/mm/yyyy					



Adult Cardiac Surgical Database Page 2; Version 1.1         Hospital number       Date of surgery       dd/mm / yyyy         Previous PCI       0. No previous PCI       0. A to previous admission         Date of last PCI       dd/mm/yyyy       0. A to previous cardiac surgery       1. A to previous cardiac surgery       1. A totic - ascending / arch         Date of last PCI       dd/mm/yyyy       0. A top revious cardiac surgery       1. C totic + toracic         2. Valve       0. A top revious cardiac surgery       0. A totic - descending / arch         2. Valve       0. A totic - ascending / arch         0. A totic - additic       0. A totic - ascending / arch         0. A totic - additic       0. A totic - ascending / arch         0. A totic - additic       0. A totic - additic         0. A totic - additic       0. A totic - additic         0. A totic - additic       0. A totic - additic         0. B totic       0. A totic - additic         0. A totic - additic       0. A totic - additic         0. A totic - additic       0. A totic - add										
Page 2; Version 1.1         Proprint number       Date of surgery       dd /mm /yyyy         Previous PCI       0. No previous PCI       0. PCI < 24 hours before surgery; same admission										
Previous Interventions         Previous PCI       0. No previous PCI         1. PCI < 24 hours before surgery;										
Previous PCI       0. No previous PCI         0. PCI > 24 hours before surgery;       2. PCI > 24 hours before surgery; same admission         0. Date of last PCI       dd/mm/yyyy         Previous cardiac surgery       5. Aortic - ascending / arch         1. CABG       6. Aortic - descending / abdominal         2. PCI > 24 hours before surgery; previous admission       6. Aortic - descending / abdominal         1. CABG       6. Aortic - descending / abdominal         2. Valve       7. Other thoracic         2. Valve       7. Other thoracic         3. Congenital cardiac       8. Carotid endarterectomy         4. Other cardiac       9. Other peripheral vascular         Date of last cardiac operation       dd/mm/yyyy         Risk factors for acquisition of coronary disease       0. No to diabetic       2. Current smoker         0. Diabetes       0. No Never smoked       1. Ex smoker       2. Current smoker         Hypercholesterolaemia       0. No       1. Yes       1. Functioning transplant _         Family history of Hyp       0. No       1. Yes         Renal function / dialysis       0. No       1. Yes         History of pulmonary disease       3. Jalaysis for chronic renal failure; onset within 6 weeks of cardiac surgery         4. Dialysis for chronic renal failure; onset within 6 weeks prior to car	Hospital number   Date of surgery     dd / mm / yyyy									
0       1. PCl < 24 hours before surgery; same admission         0       2. PCl > 24 hours before surgery; same admission         0       3. PCl > 24 hours before surgery; previous admission         Date of last PCl       dd/mm/yyyy         Previous cardiac surgery       5. Aortic - ascending/arch         1. CABG       6. Aortic - descending/abdominal         2. Valve       7. Other thoracic         3. Congenital cardiac       8. Carotid endarterectomy         4. Other cardiac       9. Other peripheral vascular         Date of last cardiac operation       dd/mm/yyyy         Date of last cardiac operation       dd/mm/yyyy         Ibabetes       0. No to diabetic       2. Current smoker         0       0. Not diabetic       2. Current smoker         1. Ex smoker       2. Current smoker         Hypercholesterolaemia       0. No       1. Yes         History of hypertension       1. Texet dor BP>140/90 on >1 occasion prior to admission         0. Unknown       1. Functioning transplant ;       2. Creatinne >200 µmol 8         3. Jolayisis for acute renal failure; onset within 6 weeks of cardiac surgery       4. Dialysis for acute renal failure; onset more than 6 weeks prior to cardiac surgery         4. Dialysis for acute renal failure; onset within 6 weeks of cardiac surgery       5. Unknown       1. Yes		Previous interventions								
Previous cardiac surgery       5. Aortic - ascending/arch         1. CABG       6. Aortic - descending/abdominal         2. Valve       7. Other thoracic         3. Congenital cardiac       8. Carotid endarterectomy         4. Other cardiac       9. Other peripheral vascular         Date of last cardiac operation       dd/mm/yyyy         Risk factors for acquisition of coronary disease         Diabetes       0. Not diabetic       2. Oral therapy         0. Not diabetic       2. Current smoker       2. Current smoker         1. Diet       3. Insulin         Cigarette smoking history       0. No       1. Yes         History of hypertension       0. No       1. Yes         Renal function / dialysis       0. No       1. Yes         Renal function / dialysis       0. No       1. Yes         History of pulmonary disease       3. Dialysis for chronic renal failure; onset within 6 weeks of cardiac surgery       3. Dialysis for chronic renal failure; onset within 6 weeks of cardiac surgery         5. Unknown       1. Yes       3. Dialysis for chronic renal failure; onset within 6 weeks of cardiac surgery         6. Unknown       1. Yes       3. Dialysis for chronic renal failure; onset within 6 weeks of cardiac surgery         6. Unknown       1. Yes       3. Dialysis for chronic renal failure; onset withi	Previous PCI	<ul> <li>O 1. PCI &lt; 24 hours before surgery</li> <li>O 2. PCI &gt; 24 hours before surgery; same admission</li> </ul>								
Image: Construct on the second seco	Date of last PCI	dd / mm / yyyy								
Risk factors for acquisition of coronary disease         Diabetes       0. Not diabetic       2. Oral therapy         1. Diet       3. Insulin         Cigarette smoking history       0. Never smoked         0. 1. Ex smoker       2. Current smoker         Hypercholesterolaemia       0. No       1. Yes         History of hypertension       0. No hypertension       1. Treated or BP>140/90 on >1 occasion prior to admission         9. Unknown       1. Yes         Family history of IHD       0. No       1. Yes         Renal function/dialysis       0. None       1. Yes         1. Functioning transplant       2. Creatinine >200 µmol ℓ       3. Dialysis for acute renal failure; onset within 6 weeks of cardiac surgery         4. Dialysis for chronic renal failure; onset more than 6 weeks prior to cardiac surgery       5. Unknown         History of pulmonary disease       3. Neoplasm         1. COAD/ephysema       4. Infective lung disease         1. COAD/ephysema       4. Infective lung disease         2. Asthma       9. Other         Neurological dysfunction       0. No       1. Yes	Previous cardiac surgery	<ul> <li>1. CABG</li> <li>2. Valve</li> <li>3. Congenital cardiac</li> </ul>	<ul> <li>6. Aortic - descending / abdominal</li> <li>7. Other thoracic</li> <li>8. Carotid endarterectomy</li> </ul>							
Diabetes00. Not diabetic02. Oral therapy01. Diet3. InsulinCigarette smoking history0. Never smoked2. Current smokerHypercholesterolaemia0. No1. Ex smoker2. Current smokerHistory of hypertension0. No hypertension1. Treated or BP>140/90 on >1 occasion prior to admission9. Unknown9. Unknown1. YesRenal function / dialysis0. No1. YesCreatine >200 µmol l3. Dialysis for acute renal failure; onset within 6 weeks of cardiac surgery4. Dialysis for chronic renal failure; onset more than 6 weeks prior to cardiac surgery5. Unknown1. YesHistory of pulmonary disease3. Neoplasm1. COAD / ephysema4. Infective lung disease2. Asthma9. OtherNeurological dysfunction0. No1. Yes	Date of last cardiac operation	dd / mm / yyyy								
Diabetes00. Not diabetic02. Oral therapy01. Diet3. InsulinCigarette smoking history0. Never smoked2. Current smokerHypercholesterolaemia0. No1. YesHistory of hypertension0. No hypertension1. Treated or BP>140/90 on >1 occasion prior to admission9. Unknown1. Treated or BP>140/90 on >1 occasion prior to admissionFamily history of IHD0. No1. YesRenal function / dialysis0. None1. Yes1. Functioning transplant2. Creatinine >200 µmol l²3. Dialysis for acute renal failure; onset within 6 weeks of cardiac surgery4. Dialysis of or chronic renal failure; onset more than 6 weeks prior to cardiac surgery5. Unknown1. YesHistory of pulmonary disease3. Neoplasm1. COAD / ephysema4. Infective lung disease2. Asthma9. OtherNeurological dysfunction0. No1. Yes		Risk factors for acquisition of cor	onary disease							
Image: Constant of the service of t	Diabetes									
History of hypertension       0. No hypertension         0. 1. Treated or BP>140/90 on >1 occasion prior to admission         9. Unknown         Family history of IHD         0. No       1. Yes         Renal function/dialysis         0. None         1. Functioning transplant         2. Creatinine >200 µmol ℓ         3. Dialysis for acute renal failure; onset within 6 weeks of cardiac surgery         4. Dialysis for chronic renal failure; onset more than 6 weeks prior to cardiac surgery         5. Unknown         Hyperthyroidism         0. No       1. Yes         History of pulmonary disease         1. COAD/ephysema       3. Neoplasm         2. Asthma       9. Other         Neurological dysfunction       0. No       1. Yes	Cigarette smoking history		O 2. Current smoker							
<ul> <li>i. Treated or BP&gt;140/90 on &gt;1 occasion prior to admission</li> <li>9. Unknown</li> <li>i. Treated or BP&gt;140/90 on &gt;1 occasion prior to admission</li> <li>9. Unknown</li> <li>i. Yes</li> <li>Renal function / dialysis</li> <li>0. None</li> <li>1. Functioning transplant</li> <li>2. Creatinine &gt;200 µmol l</li> <li>3. Dialysis for acute renal failure; onset within 6 weeks of cardiac surgery</li> <li>4. Dialysis for chronic renal failure; onset more than 6 weeks prior to cardiac surgery</li> <li>5. Unknown</li> <li>1. Yes</li> <li>History of pulmonary disease</li> <li>1. COAD/ephysema</li> <li>4. Infective lung disease</li> <li>2. Asthma</li> <li>9. Other</li> <li>0. No</li> <li>1. Yes</li> </ul>	Hypercholesterolaemia	O 0. No	O 1. Yes							
Renal function / dialysis       0. None         1. Functioning transplant       2. Creatinine >200 µmol ℓ         3. Dialysis for acute renal failure; onset within 6 weeks of cardiac surgery       4. Dialysis for chronic renal failure; onset more than 6 weeks prior to cardiac surgery         4. Dialysis for chronic renal failure; onset more than 6 weeks prior to cardiac surgery       5. Unknown         Hyperthyroidism       0. No       1. Yes         History of pulmonary disease       3. Neoplasm         1. COAD / ephysema       4. Infective lung disease         2. Asthma       9. Other         Neurological dysfunction       0. No       1. Yes	History of hypertension	O 1. Treated or BP>140/90 on >1	occasion prior to admission							
<ul> <li>□ 1. Functioning transplant</li> <li>□ 2. Creatinine &gt;200 µmol ℓ</li> <li>□ 3. Dialysis for acute renal failure; onset within 6 weeks of cardiac surgery</li> <li>□ 4. Dialysis for chronic renal failure; onset more than 6 weeks prior to cardiac surgery</li> <li>□ 5. Unknown</li> <li>○ 0. No</li> <li>○ 1. Yes</li> <li>History of pulmonary disease</li> <li>□ 1. COAD / ephysema</li> <li>□ 4. Infective lung disease</li> <li>□ 2. Asthma</li> <li>□ 9. Other</li> <li>○ 0. No</li> <li>○ 1. Yes</li> </ul>	Family history of IHD	O 0. No	O 1. Yes							
History of pulmonary disease       0       0. No pulmonary disease       3. Neoplasm         1. COAD/ephysema       4. Infective lung disease       9. Other         Neurological dysfunction       0       0. No       1. Yes	Renal function / dialysis	<ul> <li>1. Functioning transplant</li> <li>2. Creatinine &gt;200 µmol l</li> <li>3. Dialysis for acute renal failure; onset within 6 weeks of cardiac surgery</li> <li>4. Dialysis for chronic renal failure; onset more than 6 weeks prior to cardiac surgery</li> </ul>								
Image: COAD/ephysema       Image: 4. Infective lung disease         Image: 2. Asthma       9. Other         Neurological dysfunction       0. No       1. Yes	Hyperthyroidism	O 0. No	O 1. Yes							
	History of pulmonary disease	1. COAD / ephysema	4. Infective lung disease							
Extra-cardiac arteriopathy O 0. No O 1. Yes	Neurological dysfunction	O 0. No	O 1. Yes							
	Extra-cardiac arteriopathy	O 0. No	O 1. Yes							



Queen Mary Hospital, Hong Kong Adult Cardiac Surgical Database Page 3; Version 1.1								
Hospital number Date of surgery dd / mm / yyyy								
	Additional medical history and risk fac	tors						
History of gastrointestinal disease	<ul> <li>0. None</li> <li>1. Peptic ulcer disease</li> <li>2. Malignancy</li> </ul>	<ul> <li>3. Inflammatory bowel disease</li> <li>4. Major abdominal surgery</li> <li>5. Other</li> </ul>						
Major abdominal surgery	O 0. No	O 1. Yes						
Pre-operative heart rhythm	<ul> <li>O. Sinus rhythm</li> <li>I. Atrial fibrillation / flutter</li> <li>2. Complete heart block / pacing</li> </ul>	<ul><li>3. VF /VT</li><li>4. Other abnormal rhythm</li></ul>						
Saphenous vein	<ul> <li>0. Normal</li> <li>1. Minor varicosites</li> <li>2. Major varicosites</li> </ul>	<ul><li>3. Previous varicose vein surgery</li><li>4. Previous DVT</li></ul>						
Capillary refill (non-dominant hand)	<ul> <li>0. &lt;5 seconds</li> <li>1. 5-10 seconds</li> </ul>	O 2. >10 seconds						
Pre-operative haemoglobin	globin g d $\ell^{-1}$							
Pre-operative creatinine $\mu mol \ \ell^{-1}$								
Cardiac investigations								
Left- or right-heart catheterisation	<ul><li>O. Normal</li><li>I. Minor varicosites</li></ul>	3. Previous varicose vein surgery						
Date of last catheterisation	dd/mm/yyyy							
Extent of coronary vessel disease	<ul> <li>0. No vessel with &gt;50% diameter stenosis</li> <li>1. One vessel with &gt;50% diameter stenosis</li> <li>2. Two vessels with &gt;50% diameter stenosis</li> <li>3. Three vessels with &gt;50% diameter stenosis</li> <li>9. Not investigated</li> </ul>							
Left main stem disease	<ul> <li>0. No LMS disease or LMS disease &lt;= 50</li> <li>1. LMS &gt;50% diameter stenosis</li> <li>9. Not investigated</li> </ul>	0% diameter stenosis						
Left ventricular function	%							
Ejection fraction category	<ul> <li>O 1. Good (LVEF &gt; 50%)</li> <li>O 2. Fair (LVEF 30-50%)</li> </ul>	<ul> <li>3. Poor (LVEF &lt; 30%)</li> <li>9. Not measured</li> </ul>						
Ejection fraction estimate based upon	<ul><li>1. Left ventriculogram</li><li>2. Echocardiogram</li></ul>	<ul><li>3. MR scan</li><li>3. Other investigation</li></ul>						
PA systolic	mm Hg							
AV gradient	mm Hg							
LVEDP	mm Hg							
Mean PAWP LA	mm Hg							



Appendices

		Queen Mary H dult Cardiac Page 4			63		
Hospital number				Date of surg	dd / mm / yyyy		
	Pre-op	erative status a	nd support				
IV nitrates or any heparin	-	). Never smoked . Until operation		0	2. Within one week of surgery		
Pre-operative aspirin <sup>1</sup>	0 0	). No		0	1. Yes		
Pre-operative clopidogrel <sup>1</sup>	0 0	). No		0	1. Yes		
Other anticoagulant	0 0	). No		0	1. Yes		
V inotropes prior to anaesthesia	0 0	). No		0	1. Yes		
Ventilated	0 0	). No		0	1. Yes		
Cardiogenic shock	0 0	). No		0	1. Yes		
	Operat	tion data					
Operative urgency     O     1. Elective     O     3. Emergency       O     2. Urgent     O     4. Salvage							
Number of previous heart operations							
Responsible consultant anaesthetist select from list							
First oper	ator	select from list					
First operator: grade	O 2 O 3	. Consultant 2. Professor 3. Associate profess 4. Specialist	sor	0	5. Associate consultant 6. HST 9. Other		
First operator: year of HST	0 2	. Year 1 2. Year 2 3. Year 3		0	4. Year 4 5. Year 5 6. Year 6 8. Not applicable		
First assis	tant		select from list				
First assistant: grade	O 2 O 3	. Consultant 2. Professor 8. Associate profess 4. Specialist	sor	Ō	5. Associate consultant 6. HST 9. Other		
First assistant: year of HST	0 2	. Year 1 2. Year 2 3. Year 3		0 0	4. Year 4 5. Year 5 6. Year 6 8. Not applicable		
This form is designed so that question		1. Within the					



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Appendices

	Queen Mary Hosp <b>Adult Cardiac Su</b> Page 5; Ver	rgical Database							
Hospital number   Date of surgery     dd / mm / yyyy									
	Operation data continued	I							
Chest opened by	select from list								
IMA harvested by	select from list								
Vein harvested by	select from list								
Radial artery harvest by	select from list								
Cannulation by	select from list								
Chest closed by	select from list								
Endoscopic vein harvest	<ul><li>O. No</li><li>O. 1. Yes</li></ul>	O 2. Converted							
Endoscopic radial artery harvest	<ul><li>O. No</li><li>O. 1. Yes</li></ul>	O 2. Converted							
Arterial cannulation	<ul><li>O. Not applicable</li><li>1. Ascending aorta</li><li>2. Arch</li></ul>	<ul> <li>3. Axillary / subclavian</li> <li>4. Femoral</li> <li>5. Other</li> </ul>							
Venous cannulation	<ul> <li>0. Not applicable</li> <li>1. Right atrial</li> <li>2. RA/IVC 2-stage</li> </ul>	<ul><li>3. Bicaval</li><li>4. Femoral</li><li>5. Other</li></ul>							
	Procedures classified by g	Iroup							
Cardiac procedures	<ul> <li>1. CABG alone</li> <li>2. CABG &amp; valve</li> <li>3. CABG, valve &amp; other</li> </ul>	<ul> <li>4. CABG &amp; other</li> <li>5. Valve alone</li> <li>6. Valve &amp; other</li> <li>7. Other</li> </ul>							
Other cardiac procedures	<ul> <li>O. None</li> <li>1. LV aneurysmectomy</li> <li>2. Acquired VSD</li> <li>3. Atrial myxoma</li> <li>4. Pulmonary embolect</li> <li>5. Cardiac transplant</li> <li>6. Pulmonary transplant</li> </ul>	12. Atrial ablation							
Other thoracic & vascular procedures	<ul><li>O. None</li><li>1. Aortic</li><li>2. Peripheral vascular</li></ul>	<ul><li>3. Carotid endarterectomy</li><li>4. Other thoracic</li></ul>							



		t Cardiac		long Kong I <b>Databas</b> 1			3
Hospital number				Date of s	urgery	dd / mm /	уууу
	Coronary	artery surg	ery				
Number of DCAs <sup>1</sup>							
	Graft 1	Graft 2	Graft 3	Graft 4	Graft 5	Graft 6	
Graft site	code	code	code	code	code	code	see below
Coronary quality <sup>3</sup>	code	code	code	code	code	code	see below
Coronary lumen at anastomosis	code	code	code	code	code	code	see below
Graft conduit	code	code	code	code	code	code	see below
Conduit quality	code	code	code	code	code	code	see below
Graft anastomosiscodecodeCABG: Graft sites $1 \rightarrow 1$ . Prox RCA $2 \rightarrow 2$ . Mid RCA $3 \rightarrow 3$ . Distal RCA $4 \rightarrow 4$ . RCA-PDA $5 \rightarrow 5$ . RCA-LV $6 \rightarrow 6$ . LMS $7 \rightarrow 7$ . Prox LAD $8 \rightarrow 8$ . Mid LAD $9 \rightarrow 9$ . Distal LAD $10 \rightarrow 10$ . Diag 1 $11 \rightarrow 11$ . Diag 2 $12 \rightarrow 12$ . Prox Cx $13 \rightarrow 13$ . Int $14 \rightarrow 14$ . OM1 $15 \rightarrow 15$ . OM2 $16 \rightarrow 16$ . Distal Cx $15 \rightarrow 17$ . Cx-PDACABG: Coronary quality $1 \rightarrow 1$ . Good $2 \rightarrow 2$ . Moderate/patchyce $3 \rightarrow 3$ . Severe/diffuse diss $4 \rightarrow 4$ . Endarterectomy rest				$1 \rightarrow 1.$ $2 \rightarrow 2.$ $3 \rightarrow 3.$ CABG: Gr $1 \rightarrow 1.$ $2 \rightarrow 2.$ $4 \rightarrow 4.$ $5 \rightarrow 5.$ $7 \rightarrow 7.$ $8 \rightarrow 8.$ $9 \rightarrow 9.$ $11 \rightarrow 11.$ $12 \rightarrow 12.$ CABG: Co $1 \rightarrow 1.$ $2 \rightarrow 2.$ $3 \rightarrow 3.$ CABG: Gr $2 \rightarrow 2.$	1.5-2.0 mm >2.0 mm <b>aft conduits</b> Pedicle LIMA Free LIMA Free RIMA Radial artery Long SV Short SV Other artery Other vein <b>nduit quality</b> Good Moderate	y	
This form is designed so that questio		3. At and be	onary anastom yond the anast	tomosis			



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		e 7; Version 1.1	Dase	8	
Hospital number		Date	of surgery dd / r	mm / yyyy	
Valve surgery					
Number replaced / repaired					
	Aortic	Mitral	Tricuspid	Pulmonary	
Haemodynamic pathology	<ul><li>1. Stenosis</li><li>2. Regurgitation</li><li>3. Mixed</li></ul>	<ul><li>1. Stenosis</li><li>2. Regurgitation</li><li>3. Mixed</li></ul>	<ul><li>1. Stenosis</li><li>2. Regurgitation</li><li>3. Mixed</li></ul>	<ul><li>1. Stenosis</li><li>2. Regurgitation</li><li>3. Mixed</li></ul>	
Native valve pathology	code	code	code	code	
Other native valve pathology	text	text	text	text	
Explant valve type	<ul> <li>1. Native valve</li> <li>2. Mechanical</li> <li>3. Biological</li> <li>4. Homograft</li> <li>5. Autograft</li> <li>6. Ring</li> </ul>	<ul> <li>1. Native valve</li> <li>2. Mechanical</li> <li>3. Biological</li> <li>4. Homograft</li> <li>5. Autograft</li> <li>6. Ring</li> </ul>	<ul> <li>1. Native valve</li> <li>2. Mechanical</li> <li>3. Biological</li> <li>4. Homograft</li> <li>5. Autograft</li> <li>6. Ring</li> </ul>	<ul> <li>1. Native valve</li> <li>2. Mechanical</li> <li>3. Biological</li> <li>4. Homograft</li> <li>5. Autograft</li> <li>6. Ring</li> </ul>	
Reason for repeat valve operation	<ul> <li>1. Thrombosis</li> <li>2. Dehiscence</li> <li>3. Embolism</li> <li>4. Infection</li> <li>5. Intrinsic failure</li> <li>6. Haemolysis</li> <li>19. Other reason</li> </ul>	<ul> <li>1. Thrombosis</li> <li>2. Dehiscence</li> <li>3. Embolism</li> <li>4. Infection</li> <li>5. Intrinsic failure</li> <li>6. Haemolysis</li> <li>19. Other reason</li> </ul>	<ul> <li>1. Thrombosis</li> <li>2. Dehiscence</li> <li>3. Embolism</li> <li>4. Infection</li> <li>5. Intrinsic failure</li> <li>6. Haemolysis</li> <li>19. Other reason</li> </ul>	<ul> <li>1. Thrombosis</li> <li>2. Dehiscence</li> <li>3. Embolism</li> <li>4. Infection</li> <li>5. Intrinsic failure</li> <li>6. Haemolysis</li> <li>19. Other reason</li> </ul>	
Other reason for repeat	text	text	text	text	
Valve procedure	<ul><li>O 1. Replacement</li><li>O 2. Repair</li></ul>	<ul><li>O 1. Replacement</li><li>O 2. Repair</li></ul>	<ul><li>O 1. Replacement</li><li>O 2. Repair</li></ul>	<ul><li>1. Replacement</li><li>2. Repair</li></ul>	
Valve repair procedures	codes	codes	codes	codes	
Valve implant type	<ul> <li>2. Mechanical</li> <li>3. Biological</li> <li>4. Homograft</li> <li>5. Autograft</li> <li>6. Annulopl. ring</li> </ul>	<ul> <li>2. Mechanical</li> <li>3. Biological</li> <li>4. Homograft</li> <li>5. Autograft</li> <li>6. Annulopl. ring</li> </ul>	<ul> <li>2. Mechanical</li> <li>3. Biological</li> <li>4. Homograft</li> <li>5. Autograft</li> <li>6. Annulopl. ring</li> </ul>	<ul> <li>2. Mechanical</li> <li>3. Biological</li> <li>4. Homograft</li> <li>5. Autograft</li> <li>6. Annulopl. ring</li> </ul>	
implant prosthesis name					
implant prosthesis model					
Implant prosthesis serial number					
implant prosthesis size	mm	mm	mm		
Valve surgery: native valve par $0 \rightarrow 0$ . Native valve not present $1 \rightarrow 1$ . Congenital $2 \rightarrow 2$ . Degenerative $3 \rightarrow 3$ . Active infective endocar $4 \rightarrow 4$ . Previous infective endocar $5 \rightarrow 5$ . Rheumatic $6 \rightarrow 6$ . Annuloaortic ectasia $7 \rightarrow 7$ . Calcific degeneration	schaemic Functional regurgitation Other native valve patholog Jery: valve repairs Commisurotomy Annuloplasty (ring) Annuloplasty (suture) .eaflet resection .eaflet extension	9 → 9. Papil	dal shortening cial chord lary muscle repositioning lcification / debridement et patch valvar release ispension		



		en Mary Hos <b>t Cardiac Su</b> Page 8; Ve	irgical D			E
Hospital number				Date of surge	y dd /	mm / yyyy
	Major aor	tic procedure d	lata			
Number of aorta segments						
	Root	Ascending	Arch	Descending	Abdominal	
Aortic pathology	code	code	code	code	code	see below
Aortic procedure	code	code	code	code	code	see below
	4 → 4. Transection 5 → 5. Coarctation 6 → 6. Atheromatous 7 → 7. Marfan's 9 → 9. Mycotic 10 → 10. Other connective tissue disorders 11→ 11. Congenital 12 → 12. Infection - native 13 → 12. Infection - graft 99 → 99. Unknown <b>Major aortic: procedure</b> 1 → 1. Interposition tube graft 2 → 2. Tube graft + separate AVR 3 → 3. Root replacement + composite valve graft & coronary re-implantation 4 → 4. Root replacement + preservation of native valve & coronary re-implantation 5 → 5. Homograft root replacement 6 → 6. Autograft root replacement (Ross procedure) 7 → 7. Aortic patch graft 8 → 8. Sinus of Valsalva repair 9 → 9. Reduction aortoplasty					
Nativo shuthm		he end of the p				
Native rhythm	<ul><li>O 1. Sin</li><li>O 2. Atr</li></ul>	<b>he end of the p</b> us rhythm ial fibrillation / flu	procedure	-	I. Heart block 5. Other	
Native rhythm Pacing	<ul><li>O 1. Sin</li><li>O 2. Atr</li></ul>	<b>he end of the p</b> us rhythm ial fibrillation / flu dal rhythm ne	procedure	0 5		۲
	<ul> <li>1. Sin</li> <li>2. Atr</li> <li>3. No</li> <li>0. No</li> <li>1. Atr</li> <li>0. No</li> </ul>	<b>he end of the p</b> us rhythm ial fibrillation/flu dal rhythm ne ial	orocedure	0 2 0 3	5. Other 2. Ventricular	
Pacing	<ul> <li>1. Sin</li> <li>2. Atr</li> <li>3. No</li> <li>0. No</li> <li>1. Atr</li> <li>0. No</li> </ul>	he end of the p us rhythm ial fibrillation / flu dal rhythm ne ial	orocedure	0 2 0 3	5. Other 2. Ventricular 3. Dual chambe	
Pacing	<ul> <li>1. Sin</li> <li>2. Atr</li> <li>3. No</li> <li>0. No</li> <li>1. Atr</li> <li>0. No</li> <li>1. Low</li> </ul>	he end of the p us rhythm ial fibrillation / flu dal rhythm ne ial ne v dose (<10 ml hr t applicable	orocedure	0 2	5. Other 2. Ventricular 3. Dual chambe	
Pacing	<ul> <li>○ 1. Sin</li> <li>○ 2. Atr</li> <li>○ 3. No</li> <li>○ 0. No</li> <li>○ 1. Atr</li> <li>○ 0. No</li> <li>○ 1. Lov</li> </ul> Closure           ○ 0. No           ○ 1. Atr           ○ 0. No           ○ 1. Atr	he end of the p us rhythm ial fibrillation / flu dal rhythm ne ial ne v dose (<10 ml hr t applicable	orocedure		5. Other 2. Ventricular 8. Dual chambe 2. High dose (>	10 ml hr -1)
Pacing Inotropes Arterial cannulation	<ul> <li>○ 1. Sin</li> <li>○ 2. Atr</li> <li>○ 3. No</li> <li>○ 0. No</li> <li>○ 1. Atr</li> <li>○ 0. No</li> <li>○ 1. Lov</li> </ul> Closure           ○ 0. No           ○ 1. Atr           ○ 0. No           ○ 1. Atr	he end of the p us rhythm ial fibrillation/flu dal rhythm ne ial ne v dose (<10 ml hu t applicable ial diastinal icardial	orocedure		5. Other 2. Ventricular 3. Dual chambe 2. High dose (> 2. Ventricular 3. Left pleural	10 ml hr -1)
Pacing Inotropes Arterial cannulation Drains	<ul> <li>○ 1. Sin</li> <li>○ 2. Atr</li> <li>○ 3. No</li> <li>○ 0. No</li> <li>○ 1. Atr</li> <li>○ 0. No</li> <li>○ 1. Low</li> </ul> Closure Closure <ul> <li>○ 0. No</li> <li>□ 1. Atr</li> <li>□ 1. Me</li> <li>□ 2. Per</li> </ul>	he end of the p us rhythm ial fibrillation/flu dal rhythm ne ial ne v dose (<10 ml hi t applicable ial diastinal icardial utine	orocedure		5. Other 2. Ventricular 3. Dual chambe 2. High dose (> 2. Ventricular 3. Left pleural 4. Right pleural 2. Modified	10 ml hr -1)



Cardioplegia: temperatureI. Cold2. Warm8. Not applicationCardioplegia: infusion mode1. Antegrade2. Retrograde8. Not application	Queen Mary Hospital, Hong Kong Adult Cardiac Surgical Database							
Cardiopulmonary typass       0. No       1. Yes         Conversion to off pump       0. No       1. Yes         Predominant method of myocardial protection       0. Non-cardioplegic       1. Cardioplegia         Cardioplegia: totion       0. Non-cardioplegic       1. Cardioplegia         Cardioplegia: totion       0. Non-cardioplegic       0. Correction         Cardioplegia: totion       0. Non-cardioplegic       0. Correction         Cardioplegia: timing       0. Non-cardioplegic       2. Crystalloid       8. Not applica         Cardioplegia: timing       0. No       0. No       8. Not applica         Cardioplegia: timing       0. No       0. No       8. Not applica         Cardioplegia: timing       0. No       0. No       8. Not applica         Mon-cardioplegia       0. No       0. No       8. Not applica         Mon-cardioplegia       0. No       0. No       8. Not applica         Mon-cardioplegia       0. No       1. Yes       8. Not applica         Mon-cardioplegia       0. No       1. Yes       8. Not applica         Mon-cardioplegia       0. No       1. Yes       8. Not applica         Mategrade highest concentration       1. Aeros clamp and beating heart       3. Post-operation         Intra-aortic bal		Pa	age 9; version 1	.1				
Cardiopulmonary bypass Conversion to off pump Predominant method of myocardial protection Cardioplegia: isolution Cardioplegia: infusion mode Cardioplegia: infusion mode Chest opened by select from list Lowest systemic temperature Chest opened by Select from list Constant constant con	Hospital number			Date	of surgery	dd /	mm / yyyy	
Conversion to of pump Predominant method of myocardial protection Cardioplegia: solution       0       0. No       1. Yes         Cardioplegia: solution Cardioplegia: timusion mode       1. Cold       2. Crystalloid       8. Not applicat         Cardioplegia: infusion mode Cardioplegia: timuing       0       1. Antegrade       2. Retrograde       8. Not applicat         Cardioplegia: timuing       0       1. Intermittent       0       2. Continuous       8. Not applicat         Cardioplegia: timing       0       1. Antegrade       2. Retrograde       8. Not applicat         Mon-cardioplegia       0       0. No       1. Yes       8. Not applicat         Mon-cardioplegia: timing       0       1. Nortic cross clamping with fibrillation       8. Not applicat         Mon-cardioplegia:       0       0. No       1. Yes       9.         Chest opened by       select from list       9.       9.       9.         Lowest systemic temperature       °C       1. Haemodynamic instability       3. CPB wean       3. CPB wean         0       0. No       1. Yes       9.       9.       9.         Intra-aortic balloon pump used       0       1. Haemodynamic instability       3. CPB wean       3. CPB wean         0       1. Ne       0       1. Yes		Cardiopulmonar	y support					
Predominant method of myocardial protection <ul> <li>O</li> <li>O. Non-cardioplegic</li> <li>O</li> <li>Cardioplegia: solution</li> <li>Cardioplegia: solution</li> <li>Cardioplegia: temperature</li> <li>I. Cold</li> <li>Z. Warm</li> <li>S. Not applica</li> </ul> Cardioplegia: infusion mode         I. Antegrade         Z. Retrograde         S. Not applica           Cardioplegia: timing         I. Intermittent         Z. Continuous         S. Not applica           Mot shot         O         O. No         I. Yes         S. Not applica           Mon-cardioplegia         I. Anotic cross clamping with fibrillation         S. Cross clamp and beating heart         S. Continuous         S. Not applica           Non-cardioplegia         I. Anotic cross clamp and beating heart         S. Constant preduction         S. Cross clamp         S. Constant preduction         S. Cross clamp         S. Cross clamp <th>Cardiopulmonary bypass</th> <th>O 0. No</th> <th>0</th> <th>1. Yes</th> <th></th> <th></th> <th></th>	Cardiopulmonary bypass	O 0. No	0	1. Yes				
myocardial protection Cardioplegia: solution       0       0. Non-cardioplegia       0       2. Crystalloid       0       8. Not applica         Cardioplegia: temperature       1. Cold       2. Warm       0       8. Not applica         Cardioplegia: temperature       1. Antegrade       2. Retrograde       0       8. Not applica         Cardioplegia: timing       0       1. Intermittent       2. Continuous       0       8. Not applica         Kono-cardioplegic       0       1. Antegrade       2. Continuous       0       8. Not applica         Non-cardioplegic       0       1. Antic cross clamping with fibrillation       0       8. Not applica         Non-cardioplegic       0       1. Aortic cross clamp and beating heart       0       3. Cross clamp and beating heart       0       3. Cross clamp and beating heart       0       0. No       0       1. Pre-operation       1       2. Intra-operation       1 <td< th=""><th></th><th>O 0. No</th><th>0</th><th>1. Yes</th><th></th><th></th><th></th></td<>		O 0. No	0	1. Yes				
Cardioplegia: temperature <ul> <li>1. Cold</li> <li>2. Warm</li> <li>8. Not applica</li> </ul> Cardioplegia: timing <ul> <li>1. Antegrade</li> <li>2. Retrograde</li> <li>8. Not applica</li> </ul> Cardioplegia: timing <ul> <li>1. Intermittent</li> <li>2. Continuous</li> <li>8. Not applica</li> </ul> Mon-cardioplegia: timing <ul> <li>1. Aortic cross clamping with fibrillation</li> <li>2. Fibrillation with perfusion</li> <li>3. Cross clamp and beating heart</li> <li>5. Beating heart without cross clamp</li> </ul> Chest opened by         select from list           Lowest systemic temperature <ul> <li>Cross clamp and beating heart</li> <li>5. Beating heart without cross clamp</li> </ul> Intra-aortic balloon pump used <ul> <li>0. No</li> <li>1. Pre-operation</li> <li>3. Post-operation</li> </ul> Intra-aortic balloon pump used <ul> <li>1. Pre-operation</li> <li>3. Post-operation</li> <li>1. Pre-operation</li> <li>3. Post-operation</li> <li>4. Prophylactic</li> </ul> Intra-aortic balloon pump used <ul> <li>0. No</li> <li>1. Yes</li> <li>4. Prophylactic</li> </ul>		O 0. Non-cardic	oplegic O	1. Cardio	oplegia			
Cardioplegia: infusion mode       1. Antegrade       2. Retrograde       8. Not application and the second secon	Cardioplegia: solution	O 1. Blood	0	2. Crysta	lloid	0	8. Not applicable	
Cardioplegia: timing 1. Intermittent 2. Continuous 8. Not applied   Hot shot 0. No 1. Yes   Non-cardioplegia: myocardial protection 1. Aortic cross clamping with fibrillation 0. 2. Fibrillation with perfusion 0. 3. Cross clamp and beating heart 0. 5. Beating heart without cross clamp 1. Yes   Chest opened by select from list   Lowest systemic temperature °C   Antegrade highest concentration 3. Cross clamp with direct coronary perfusion 0. 8. Not applied 0. Select from list   Intra-aortic balloon pump used 0. No   0 0. No   1. Haemodynamic instability 3. CPB wean 0. 2. Unstable angina   0 0. No   1. Haemodynamic instability 3. CPB wean 0. 2. Unstable angina   0 0. No   1. Haemodynamic instability 3. CPB wean 0. 1. Yes   0 0. No   1. Haemodynamic instability 3. CPB wean 0. 1. Yes   0 0. No   1. Haemodynamic instability 3. CPB wean 0. 1. Yes   0 0. No   1. Haemodynamic instability 1. Yes   0 0. No   1. Yes   0 0. No   1. Yes   1. Heighti	Cardioplegia: temperature	1. Cold		2. Warm		0	8. Not applicable	
Hot shot       0. No       1. Yes         Non-cardioplegic myocardial protection       0. Aortic cross clamping with fibrillation       0. So         0. S. Cross clamp and beating heart       0. So cross clamp and beating heart       0. So cross clamp and beating heart         0. S. Beating heart without cross clamp       S. Beating heart without cross clamp       0. No         Chest opened by       select from list         Lowest systemic temperature       °C         Intra-aortic balloon pump used       0. No       1. Pre-operation         1. Haemodynamic instability       3. CPB wean         0. 2. Unstable angina       4. Prophylactic         IABP serial number       0. No       1. Yes         Volume filtered       0. No       1. Yes         Filtration       0. No       1. Yes         Filtration       0. No       1. Yes         Weight       Com       1. Yes	Cardioplegia: infusion mode	1. Antegrade		2. Retrog	grade	0	8. Not applicable	
Non-cardiopleginmyocardial protection <ul> <li>1. Aortic cross clamping with fibrillation</li> <li>2. Fibrillation with perfusion</li> <li>3. Cross clamp and beating heart</li> <li>5. Beating heart without cross clamp</li> </ul> Chest opened by           Schest opened by           Select from list           Lowest systemic temperature           Intra-aortic balloon pump used           0         0. No           1. Pre-operation         3. CPB wean           0         2. Unstable angina         4. Prophylactic           IABP serial number <ul> <li>2. Unstable angina</li> <li>3. Post-operation</li> <li>3. Post-operation</li> <li>3. Post-operation</li> <li>3. Post-operation</li> <li>4. Prophylactic</li> </ul> <li>IABP serial number</li> <li>2. Unstable angina</li> <li>4. Prophylactic</li> <li>3. CPB wean</li> <li>0. No</li> <li>1. Yes</li> <li< th=""><th>Cardioplegia: timing</th><th>O 1. Intermitter</th><th>nt O</th><th>2. Contir</th><th>nuous</th><th>0</th><th>8. Not applicable</th></li<>	Cardioplegia: timing	O 1. Intermitter	nt O	2. Contir	nuous	0	8. Not applicable	
myocardial protection <ul> <li>2. Fibrillation with perfusion</li> <li>3. Cross clamp and beating heart</li> <li>4. Cross clamp and beating heart</li> <li>5. Beating heart without cross clamp</li> </ul> Chest opened by select from list   Lowest systemic temperature <ul> <li>C</li> </ul> Antegrade highest concentration Intra-aortic balloon pump used 0 0. No 1. Haemodynamic instability 3. CPB wean 3. CPB wean 3. CPB wean 4. Prophylactic 3. CPB wean 4. Prophylactic 3. CPB wean 4. Prophylactic 1. Haemodynamic instability 3. CPB wean 3. CPB wean 4. Prophylactic 1. Haemodynamic instability 3. CPB wean 4. Prophylactic 1. Haemodynamic instability 3. CPB wean 4. Prophylactic 1. Haemodynamic instability 3. CPB wean 3. CPB wean 4. Prophylactic 1. Haemodynamic instability 3. CPB wean 4. Prophylactic 1. Haemodynamic instability 3. CPB wean 4. Prophylactic 1. Height 6. 0. No 1. Yes	Hot shot	O 0. No	0	1. Yes				
Lowest systemic temperature Covernment of the systemic temperature Covernment of temperature Covernment of temperature Covernment of te		<ul> <li>2. Fibrillation with perfusion</li> <li>3. Cross clamp with direct coronary perfusion</li> <li>4. Cross clamp and beating heart</li> </ul>						
Antegrade highest concentration   Intra-aortic balloon pump used   0   0   0   1   Pre-operation   3   Post-operation   3   C   1   1   Pre-operation   3   2   1   1   Pre-operation   3   2   1   1   Pre-operation   3   2   1   1   1   1   Pre-operation   3   2   1 <th>Chest opened by</th> <th>select from list</th> <th>t</th> <th></th> <th></th> <th></th> <th></th>	Chest opened by	select from list	t					
Intra-aortic balloon pump used <ul> <li>0. No</li> <li>1. Pre-operation</li> <li>3. Post-operation</li> <li>3. Post-operation</li> </ul> Reason for IABP use <ul> <li>1. Haemodynamic instability</li> <li>3. CPB wean</li> <li>4. Prophylactic</li> </ul> IABP serial number <ul> <li>2. Unstable angina</li> <li>4. Prophylactic</li> </ul> Date IABP removed              dd/mm/yyyy             Transamin              0. No              1. Yes           Novo 7              0. No              1. Yes           Filtration              0. No              1. Yes               Volume filtered              ml               Height              ccm               Weight              kg               Cumulative bypass time              min	Lowest systemic temperature		°C					
Image: Second for IABP use 1. Pre-operation 3. Post-operation   Reason for IABP use 1. Haemodynamic instability 3. CPB wean   2. Unstable angina 4. Prophylactic   IABP serial number dd/mm/yyyy   Transamin 0. No 1. Yes   Novo 7 0. No 1. Yes   Filtration 0. No 1. Yes   Wolume filtered ml   Height cm   Weight kg   Cumulative bypass time min	ntegrade highest concentration							
O 2. Unstable angina O 4. Prophylactic   IABP serial number dd/mm/yyyy   Date IABP removed dd/mm/yyyy   Transamin O 0. No   1. Yes O 1. Yes   Novo 7 O 0. No   Filtration O 0. No   Volume filtered ml   Height cm   Weight kg   Cumulative bypass time min	Intra-aortic balloon pump used	· · · · · · · · · · · · · · · · · · ·	ion					
Date IABP removed   Image: dd/mm/yyyy   Transamin   Image: Dolored image: Do	Reason for IABP use							
TransaminO0. NoO1. YesNovo 7O0. NoO1. YesFiltrationO0. NoO1. YesVolume filteredImlImlHeightcmcmWeightkgImlCumulative bypass timeIml	IABP serial number							
Novo 7 O 0. No O 1. Yes   Filtration O 0. No O 1. Yes   Volume filtered ml   Height cm   Weight kg   Cumulative bypass time min	Date IABP removed	dd/mm/yyyy	/					
Filtration     O     0. No     O     1. Yes       Volume filtered     ml       Height     cm       Weight     kg       Cumulative bypass time     min	Transamin	O 0. No			O 1. Ye	S		
Volume filtered     ml       Height     cm       Weight     kg       Cumulative bypass time     min	Novo 7	O 0. No			O 1. Ye	S		
HeightcmWeightkgCumulative bypass timemin	Filtration	O 0. No			O 1. Ye	S		
Weight     kg       Cumulative bypass time     min	Volume filtered		ml					
Cumulative bypass time min	Height		cm					
	Weight		kg					
Cumulative cross clamp time min	Cumulative bypass time	min						
	Cumulative cross clamp time		min					
Total circulatory arrest time min	Total circulatory arrest time		min					



Queen Mary Hospital, Hong Kong Adult Cardiac Surgical Database Page 10; Version 1.1							
Hospital number		Date of	surgery	dd / mm / yyyy			
	Cardiopulmonary	y support continued					
Cerebral perfusion during HCA	<ul><li>O. None</li><li>O. Antegrade</li></ul>		O 2. R	etrograde			
Cell salvage used	O 0. No		O 1. Ye	25			
Volume heparinized saline		ml					
Other volume (blower mister etc)		ml					
Blood from circuit		ml					
Volume processed		ml					
Volume re-infused		ml					
Perfusion notes							
	Blood products u	sed					
Blood		units					
Platelets		units					
FFP		units					
Cryoprecipitate		units					
	Post-operative co	ourse (CCU)					
PA catheter	O 0. No		O 1. Ye	25			
Inotropes	<ul> <li>O. None</li> <li>1. Dopamine</li> <li>2. Dobutamin</li> <li>3. Adrenaline</li> </ul>		5. Va	oradrenaline asopressin Iilrinone noxamine			
Inotropes >5 ml hour <sup>-1</sup>	O 0. No		O 1. Ye	25			
Vasoconstrictor >5 ml hour <sup>-1</sup>	O 0. No		O 1. Ye	25			
Chest drainage (first 24 hours)		ml					
Date of discharge from CCU	dd/mm/yyyy						



	Queen Mary Hospital, Hong F Adult Cardiac Surgical Data Page 11; Version 1.1	-		
Hospital number	Dat	e of surgery dd / mm / yyyy		
Post-operative course				
Post-operative complications	O 0. No	O 1. Yes		
Re-admission to CCU	O 0. No	O 1.Yes		
Return to theatre	<ul> <li>0. No re-operation necessary</li> <li>1. Re-operation for bleeding or tampon</li> <li>2. Re-operation for valvular problems</li> <li>3. Re-operation for graft problems</li> <li>4. Re-operation for other cardiac probl</li> <li>5. Sternum resuturing (sterile)</li> <li>6. Surgery for deep sternal wound infer</li> </ul>	ems		
Arrhythmias requiring intervention	<ul> <li>0. None</li> <li>1. Atrial fibrillation / flutter</li> <li>2. VT</li> </ul>	<ul> <li>3. VF</li> <li>4. Heart block</li> <li>5. Other</li> </ul>		
Intervention	<ul><li>1. Pharmocological</li><li>2. Electrical cardioversion</li></ul>	<ul><li>3. Permanent pacemaker</li><li>4. Other</li></ul>		
Secondary airway support	<ul><li>O. None</li><li>1. Mini-tracheostomy</li><li>2. Facial CPAP</li></ul>	<ul><li>3. Re-intubation</li><li>4. Tracheostomy</li></ul>		
Pulmonary complications requiring intervention	<ul><li>O. None</li><li>1. Chest infection</li><li>2. Pleural effusion</li></ul>	<ul><li>3. Pneumothorax</li><li>4. Pulmonary embolus</li><li>5. Other</li></ul>		
Infective complications	<ul> <li>O. None</li> <li>1. Superficial sternal</li> <li>2. Deep sternal / mediastinal</li> <li>3. Pulmonary</li> </ul>	<ul> <li>4. Leg or arm wound</li> <li>5. Septicaemia</li> <li>6. Other</li> </ul>		
Post-operative fever	O 0. No	O 1.Yes		
Gastro-intestinal complications	<ul> <li>O. None</li> <li>1. GI bleed</li> <li>2. Perforated peptic ulcer</li> <li>3. lschaemic bowel</li> </ul>	<ul><li>4. Pancreatitis</li><li>5. Ileus requiring intervention</li><li>6. Other</li></ul>		
Renal impairment	O 0. No	O 1. Yes		
New HF / dialysis post-operatively	O 0. No	O 1. Yes		
Renal replacement therapy	O 0. No	O 1. Yes		
Type of renal replacement therapy	<ul><li>1. Peritoneal dialysis</li><li>2. CWH</li></ul>	3. HD		
Peak post-operative creatinine	µmol ℓ⁻¹			
New post-operative stroke	<ul><li>O. None</li><li>O. 1. Yes (prophylatic)</li></ul>	O 2. Yes (clinically indicated)		
Post-operative antibiotics	<ul><li>O. None</li><li>O. 1. Transient stoke</li></ul>	O 2. Permanent stroke		
Complication notes				



Queen Mary Hospital, Hong KongAdult Cardiac Surgical DatabasePage 12; Version 1.1				
Hospital number		Date of surgery	dd / mm / yyyy	
	Discharge			
Pre-discharge haemoglobin	g dℓ-¹			
Pre-discharge creatinine	µmol ℓ⁻¹			
Aspirin	<ul><li>O. Not given</li><li>O. 1. Given</li><li>O. 2. Contra-indicated</li></ul>		Other antiplatelet given Unknown	
Statin	<ul><li>O. Not given</li><li>O. 1. Given</li></ul>	· · · · · · · · · · · · · · · · · · ·	Contra-indicated Unknown	
Warfarin	O 0. No	O 1.1	Yes	
Discharge destination from cardiothoracic ward	<ul> <li>O 1. Home</li> <li>O 2. Convalescence (Non ad O 3. Other hospital</li> </ul>		Not applicable - patient deceased Other specialty	
Patient status at discharge	<ul><li>O. Alive</li><li>O. 1. Dead</li><li>O. 2. Dead (theatre)</li></ul>	O 4.	Dead (ICU) Dead (cardiothoracic ward) Dead (other wards / hospital)	
Date of discharge from CTS	dd/mm/yyyy			
Date of discharge / death	dd/mm/yyyy			



## QMH Score - Components of the additive model

Risk Factor		Score
Age		
	<60	0
	60-64	2
	65-69	2.5
	70-74	3
	>74	4
Renal failure		4
EF<30%		3
Pre-operative Cardiac Conditions		2
Transmyocardial infarction <48hrs		3.5
Congestive cardiac failure		1.5
Endocarditis		2.5
Pulmonary hypertension		2.5
Redo operation		2
Emergency surgery		2
Valves & CABG		2

Total \_\_\_\_\_



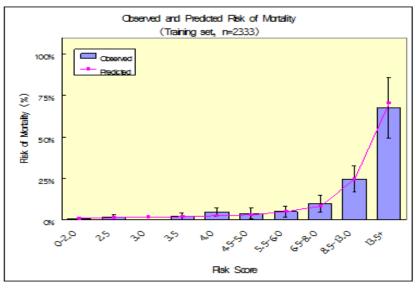
## Predicted risk of mortality with individual Risk Scores and the distribution of Total Risk Score among patients of Training set (n=2333)

Total Risk Score	Predicted Risk	No. of Patients	Cumulative % of Patients with this risk score or less	Total Risk Score	Predicted Risk	No. of Patients	Cumulative % of Patients with this risk score or less
0	0.47%	426	18.26%	10	23.42%	18	97.34%
1.5	0.87%	319	31.93%	10.5	27.37%	8	97.69%
2	1.07%	182	39.73%	11	31.72%	6	97.94%
2.5	1.31%	199	48.26%	11.5	36.41%	6	98.20%
3	1.61%	182	56.07%	12	41.37%	7	98.50%
3.5	1.98%	195	64.42%	12.5	46.51%	7	98.80%
4	2.43%	271	76.04%	13	51.73%	3	98.93%
4.5	2.98%	106	80.58%	13.5	56.91%	6	99.19%
5	3.64%	24	81.61%	14	61.94%	4	99.36%
5.5	4.45%	83	85.17%	14.5	66.73%	0	99.36%
6	5.43%	79	88.56%	15	71.20%	6	99.61%
6.5	6.61%	68	91.47%	15.5	75.29%	2	99.70%
7	8.03%	18	92.24%	16	78.97%	1	99.74%
7.5	9.71%	25	93.31%	16.5	82.23%	1	99.79%
8	11.70%	21	94.21%	17	85.08%	2	99.87%
8.5	14.04%	30	95.50%	17.5	87.55%	1	99.91%
9	16.76%	8	95.84%	18	89.65%	0	99.91%
9.5	19.88%	17	96.57%	18.5+	>90.00%	2	100.00%

\*The highest observed total risk score was 18.5, and there were no patients had total risk score of 14.5 and 18.

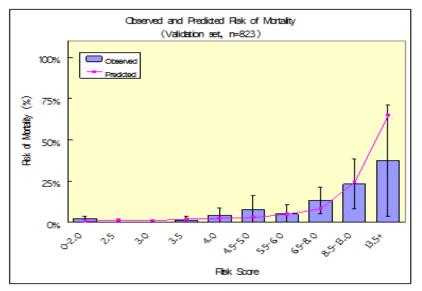


### Graph of a comparison of Predicted and Observed mortality (Training set, n=2333)



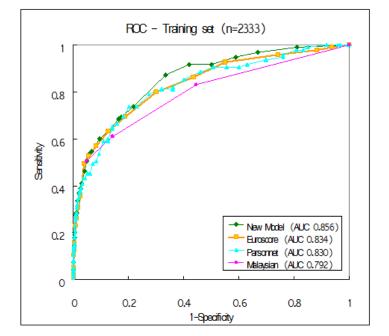
\*Narrow vertical bars depict the 95% CI for each observed value

## Graph of a comparison of Predicted and Observed mortality (Validation set, n=823)



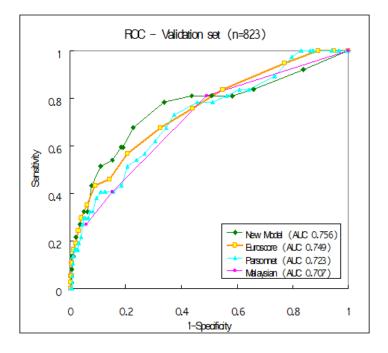
<sup>\*</sup>Narrow vertical bars depict the 95% CI for each observed value





### **Comparing different Model ROC – Training Set**

### **Comparing Model ROC – Validation Set**





Hong Kong Med J 2009 Jun;15(3):173-8

WK Au; KT Lam; LC Cheng; SW Chiu Division of Cardiothoracic Surgery, Department of Surgery, University of Hong Kong, Grantham Hospital, Hong Kong

**Objective:** To determine the impact of diabetes on early and mid-term survival in the Hong Kong Chinese population undergoing coronary artery bypass graft surgery.

**Design.** Prospective study.

Setting. Regional hospital, Hong Kong.

**Patients.** A total of 904 consecutive patients following coronary artery bypass graft surgery from November 1999 to December 2003 were prospectively analysed. Among them, 377 (42%) diabetic and 527 (58%) non-diabetic patients were evaluated.

**Main outcome measures.** Hospital mortality, mid-term mortality, and percutaneous coronary intervention–free survival.

**Results:** The diabetic group had a higher risk score than the non-diabetic group (mean+/-standard deviation: EuroScore 4.7+/-3.4 and 3.6+/-3.4, respectively; P<0.001). Hospital mortality was 3.4% in the diabetic group compared to 2.9% in the non-diabetic group (P=0.37). Multiple logistic regression analysis identified left ventricular ejection fraction of less than 30% and preoperative intubation as independent risk factors for early hospital death. There were 81 late deaths and the actuarial survival at 48 months for the diabetic and non-diabetic patients were 86% and 90%, respectively (P=0.298). The angina-free survival and percutaneous coronary intervention–free survival at 48 months for the diabetic and non-diabetic patients were 86% and 90%, respectively (P=0.298). The angina-free survival and percutaneous coronary intervention–free survival at 48 months for the diabetic and non-diabetic patients yielded no statistically significant difference.

**Conclusions:** Diabetes mellitus was not a predictor of early and midterm mortality after coronary artery bypass graft surgery in our Chinese



population. Furthermore, diabetes did not affect angina recurrence or intervention free–survival up to 4 years.

### Key words

Coronary artery bypass; Coronary disease; Diabetes mellitus; Survival analysis



## Mortality prediction in adult cardiac surgery patients: comparison of two risk stratification models.

Hong Kong Med J 2007;13:293-7

WK Au, FRCS FHKAM; SR Das2, FFARCSI FHKAM; MP Sun, BSc (RN); LC Cheng, FRCS FHKAM; SW Chiu, FRCS FHKAM. Division of Cardiothoracic Surgery, Department of Surgery, University of Hong Kong, Grantham Hospital, Hong Kong 2Department of Anaesthesia, Grantham Hospital

**Objective:** To assess and compare the two commonly applied models, EuroSCORE model and the Parsonnet model in our local adult cardiac surgery patients by quantifying risk factors and relating them to mortality using risk stratification protocol in order to assess the quality of cardiac surgical care.

**Design:** Prospective study.

**Patients:** All adult patients undergoing Coronary Artery Bypass Graft Surgery (CABG) and Heart Valve Surgery at the Grantham Hospital were collected prospectively since November 1999.

**Main outcome measures:** In-hospital mortality was the defined end-point. Statistical analysis consisted of observed vs. expected mortality, Hosmer-Lemeshow goodness-of-fit test for calibration accuracy and Receiver-Operating-Characteristic (ROC) curve for discrimination performance.

**Results:** From November 1999 to July 2005, 2653 patients underwent either CABG (1247) or heart valve surgery (1406). Observed mortalities in CABG and valve surgery patients in the study were 2.9% and 4.8% respectively. The expected mortalities of CABG and valve surgery patients as predicted by EuroSCORE were 4.0 +/- 3.3% and 5.2 +/- 3.0% respectively and by the Parsonnet model were 5.9 +/- 4.2 % and 7.3 +/- 4.4 % respectively. EuroSCORE performed better than the Parsonnet model at predicting inhospital mortality assessed by Hosmer-Lemeshow goodness-to-fit test.



Area under the ROC curves in CABG surgery was: EuroScore 0.76, Parsonnet 0.74 and ROC curve areas in valves surgery was: EuroScore 0.77, Parsonnet 0.79.

**Conclusion:** Despite significant geographic and demographic differences between European and Asian patients, EuroScore performed well with good calibration and discrimination in predicting mortality in our local adult cardiac surgery patients. There was a tendency for both models to over predict. However, EuroSCORE can serve as a baseline for the development of a local risk model.

#### Key words:

Risk stratification, cardiac surgery, Asian, EuroSCORE, prospective



## Predicting major early surgical morbidity and intensive care length of stay (ICLOS) after adult cardiac surgery using the EuroSCORE.

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**Introduction:** The ability to predict early morbidity and mortality after major cardiac surgery is known to be useful in many areas when providing quality surgical care. The European System for Cardiac Operative Risk Evaluation (EuroSCORE) has been widely validated for predicting in-hospital mortality. Few studies however exist that evaluate the performance of the EuroSCORE in predicting early morbidity. The aim of this study is to evaluate the performance of EuroSCORE in predicting major in-hospital morbidity and the need for prolonged intensive care after cardiac surgery.

**Methods:** 745 consecutive adult cardiac surgical patients who underwent CABG and/or valve surgery at the Grantham Hospital from July 2004 till December 2005 were included in this study. EuroSCOREs, predefined postoperative in-hospital major morbidity and ICLOS data were collected prospectively and analyzed using SPSS 11. The ability of the EuroSCORE to discriminate outcomes was assessed using C statistic (area under receiver operating characteristic curve) while the calibration was evaluated by the Hosmer-Lemeshow goodness-of-fit statistic.

**Results:** EuroSCORE showed good discriminatory ability and good calibration in predicting prolonged ICLOS > 3 days (C statistic: 0.713, Hosmer-Lemeshow: p = 0.43), need for ventilatory support > 2 days (C statistic: 0.829, Hosmer-Lemeshow: p = 0.2), and new postoperative renal failure (C statistic: 0.789, Hosmer-Lemeshow: p = 0.3). Its ability to predict the development of postoperative strokes, myocardial infarction, major



sepsis (pneumonia, septicaemia or mediastinitis), and need to reopen for bleeding and/or tamponade were however unsatisfactory.

**Conclusions:** EuroSCORE can be used to predict prolonged ICU stay and specific morbidities such as the development of new post-operative renal failure and prolonged ventilatory support. And this has clinical implications in terms of providing more detail informed consent, quality of care assessment and hospital resources allocation.



# A simple new risk stratification model for adult cardiac surgery in Hong Kong.

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**Objective:** Significant geographic and demographic differences are obvious between European and Asian patients in adult cardiac surgery. Western population-based risk stratification models may not be optimal for local application. This study aims to develop a simple risk model to predict the in-hospital mortality for patients undergoing coronary artery bypass graft surgery (CABG) and valves surgery in a single institution.

**Patients and methods:** Between Nov 1999 to May 2006, pre-operative risk factors and in-hospital mortality of 3156 adults undergoing CABG and valves surgery were prospectively collected for analyses. The new risk model was derived from multivariate logistic regression. The discrimination performance was evaluated using Receiver-Operating-Characteristic (ROC) curve.

**Results:** The overall in-hospital mortality was 4.18%. The risk model included 11 factors, in order of importance (all P<0.01), age, renal failure, transmyocardial infarction < 48 hours, left ventricular ejection fraction < 30%, active endocarditis, pulmonary hypertension, redo operation, emergency surgery, combined valve and CABG, critical pre-operative status and congestive heart failure. The new risk model exhibited good discriminated between high- and low-risk patients (ROC curve area 0.856).

**Conclusion:** This is the first local risk model that predicts in-hospital mortality for adult cardiac surgery in Hong Kong. This model has been



evaluated with good discrimination accuracy. This simple additive non-linear model can provide a useful tool for patient consent and institutional quality assessment. Application of this new model to other local and regional institutions shall be carrying out for further validation.

#### Key words:

Risk stratification, cardiac surgery, new model, EuroSCORE, prospective

