

Health NZ



**New Zealand Cardiac Surgery
National Report**

2017

Preface

New Zealand Cardiac Surgical Annual Report 2017

Patients undergoing cardiothoracic surgery are some of the most ill patients that our health system cares for. Suffering conditions such as heart valve disease and blocked coronary arteries. All cardiothoracic surgeons have professional responsibilities towards patients, the public, their colleagues and employers, and the wider surgical profession, and to that end Cardiothoracic surgeons have championed the collection and publication of clinical outcomes data and since monitoring and publishing survival rates for adult cardiac surgery in New Zealand in the past 3 years; cardiothoracic units can now benchmark performance against the national average, a powerful tool for continued improvements in services to patients.

This report has been prepared by Dendrite Clinical Systems in conjunction with the National Cardiothoracic Surgery Clinical Network (NCSCN). It is based on information collected between 1 January 2017 and 31 December 2017, and presents performance and outcomes data on the most common, publicly-funded cardiac surgery procedures; mainly isolated coronary artery bypass grafting (CABG) and isolated aortic valve replacements (AVR).

The outcome of a cardiac surgical procedure is an example of teamwork. There are many factors such as the patient's general condition; the other medical staff (cardiologists, anaesthetists, intensivists and junior medical staff); the post-operative care (provided by nurses, physiotherapists, pharmacists etc.); and the hospital facilities (which impact infection rates, physical plant) that can have a bearing on the surgical outcome. Many of these factors are simply outside the day-to-day control of the Consultant Surgeon.

The sporting cliché *There is no 'I' in team* should be borne in mind when discussing hospital-specific data. We have used crude mortality, unadjusted for case mix, because our dataset is not yet big enough to apply case mix adjustment to every hospital practice.

Our patients should be reassured by the information presented within this report. Our national results compare very favourably with international standards and continue to improve.

The last 12 months has seen further progress towards improving cardiac services in New Zealand. There has been major investment nationally to improve the clinical settings in which our patients are treated, and cardiac surgery waiting times have continued to improve. Relatively fewer patients wait more than 12 weeks now for open heart surgery. Another point that is important to highlight is the extremely high standard of care that is being offered in this country. Despite ever-increasing complexity of case mix, the mortality rates here compare favourably with those elsewhere in the world. This is a tribute to the high standards of training being made available in New Zealand and to the commitment to quality that Cardiothoracic Surgeons promulgate.

Mr Adam El Gamel

Chair, National Cardiothoracic Surgery Clinical Network



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Introduction

The New Zealand National Cardiac Surgery Registry (NZCSR) has been established by the New Zealand cardiac surgical community as a quality assurance tool that will enable us to audit our practice, review surgical outcomes and to compare these between units in New Zealand and also to benchmark against internationally reported standards. This report is the analysis of the year 2017 of patient enrolment in all 5 cardiac surgical units in New Zealand (Auckland, Waikato, Capital and Coast, Canterbury, Southern).

At an individual Surgeon and unit level NZCSR facilitates analysis of work patterns and ensures that key performance indicators are met. It is used for regular multi-disciplinary discussion of individual patient morbidity and mortality as part of already well established peer review and audit processes. On a national level NZCSR facilitates comparison of regional variation in surgical work load, patient characteristics, risk profiles, comparison of outcomes and will better inform national planning for current and future population and individual patient needs. With time the database will mature to become an important resource for ongoing improvement of patient care and also to facilitate the implementation of quality improvement projects. It will help us plan for national variations in healthcare needs of our population and to ensure equitable access to surgical treatment across the regions. The database is a tool for surgeons, cardiac surgical units and the New Zealand community to assess surgical intervention and outcomes across the country and to ensure the highest standard of care to our patients.

All data is collated and analysed independently by Dendrite Clinical Systems, an internationally respected specialist supplier of clinical database and analysis software. The project is overseen by the National Cardiac Surgery Clinical Network which comprises members from each of the 5 public cardiac surgical units in New Zealand along with members from the Ministry of Health, the National Cardiac Network and community representatives. We aim to provide a patient focused, accurate and transparent report of outcomes for cardiac surgery in New Zealand. The 2 most common category of operation performed in New Zealand are reported: Isolated Coronary Artery Bypass Grafting (CABG) and Isolated Aortic Valve replacement (AVR). Volume of procedure, patient characteristics, morbidity and mortality indicators of resource utilisation are presented. These 2 groups combined make up approximately 65% of all cardiac surgery performed in New Zealand and are reflective of national surgical practice and results. Outcomes for individual patients are heavily influenced by factors such as overall health, age, co-existing medical conditions, acuity and magnitude of surgery. Therefore, major outcomes such as mortality will be risk adjusted using internationally validated and accepted risk scoring tools. Also we will compare outcomes in New Zealand by benchmarking against other internationally reported cardiac surgical registries. In comparison to these other registries New Zealand is a small surgical community, to ensure that reporting of outcomes does not reflect statistically insignificant variation we aim to produce a National yearly report. Ultimately our goal is to provide the highest standard of medical and surgical care to the population of New Zealand and to continue to reflect on and to improve our practice for the good of our patients.

Over the coming years as the registry grows we expect it will form the framework for development and ongoing reporting of a number of quality improvement programmes.

Finally, it is important to stress that a cardiac surgical team is an extensive one and numerous medical professionals' support and provide care to each individual patient through their journey. Whilst the operation is ultimately the largest intervention undertaken it is important to stress that each of the medical professions involved (cardiologist, surgeon, perfusionist, intensive care specialist, anaesthetist, junior doctor, nurse, social worker, physiotherapist, pharmacist and occupational therapist) play an important role in the care provided to, and the outcomes for each individual patient. When we report outcomes these are collectively shared by all members of the team. The development of a robust and accurate database allows us to identify where the team is doing well but also where there is room for the team to improve. The national database is supported by a rigorous governance structure, each individual surgeon maintains professional development and practice audit in keeping with standards set by the New Zealand Medical Council (NZMC), The Australasian Society of Cardiac and Thoracic Surgeons (ANZCTS) and The Royal Australasian College of Surgeons (RACS). Whilst the database and our regulatory bodies (NZMC, RACS) have processes in place to identify and further assess underperforming individuals an important aspect of a national report is that it remains confidential at an individual surgeon and patient level. In reporting unit results we are acknowledging that the outcomes presented are not just attributable to individuals but are a product of the collaboration between and the contributions made by all members of the cardio surgical team.

Mr Sean Galvin

on behalf of the National Cardiothoracic Surgery Clinical Network (NCSCN)



Data presentation

- The data presented within this report is for the period 1 January 2017 to 31 December 2017.
- It includes all public funded cardiac surgical procedures performed nationally.
- In this report we have analysed the risk factors and their impact on outcomes.
- The two standardised operations included are coronary artery bypass grafting (CABG) and aortic valve replacement (AVR) these account for over 65% of the workload of all cardiac surgical units.
- The data has been collected using Dendrite Clinical Systems clinical database.
- The definitions used in this database have been aimed to be identical with international definitions so a realistic comparison can be made with other international standards.

Overview of people who had cardiac surgery

In the 12-month period (2017) a total of 2,727 cardiac surgical procedures were performed across the five publicly-funded cardiac surgery centres ¹ (Table 1).

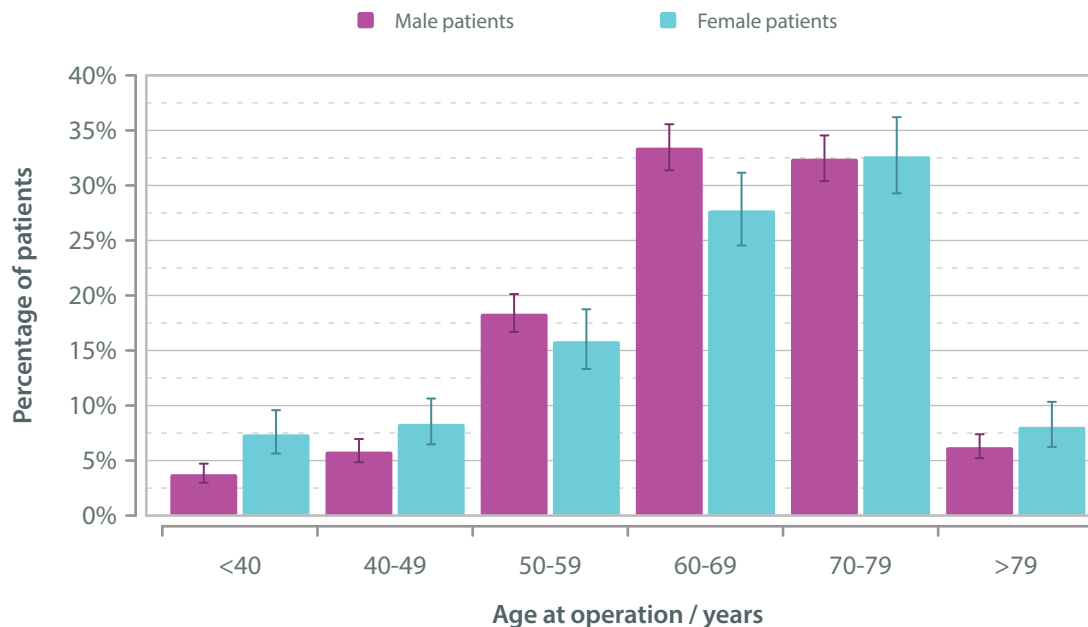
70% of cardiac surgical procedures were for patients aged 60 years or over with 73% of the total number of patients male (Fig. 1).

Table 1. All cardiac surgery patients in 2017: age and gender

Age at operation / years	Gender		All
	Male	Female	
<40	75	54	129
40-49	116	61	177
50-59	366	116	482
60-69	667	203	870
70-79	647	239	886
>79	124	59	183
Unspecified	0	0	0
All	1,995	732	2,727

Fig. 1

All cardiac surgery patients: Age & gender distributions; calendar year 2017 (n=2,727)



1. The 5 District Health Boards (DHB) undertaking Cardiac surgery: Auckland DHB, Waikato DHB, Capital and Coast DHB, Canterbury DHB, Southern DHB.



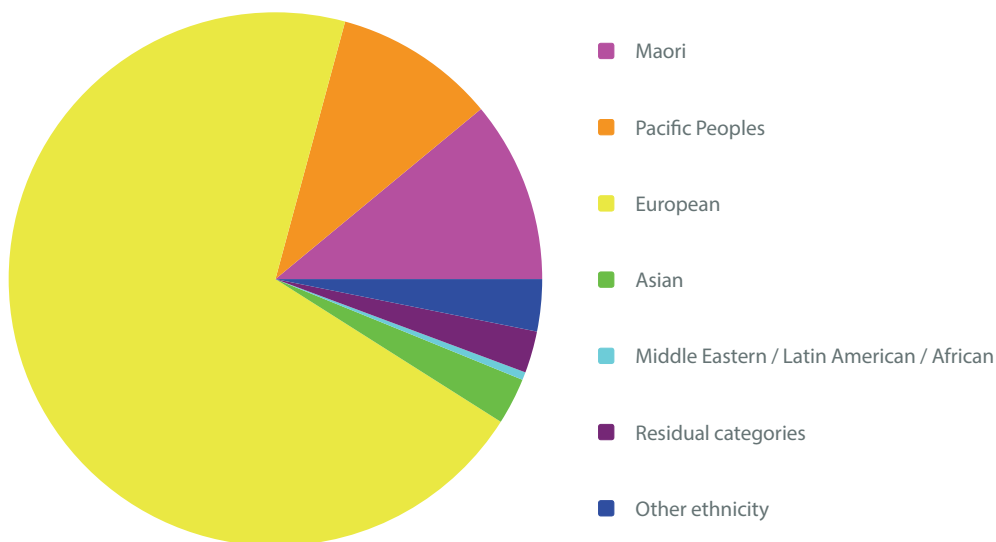
Ethnicity

70.2% of patients were of European ethnicity with Maori accounting for 11.0% and Pacific Peoples 9.8%; (Table 2)

Table 2. Ethnicity of patients undergoing cardiac surgery in 2017

	Count	Percentage
Maori	300	11.0%
Pacific Peoples	267	9.8%
European	1,915	70.2%
Asian	77	2.8%
Middle Eastern / Latin American / African	13	0.5%
Residual categories	69	2.5%
Other ethnicity	86	3.2%
Unspecified	0	
All	2,727	

Fig. 2 All cardiac surgery patients: Ethnicity; calendar year 2017 (n=2,727)



Risk factors

The risk of heart disease is influenced by a number of factors. These include age, sex, lifestyle choices (e.g., smoking), elevated cholesterol levels (familial, high cholesterol diet, lack of exercise), high blood pressure and diabetes. The risk factor spectrum continues to remain similar. Further analysis of this will need to be undertaken over the coming years to determine variation within diverse ethnic groups and areas for targeted improvement.

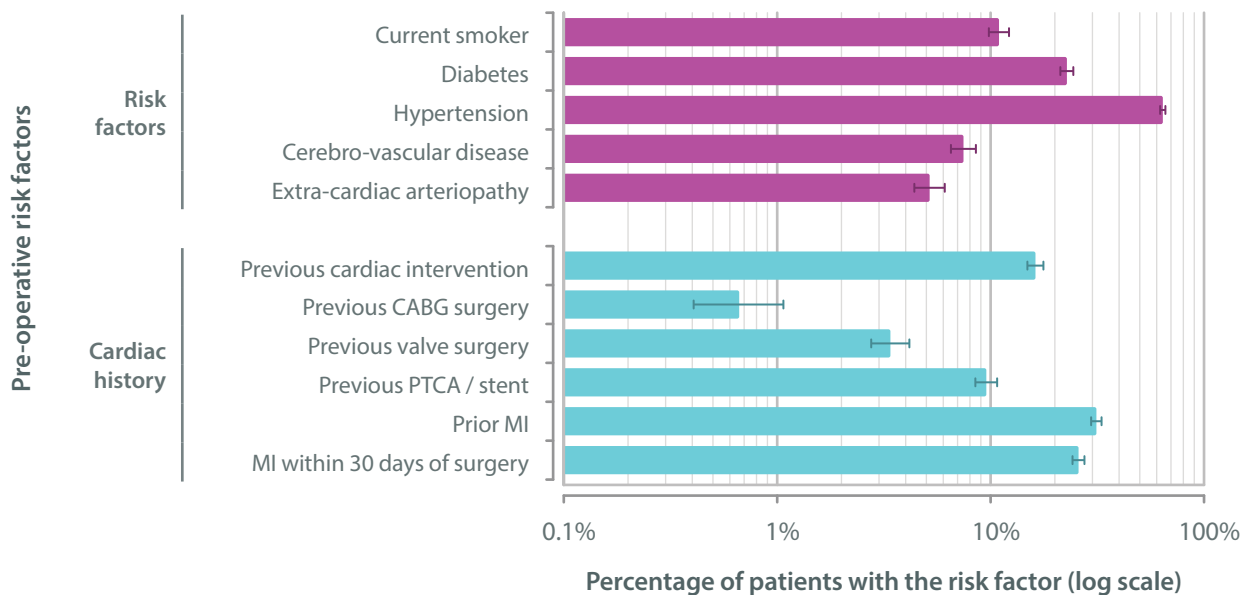
- One in 10 patients were still smoking at the time of surgery in 2017; in 2016 this figure was higher with one in eight patients still smoking at the time of surgery.
- Approximately one-quarter of the patients had diabetes.
- Over 60% of patients had high blood pressure.

Table 3. All patients in 2017: pre-operative risk factors

		Risk factor present			Percentage with the risk factor
		No	Yes	Unspecified	
Risk factors	Current smoker	2,424	298	5	10.9%
	Diabetes	2,102	620	5	22.8%
	Hypertension	977	1,746	4	64.1%
	Cerebro-vascular disease	2,517	203	7	7.5%
	Extra-cardiac arteriopathy	2,581	141	5	5.2%
Cardiac history	Previous cardiac intervention	2,282	442	3	16.2%
	Previous CABG surgery	2,694	18	15	0.7%
	Previous valve surgery	2,620	92	15	3.4%
	Previous PTCA / stent	2,463	260	4	9.5%
	Prior MI	1,871	853	3	31.3%
	Prior MI within 30 days of surgery	2,019	703	5	25.8%

Fig.3

**All cardiac surgery patients:
Risk factors; calendar year 2017**





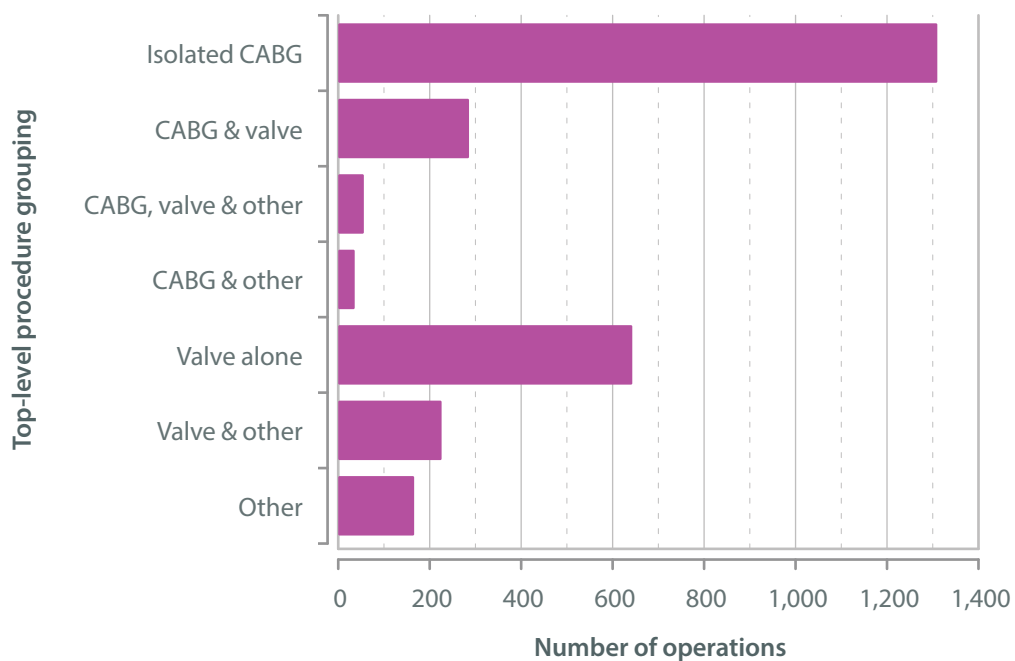
Types of operations performed

- Isolated coronary artery bypass accounted for 48.1% of the total volume of cases.
- Isolated heart valve operations were 23.6%.
- Combined valve and coronary artery bypass 10.5%.
- Approximately 15% of cases were for *Other*, less common procedures.

Table 4. Procedures performed in 2017

	Count	Percentage
CABG	1,310	48.1%
CABG & valve	286	10.5%
CABG, valve & other	56	2.1%
CABG & other	36	1.3%
Valve alone	643	23.6%
Valve & other	226	8.3%
Other	166	6.1%
Unspecified	4	
All	2,727	

Fig. 4 **Operations performed in the calendar year 2017**



Isolated coronary artery bypass surgery

Coronary artery bypass grafting (CABG) is an operation undertaken to bypass blocked arteries of the heart in patients who are not suitable for a non-surgical treatment option (stent placement) or due to failure of stents. The aim of the procedure is to improve quality of life and minimise the risk of a heart attack.

This procedure is the most commonly-performed operation by a cardiac surgeon. In the year 2017 a total of 1,307 patients underwent a publicly-funded isolated CABG operation (47.9%) of the total volume of cardiac surgery. (Table 4). The volumes of the procedure is consistent over the past three years audited.

Coronary artery disease is a condition where cholesterol deposition occurs in the arteries supplying blood to the heart. Multiple risk factors contribute to occurrence of the disease. The risk factors include diabetes, high blood pressure, smoking and obesity (Table 7; Fig. 6) or a combination of them. Some people unfortunately have a genetic predisposition. Other risk factors can enhance early progression of the disease in those with a familial predisposition. They also impact on outcome in terms of complications and early recovery from heart surgery.

Body Mass Index (BMI) is an indicator of a patients size. BMI is calculated by the patient's mass in kg ÷ height in metres squared (Table 5).

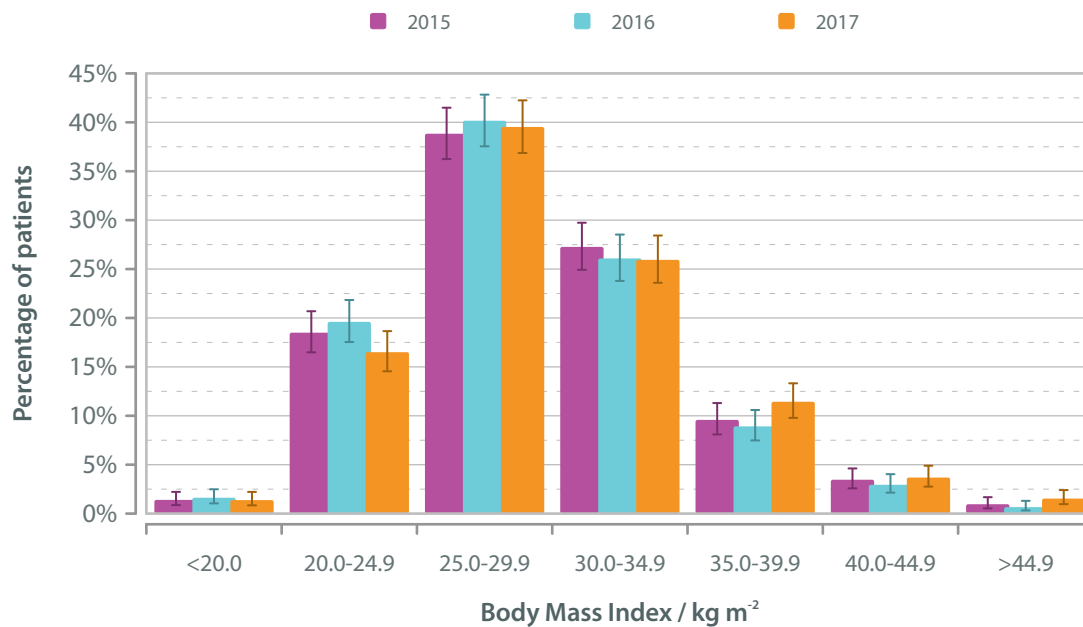
- A BMI of less than 18.5 is defined as underweight; currently less than 1.4% of the CABG patient population fall within this category.
- Approximately 16.5 % of CABG patients are classified within the healthy BMI range NZ European (18.5-24.9).
- The remaining 80% of CABG patients were in the overweight (25.0-29.9), obese (30.0-34.9) or morbidly obese (>35.0) categories.

Table 5. First-time isolated CABG: Body Mass Index in each of the last 3 calendar years

	2015		2016		2017	
	Count	Percentage	Count	Percentage	Count	Percentage
<20.0	19	1.4%	22	1.6%	18	1.4%
20.0-24.9	251	18.5%	266	19.6%	215	16.5%
25.0-29.9	527	38.8%	545	40.2%	515	39.5%
30.0-34.9	370	27.3%	354	26.1%	338	25.9%
35.0-39.9	130	9.6%	121	8.9%	149	11.4%
40.0-44.9	47	3.5%	40	2.9%	48	3.7%
>44.9	13	1.0%	9	0.7%	20	1.5%
Unspecified	12		3		4	
All	1,369		1,360		1,307	



Fig. 5 **First-time isolated CABG: BMI distributions over the last three calendar years; (n=4,017)**



Isolated CABG

BMI classifications

Ministry of Health New Zealand. Body size. Retrieved from: <http://www.health.govt.nz/ourwork/populations/maori-health/tatau-kahukura-maori-health-statistics/nga-tauwehe-tupono-me-temarumaruru-risk-and-protective-factors/body-size>.

Table 6. International BMI cut-off points for adults aged 18 years and over

Classification	BMI range (kg m ⁻²)	Risk of health conditions
Underweight	<18.5	Low risk
Normal range	18.5-24.9	Average risk
Overweight	25.0-29.9	Increased risk
Obese	>29.9	Substantially increased risk

Isolated CABG



In the New Zealand population entered in the registry, the incidence of these risk factors was:

- One in ten patients (11.2%) were current smokers.
- One-third of the patients (31.7%) were diabetic.
- More than 70% of the patients had high blood pressure (hypertension).

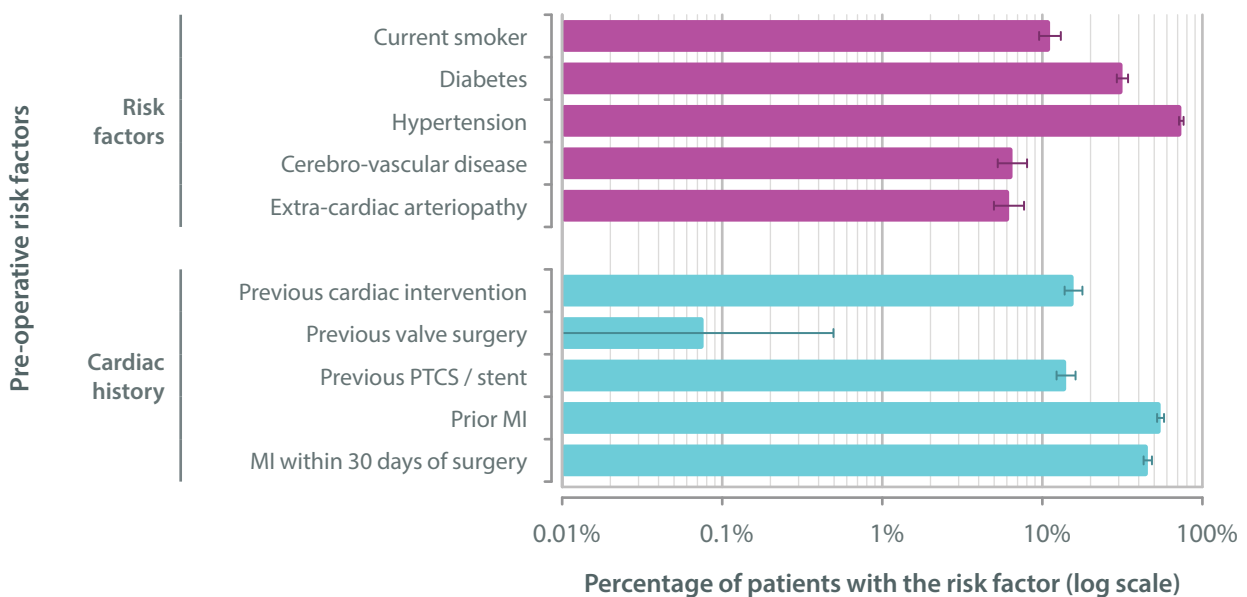
The incidence rate has remained consistent over the last 3 years of reporting.

Table 7. First-time isolated CABG in 2017: pre-operative risk factors

		Risk factor present			Percentage with the risk factor
		No	Yes	Unspecified	
Risk factors	Current smoker	1,160	146	1	11.2%
	Diabetes	892	414	1	31.7%
	Hypertension	341	966	0	73.9%
	Cerebro-vascular disease	1,220	85	2	6.5%
	Extra-cardiac arteriopathy	1,225	81	1	6.2%
Cardiac history	Previous cardiac intervention	1,102	205	0	15.7%
	Previous valve surgery	1,306	1	0	0.1%
	Previous PTCA / stent	1,122	184	1	14.1%
	Prior MI	590	717	0	54.9%
	Prior MI within 30 days of surgery	710	596	1	45.6%

Fig. 6

First-time isolated CABG: Risk factors; calendar year 2017



The most common presentation of these patients is between 50 to 75 years of age, which accounts for over 85% of the total volume. The majority of these patients present between 50 to 79 years of age, with men presenting at an earlier age than female patients. The overall male to female ratio is 4.4 : 1.

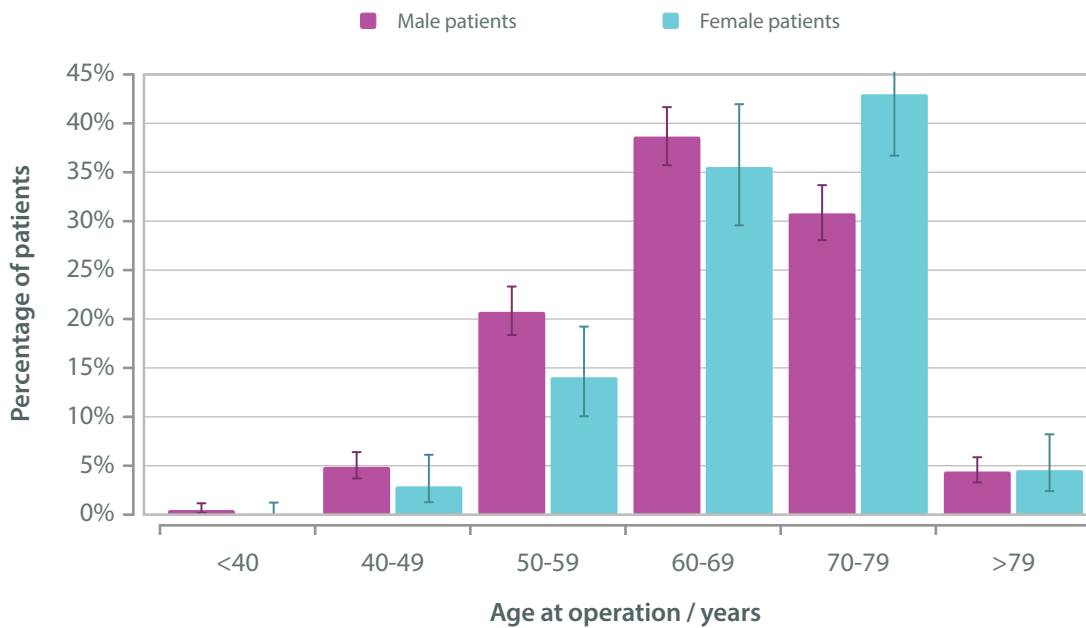
Table 8. First-time isolated CABG in 2017: age and gender

Age at operation / years	Gender		
	Male	Female	All
<40	5	0	5
40-49	52	7	59
50-59	221	34	255
60-69	412	86	498
70-79	328	104	432
>79	47	11	58
Unspecified	0	0	0
All	1,065	242	1,307

Isolated CABG

Fig. 7

First-time isolated CABG: Age & gender distributions; calendar year 2017 (n=1,307)



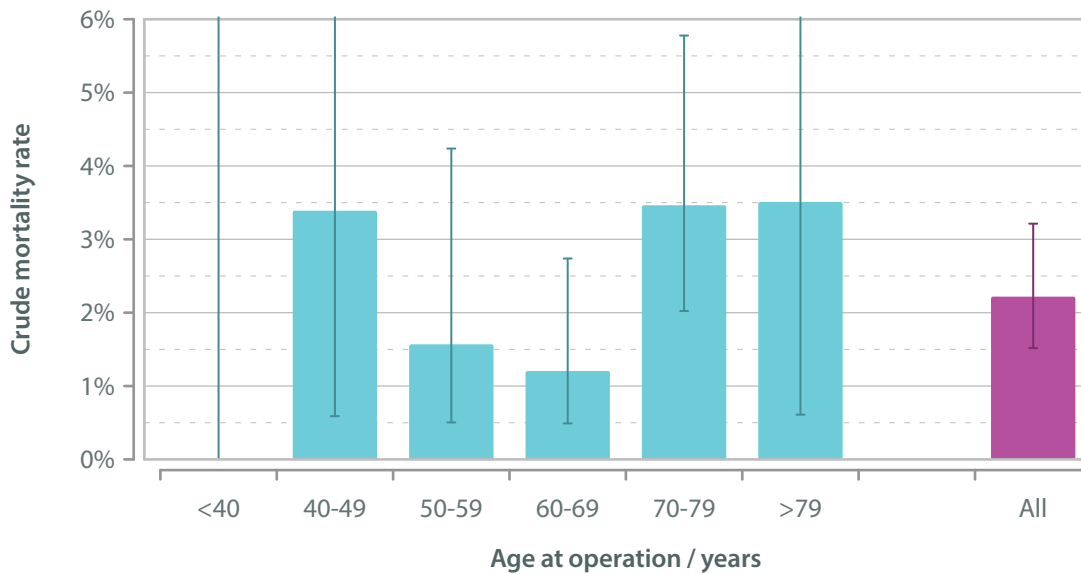


The overall survival results of isolated CABG operations nationwide is well within the International standard bench mark of care at 97.8%¹. This is similar to the previous year before.

Table 9. First-time isolated CABG in 2017: age and in-hospital mortality

Age at operation / years	In-hospital mortality			Mortality rate (95% CI)
	No	Yes	All	
<40	5	0	5	0.0% (0.0-45.1%)
40-49	57	2	59	3.4% (0.6-12.7%)
50-59	251	4	255	1.6% (0.5-4.2%)
60-69	492	6	498	1.2% (0.5-2.7%)
70-79	418	15	433	3.5% (2.0-5.8%)
>79	55	2	57	3.4% (0.6-13.2%)
Unspecified	0	0	0	NA
All	1,278	29	1,307	2.2% (1.5-3.2%)

Fig. 8 First-time isolated CABG: In-hospital mortality and age; calendar year 2017 (n=1,307)



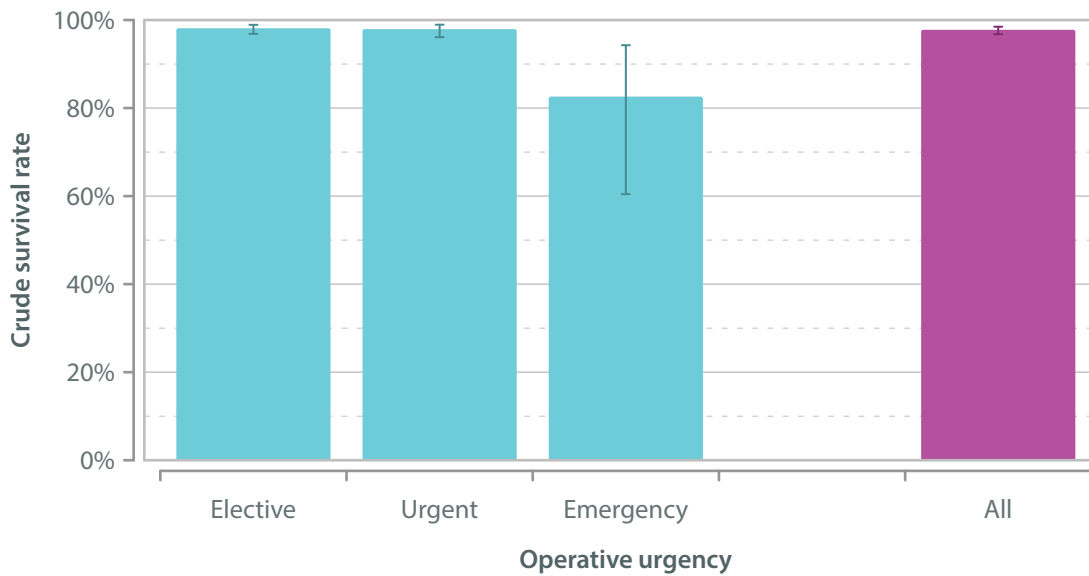
1. <http://anzscts.org/wp-content/uploads/2018/11/181024-ANZSCTS-2017-National-Annual-Report-v2.6.3-electronic-version.pdf>.

Isolated CABG

Table 10. First-time isolated CABG in 2017: operative urgency and in-hospital survival

		In-hospital survival			
		Yes	No	All	Survival rate (95% CI)
Operative urgency	Elective	786	15	801	98.1% (96.9-98.9%)
	Urgent	473	10	483	97.9% (96.1-98.9%)
	Emergency / salvage	19	4	23	82.6% (60.5-94.3%)
	All	1,278	29	1,307	97.8% (96.8-98.5%)

Fig. 9 First-time isolated CABG: In-hospital survival rates; calendar year 2017 (n=1,307)



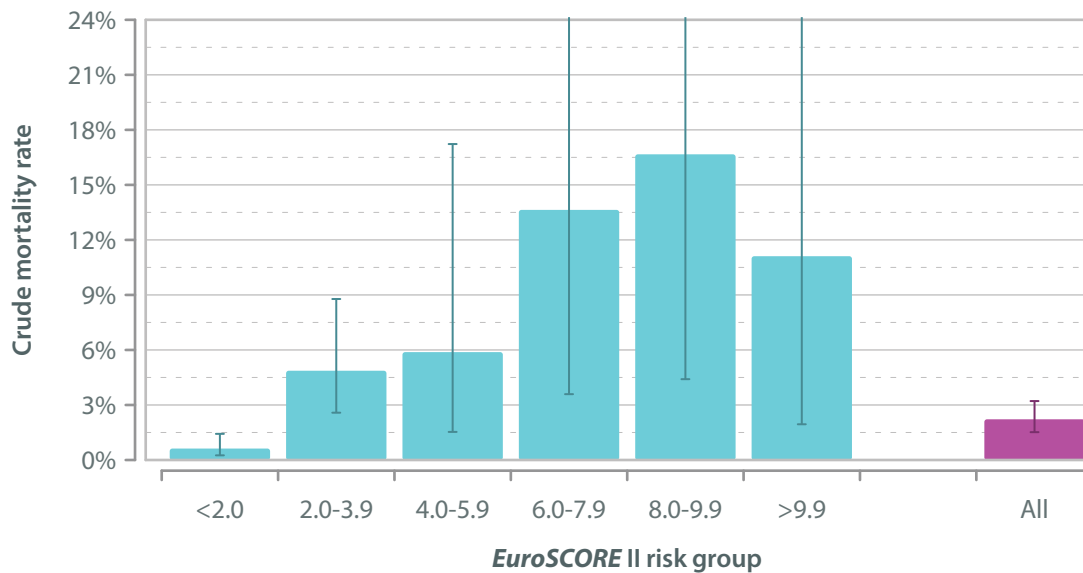


The majority of the deaths were in the high-risk group of patients. As expected the salvage and emergency procedures had a higher mortality. The risk of the patient was based on a **EuroSCORE²**. This takes into account risk factors associated with coronary artery disease and the higher the score the greater risk of morbidity and mortality post-surgery (e.g., **EuroSCORE** 6.0-7.9).

Table 11. First-time isolated CABG in 2017: **EuroSCORE** II risk score and in-hospital mortality

	In-hospital mortality			
	No	Yes	All	Mortality rate (95% CI)
EuroSCORE II				
<2.0	955	6	961	0.6% (0.3-1.4%)
2.0-3.9	215	11	226	4.9% (2.6-8.8%)
4.0-5.9	48	3	51	5.9% (1.5-17.2%)
6.0-7.9	19	3	22	13.6% (3.6-36.0%)
8.0-9.9	15	3	18	16.7% (4.4-42.3%)
>9.9	16	2	18	11.1% (1.9-36.1%)
Unspecified	10	1	11	9.1% (0.5-42.9%)
All	1,278	29	1,307	2.2% (1.5-3.2%)

Fig. 10 First-time isolated CABG: In-hospital mortality and pre-operative risk; calendar year 2017



2. **EuroSCORE** II is a method of calculating predicted operative mortality for patients undergoing cardiac surgery (Table 11). It is not a determinant factor for precluding any patient from having surgical intervention.

Quality of care of cardiac surgical patients

The success and quality of care provided for a cardiac surgical patient is determined far more on the journey of the patient. From the time of being accepted for surgery to discharge from the hospital following surgery and not only the mortality associated with the procedure. The impact of the team in delivery of a satisfactory outcome cannot be underestimated.

The registry is designed to measure these quality measures to allow us to identify and focus on specific areas and help improve quality of care. Some of these measures include mechanical ventilation, time spent in the intensive care unit, hospital stay and wound infection.

Mechanical ventilation is temporarily required following cardiac surgery. The duration of ventilated assistance is determined to a large extent by the complexity of the patient's procedure and the presence or absence of pre-existing risk factors such as obesity and lung function (Table 12). The median ventilation time for 2017 was 6 hours.

Following cardiac surgery patients usually spend a period of time in intensive care (ICU) and are transferred to the ward once fully recovered. The median time spent in ICU for 2017 was 23 hours. Time spent in ICU is determined by how quickly the patients recover, which is impacted by comorbidity conditions and complications of the procedure.

Patients' length-of-stay in hospital following a CABG procedure was on average 6 days. These all compare favourably with the international literature³. These rates have consistently been similar for the last three years. Taking these data into consideration, continued improvement initiatives for the patient journey through cardiac surgery are being considered at individual DHB levels. These pilot programs should help improve quality of care Nationally.

Table 12. First-time isolated CABG in 2017: hospital resource utilisation

		No	Yes	Rate
Resource utilisation	Same day admission	1,267	40	3.1%
		Count	Median	Inter-quartile range
	Ventilation time / hours	1,283	6.0	4-11
	Time on ICU / hours	1,284	23.0	20-44
	Post-operative stay / days	1,303	6.0	5-7
	Hospital stay / days	1,301	10.0	7-16

3. <http://anzscts.org/wp-content/uploads/2017/11/171128-ANZSCTS-2016-National-Annual-Report-FINAL-medium.pdf>.



Complications following cardiac surgery are not only determined by patient conditions, but also reflect the quality of care that the patient receives; commonly monitored by measurement of:

- Deep sternal wound infection.
- Return to theatre.
- Readmission rates following surgery.

It is encouraging to note improvement in deep sternal wound infections as compared to the previous year. The surgical site infection programme (SSI) is implemented in all cardiac surgery units. Results are reported by the Health Quality and Safety Commission New Zealand⁴. Results have improved and continue to be within internationally accepted guidelines. Changing guidelines for pre-surgical antiplatelet therapy⁵ continues to limit the anticipated improvement in post-surgical bleeding. The data so far for the three years compares favourably with international guidelines. It is the intention that this continued analysis of quality of care will ensure all New Zealanders benefit from high standards of cardiac surgery and further improvement measures can be identified. We anticipate the cardiac surgical registry will allow us to review and analyse other improvement measures.

Table 13. First-time isolated CABG in 2017: complications

		Complication			
		No	Yes	Unspecified	Rate (95% CI)
In-hospital	Deep sternal wound infection	1,297	9	1	0.7% (0.3-1.4%)
	Any return to theatre	1,237	70	0	5.4% (4.2-6.8%)
	Return to theatre for bleeding	1,261	40	6	3.1% (2.2-4.2%)
30-day	Readmission	1,161	143	3	11.0% (9.3-12.8%)
	Deep sternal wound infection	1,283	18	6	1.4% (0.8-2.2%)

4. www.hqsc.govt.nz

5. New guidelines for pre surgical cessation of anti platelet therapy <https://academic.oup.com/eurheartj/article/39/3/213/4095043#108531397>.

Aortic valve surgery

Aortic valve replacement (AVR) is undertaken to replace a diseased aortic valve. This is done with either a synthetic mechanical valve or a valve made from animal tissue. Damage to the native aortic valve leads to symptoms that may include shortness of breath, chest pain, dizziness or fainting. Internationally AVR is the most commonly performed isolated valve procedure performed by a cardiac surgeon. It is used internationally as an index procedure for benchmarking and reporting of key performance indicators and quality of care reporting.

Surgical aortic valve replacement (sAVR) is the gold standard intervention for the majority of patients with aortic valve disease and is performed by a cardiac surgical team utilising an incision in the chest and with the use of a heart and lung / cardiopulmonary bypass machine.

Transcatheter aortic valve interventions (TAVI or TAVR) are also performed in New Zealand for patients with aortic stenosis. At this time TAVR is currently performed in high-risk surgical patients and is used in a smaller patient population when compared to sAVR. The decision to perform TAVR in an individual patient is made by a multi-disciplinary team of physicians, surgeons and allied health specialists in combination with the patient and their Whanau. The outcomes of TAVR are not currently discussed in this report. A New Zealand TAVR database is currently in development and in collaboration with the National cardiac Network will be linked to the National Cardiac Surgical Registry to enable future reporting that can compare and contrast the patient populations undergoing each treatment. Future reporting of TAVR outcomes in the cardiac surgical report will allow us to gain further insights into the appropriateness of each treatment in a New Zealand context and further allow us to define the optimal patient population for treatment modality. At the current time however, all patients presented in this current report underwent standard open surgical aortic valve replacement performed by a cardiac surgical team.

Table 14. Valve surgery in 2017

	Top-level procedure classification	
	Valve alone	CABG & valve
Aortic valve alone	354	213
Mitral valve alone	159	45
Aortic & mitral valves	42	13
Mitral & tricuspid valves	52	4
Others	33	2
Unspecified	3	9
All	643	286

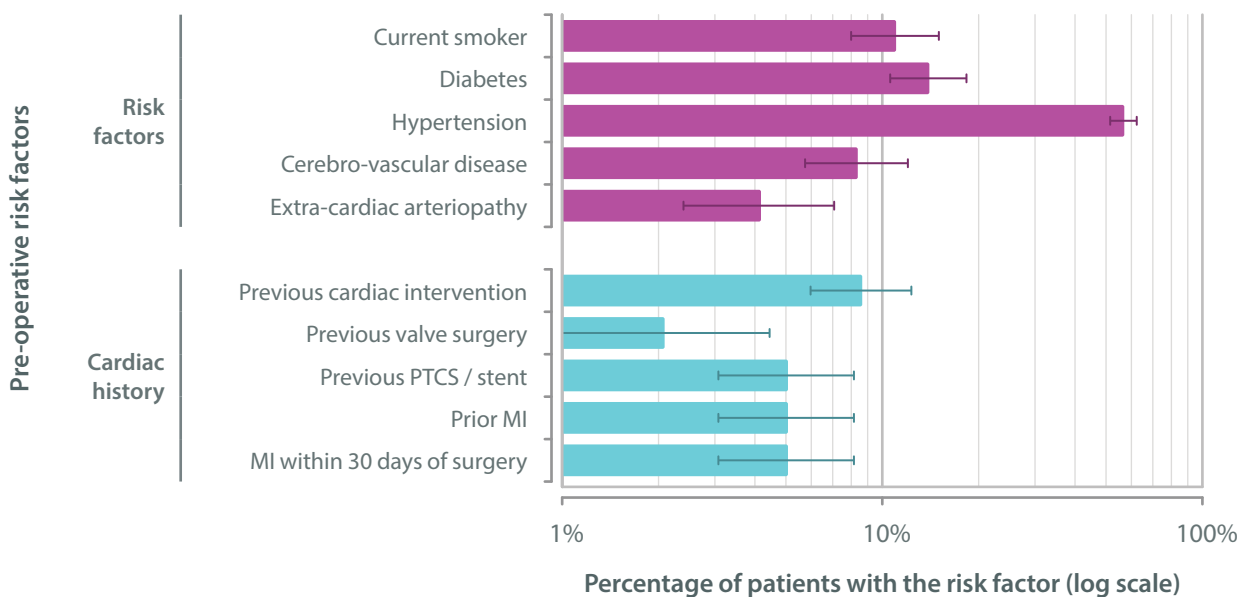


Table 15. First-time isolated AVR in 2017: pre-operative risk factors

	Risk factor present			Percentage with the risk factor	
	No	Yes	Unspecified		
Risk factors	Current smoker	298	37	0	11.0%
	Diabetes	288	47	0	14.0%
	Hypertension	144	191	0	57.0%
	Cerebro-vascular disease	306	28	1	8.4%
	Extra-cardiac arteriopathy	321	14	0	4.2%
Cardiac history	Previous cardiac intervention	306	29	0	8.7%
	Previous CABG surgery	328	7	0	2.1%
	Previous PTCA / stent	318	17	0	5.1%
	Prior MI	318	17	0	5.1%
	Prior MI within 30 days of surgery	318	17	0	5.1%

Fig. 11

**First-time isolated AVR:
Risk factors; calendar year 2017**



In the New Zealand registry 334 isolated first-time AVRs have been performed, which is approximately 12% of the overall surgical volume (Table 14). This is not significantly different when compared to the previous year's volume.

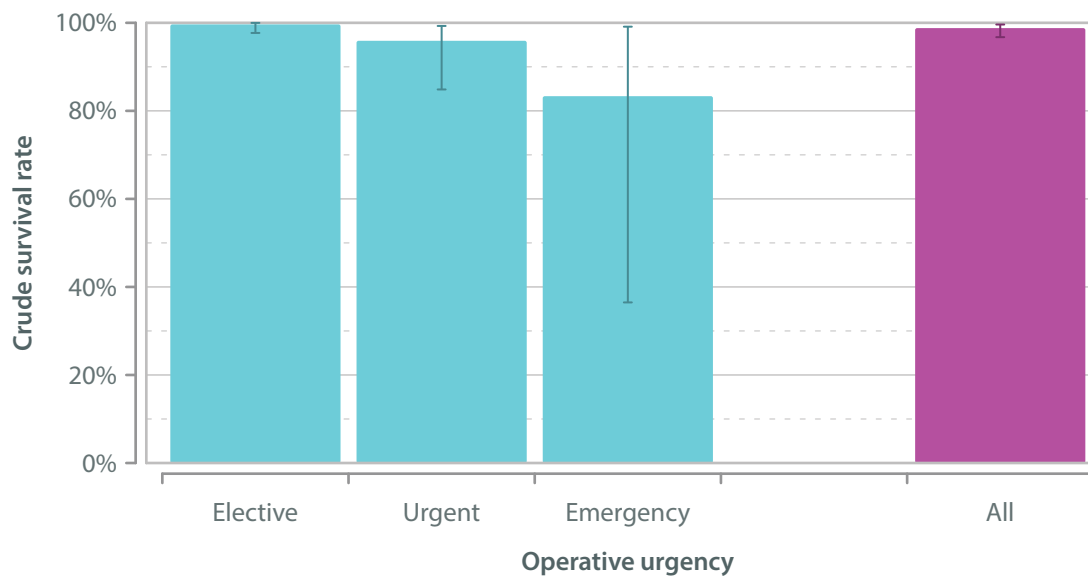
Most patients undergo surgery in a planned or elective fashion with smaller numbers undergoing urgent, emergency or salvage surgery (Table 16; Fig. 12). As expected a significant number of the patients have additional cardiovascular risk factors including: 14.0% diabetics, 57.0% with hypertension and 8.4% having had a previous cardiovascular intervention (Table 15; Fig. 11).

Aortic valve surgery may be required because of either leakage of the valve (aortic regurgitation) or blockage of the valve (aortic stenosis). These conditions can occur for a variety of reasons, the most common being degenerative age-related calcification or hardening of the valve. Dysfunction of the valve may also be due to conditions such as rheumatic fever that can damage the structure of the valve or in some cases be due to a congenital abnormality (bicuspid aortic valve) that causes it to fail at an earlier age. In some cases the valve may need to be replaced because of infection on the leaflets that lead to valve destruction.

Table 16. First-time isolated AVR in 2017: operative urgency and in-hospital survival

		In-hospital survival			
		Yes	No	Unspecified	Survival rate (95% CI)
Operative urgency	Elective	277	1	1	99.6% (97.7-100.0%)
	Urgent	47	2	1	95.9% (84.9-99.3%)
	Emergency / salvage	5	1	0	83.3% (36.5-99.1%)
	All	329	4	2	98.8% (96.7-99.6%)

Fig. 12 First-time isolated AVR: In-hospital survival rates; calendar year 2017 (n=333)



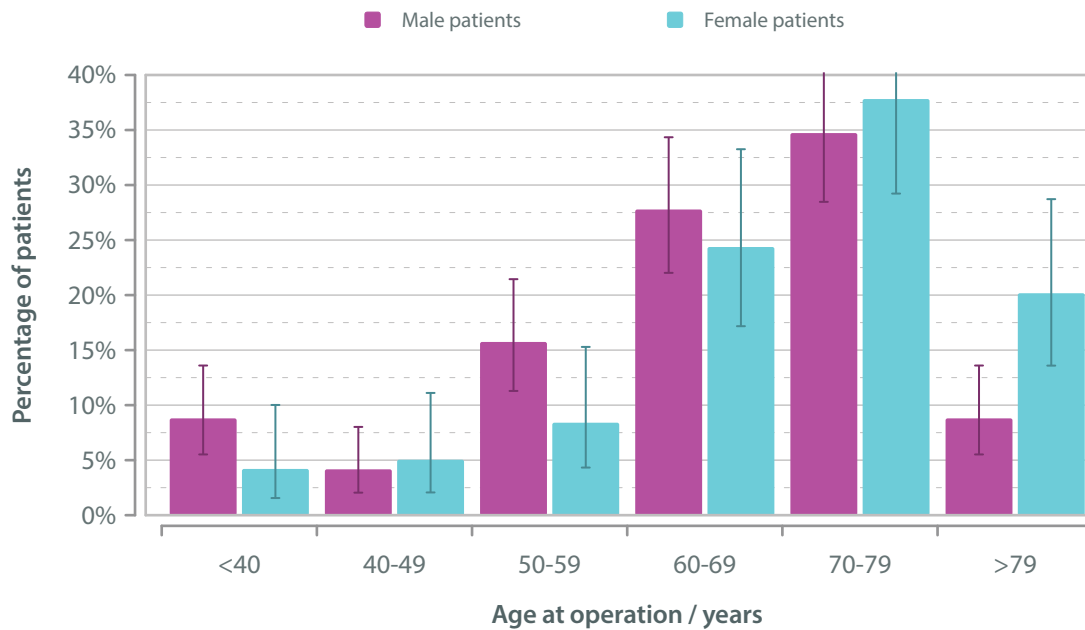


The majority of patients have age related calcific aortic stenosis and this tends to occur later in life in particular in the >70 years of age group (Table 17; Fig. 13). Younger patients are more likely to have an AVR due to rheumatic fever, a bicuspid valves or infection on the leaflets. The total male to female ratio in this report is 1.8: 1.

Table 17. First-time isolated AVR in 2017: age and gender

Age at operation / years	Gender		
	Male	Female	All
<40	19	5	24
40-49	9	6	15
50-59	34	10	44
60-69	60	29	89
70-79	75	45	120
>79	19	24	43
Unspecified	0	0	0
All	216	119	335

Fig. 13 First-time isolated AVR: Age & gender distributions; calendar year 2017 (n=335)



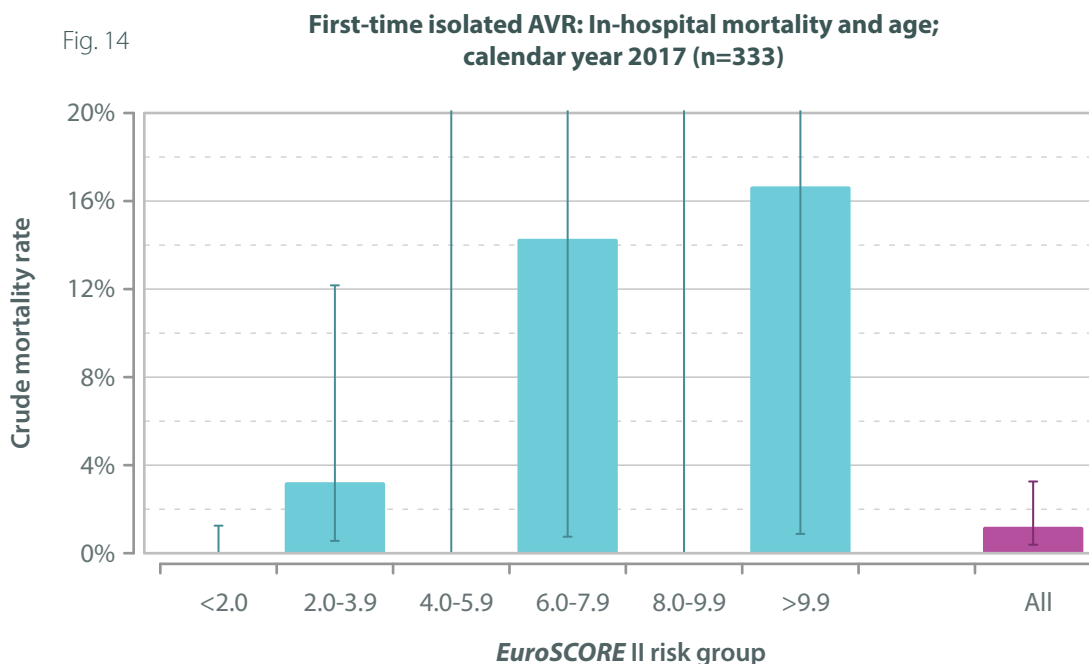
The **EuroSCORE II** is an internationally recognised tool used to predict mortality in patients undergoing cardiac surgery. It is a tool that we have used in this report to risk stratify patients undergoing AVR and to assess our performance against expected outcomes. From 2016 all units were collecting **EuroSCORE II** and the 2017 report has **EuroSCORE II** generated for all but 2 of 335 patients and gives a better reflection of risk-adjusted outcomes for the entire New Zealand surgical population.

Table 18 shows the distribution of risk profiles in patients undergoing isolated AVR and the observed mortality rate. The overall observed mortality for isolated AVR in New Zealand was extremely low (1.2%), which is favourably compared to internationally accepted outcomes. The 2017 ANZCTS publication of surgical outcomes reported a 1.6% mortality for isolated AVR in a similar cohort of patients.

As can be seen the majority of NZ patients (238) are in the low-risk category (ES <2.0% predicted mortality) with an observed mortality of 0.0% in this group of patients. Therefore, for the majority of patients in New Zealand being referred for AVR, and in the absence of major comorbidity, AVR is performed with extremely low mortality. As such AVR remains the gold standard intervention in this group with which emerging therapies such as TAVR need to be compared within a New Zealand health care context.

Table 18. First-time isolated AVR in 2017: **EuroSCORE II** risk score and in-hospital mortality

	In-hospital mortality			
	No	Yes	Unspecified	Mortality rate (95% CI)
EuroSCORE II <2.0	238	0	0	0.0% (0.0-1.3%)
2.0-3.9	60	2	0	3.2% (0.6-12.2%)
4.0-5.9	12	0	0	0.0% (0.0-22.1%)
6.0-7.9	6	1	0	14.3% (0.8-58.0%)
8.0-9.9	3	0	0	0.0% (0.0-63.2%)
>9.9	5	1	0	16.7% (0.9-63.5%)
Unspecified	5	0	2	0.0% (0.0-45.1%)
All	329	4	2	1.2% (0.4-3.3%)



6. <http://anzscts.org/wp-content/uploads/2018/11/181024-ANZSCTS-2017-National-Annual-Report-v2.6.3-electronic-version.pdf>



Reported outcomes in groups with fewer numbers of patients are heavily influenced by those small numbers and therefore mortality rates have to be interpreted in the context of statistical variance. It was however reassuring to see very low mortality in the higher-risk cohort of patients.

There were 2 deaths recorded in patients with a **EuroSCORE II** >4.0.

Major morbidity compares favourably to international reported results⁷. This includes low incidence of deep sternal wound infection rates (0.3%) and return to theatre for bleeding (3.9%).

Reported results suggest that for isolated AVR all DHBs and NZ surgeons as a collective group are performing within accepted standards when benchmarked to results observed within the United Kingdom and Australia.

Table 19. First-time isolated AVR in 2017: hospital resource utilisation

Resource utilisation	No	Yes	Rate
	Count	Median	Inter-quartile range
Same day admission	328	7	2.1%
Ventilation time / hours	332	5.5	4.0-11.0
Time on ICU / hours	330	23.0	20.0-41.5
Post-operative stay / days	334	6.0	5.0-8.0
Hospital stay / days	333	8.0	7.0-12.0

Table 20. Isolated aortic valve surgery in 2017: complications

	Complication				
	No	Yes	Unspecified	Rate (95% CI)	
In-hospital	Deep sternal wound infection	334	1	0	0.3% (0.0-1.9%)
	Any return to theatre	316	19	0	5.7% (3.5-8.9%)
	Return to theatre for bleeding	322	13	0	3.9% (2.2-6.7%)
30-day	Readmission	296	39	0	11.6% (8.5-15.7%)
	Deep sternal wound infection	333	1	1	0.3% (0.0-1.9%)

7. <http://anzscts.org/wp-content/uploads/2018/11/181024-ANZSCTS-2017-National-Annual-Report-v2.6.3-electronic-version.pdf>

Summary

I would sincerely like to thank all New Zealand cardiothoracic units and the Ministry of Health (MOH) for the result of this enormous effort. The responsible collection and analysis of valid data by a national sub-specialty group has enhanced the monitoring of quality of care, contributed to improved outcomes and resource utilisation, and in a challenging healthcare economic environment has provided cardiothoracic surgeons with a tool to help protect our patients from poor outcomes. The National Adult Cardiac Surgery Database is increasingly utilised by the district health boards (DHB) and the MOH and by our colleagues as a tool and platform for the conduct of important clinical outcomes research.

Various local investigations by the DHBs and HDC for poor outcomes, such as the Bristol, United Kingdom (UK) Inquiry, often follow upon publication in the lay press of raw outcome data, cited out of context, often incomplete if not inaccurate, and virtually never enhanced by information regarding risk stratification. Furthermore, after the inquiry is completed and findings published, the lay press may never correct the original allegations and usually does not publish the findings of the inquiry, which may be unfavourable towards the surgeons and physicians. In the final analysis, the upshot of lengthy expensive inquiries of this sort uniformly has been the recommendation to establish and maintain a reliable registry database.

The National Adult Cardiac Surgery Database has numerous disease entities to analyse and by the nature of the sub-specialty requires increased complexity in data analysis in order to produce meaningful risk stratification. As the database matures in the coming years the usefulness of the data will become apparent.

First and foremost, participating units should not fear the potentially negative consequences of reporting less than stellar results. The point is to identify the problems and institute improvement initiatives, which can include inter-institutional team visits, mentoring schemes, and educational programs. Efforts must now focus on developing mechanisms for verification of data completeness and accuracy, improving and validating our methodology of complexity



Definitions


1. **Deep sternal wound infection:** is a serious post-operative complication of cardiac surgery.
2. **Elective:** the procedure could be deferred without the risk of compromised cardiac outcome.
3. **Emergency:** unscheduled surgery required in next available theatre on same day due to refractory angina or cardiac compromise.
4. **EuroSCORE II:** an internationally recognised tool used to predict mortality in patients undergoing cardiac surgery. It is a tool that is used to risk stratify patients. **EuroSCORE II** has been developed by studying large numbers of patients (22,381) undergoing cardiac surgery in 154 hospitals in 43 countries¹.
5. **ICU:** intensive care unit.
6. **MI:** myocardial infarction.
7. **Mortality:** includes all deaths at the 5 public hospitals where cardiac surgery is performed prior to discharge and within 30 days of the date of surgery.
8. **PTCA:** percutaneous transluminal coronary angioplasty.
9. **Salvage:** the patient is undergoing CPR *en route* to the operating room, that is, prior to surgical incision.
10. **Urgent:** not routine; there is a medical reason for operating this admission.

1. Nashef SA, Roques F, Sharples LD, Nilsson J, Smith C, Goldstone AR, Lockowandt U. **EuroSCORE II**. *European Journal of Cardiothoracic Surgery*. 2012; **41(4)**: 734-745.

Appendix

Appendix

New Zealand Ministry of Health
NZ Adult Cardiac Surgical Database
 Baseline section; Page 1; Version 1.0 (13 Dec 2013)



Basic demographic data

All baseline data refer to the condition of the patient when they were originally diagnosed.

Unique patient identifier

Gender Male Female

Date of birth dd/mm/yyyy

Registry data


Admission information

Date of admission dd/mm/yyyy

Ethnicity 1 European
 Maori
 Pacific peoples
 Asian
 Middle Eastern/Latin American/African
 Other ethnicity
 Residual categories

Ethnicity 2 European not further defined
 NZ European
 Other European
 NZ Maori
 Pacific Island not further defined
 Samoan
 Cook Island Maori
 Tongan
 Niuean
 Tokelauan
 Fijian
 Other Pacific Island
 Asian not further defined
 Southeast Asian
 Chinese
 Indian
 Other Asian
 Middle Eastern
 Latin American/Hispanic
 African
 Other ethnicity
 Don't know
 Refused to answer
 Response unidentifiable
 Not stated

Date of surgery dd/mm/yyyy



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New Zealand Ministry of Health
NZ Adult Cardiac Surgical Database
Baseline section; Page 2; Version 1.0 (13 Dec 2013)



Unique patient identifier
Date of surgery dd/mm/yyyy

Admission information continued ...

Elective Day of Surgery Admit Patient	<input type="radio"/> No	<input type="radio"/> Yes
Insurance	<input type="radio"/> Public	<input type="radio"/> Self funded
	<input type="radio"/> Private health insurance	<input type="radio"/> Other
Operation number	<input type="radio"/> 1	<input type="radio"/> 4
	<input type="radio"/> 2	<input type="radio"/> 5
	<input type="radio"/> 3	<input type="radio"/> 6
Height	<input type="text"/>	cm
Weight	<input type="text"/>	kg

New Zealand Ministry of Health
NZ Adult Cardiac Surgical Database
 Baseline section; Page 3; Version 1.0 (13 Dec 2013)

Unique patient identifier	<input type="text"/>
Date of surgery	<input type="text"/> dd/mm/yyyy
Patient risk factors	
Smoking history	<input type="radio"/> No <input type="radio"/> Yes
Current smoker	<input type="radio"/> No <input type="radio"/> Yes
Family history of CAD	<input type="radio"/> No <input type="radio"/> Yes <input type="radio"/> Undiscovered
Diabetes	<input type="radio"/> No <input type="radio"/> Yes
Diabetes control	<input type="radio"/> None <input type="radio"/> Oral <input type="radio"/> Diet <input type="radio"/> Insulin
Hypercholesterolaemia	<input type="radio"/> No <input type="radio"/> Yes
Renal: last pre-op creatinine	<input type="text"/> $\mu\text{mol l}^{-1}$
Renal: dialysis	<input type="radio"/> No <input type="radio"/> Yes
Renal: transplant	<input type="radio"/> No <input type="radio"/> Yes
Renal: impairment	<input type="radio"/> Normal (CC >85 ml min ⁻¹) <input type="radio"/> Moderate (CC 50-85 ml min ⁻¹) <input type="radio"/> Severe (CC <50 ml min ⁻¹)
Hypertension	<input type="radio"/> No <input type="radio"/> Yes
Cerebrovascular disease	<input type="radio"/> No <input type="radio"/> Yes
Cerebrovascular disease: type	<input type="radio"/> Coma <input type="radio"/> RIND or TIA <input type="radio"/> CVA <input type="radio"/> Carotid test
Cerebrovascular disease: when	<input type="radio"/> Recent <input type="radio"/> Remote
PVD/extra-cardiac arteriopathy	<input type="radio"/> No <input type="radio"/> Yes
Respiratory/pulmonary disease	<input type="radio"/> No <input type="radio"/> Yes
Respiratory/pulmonary disease: type	<input type="radio"/> Mild <input type="radio"/> Severe <input type="radio"/> Moderate
Infective endocarditis	<input type="radio"/> No <input type="radio"/> Treated <input type="radio"/> Active
Immunosuppressive treatment	<input type="radio"/> No <input type="radio"/> Yes
Poor mobility due to any non-cardiac reason	<input type="radio"/> No <input type="radio"/> Yes



New Zealand Ministry of Health
NZ Adult Cardiac Surgical Database
Baseline section; Page 4; Version 1.0 (13 Dec 2013)



Unique patient identifier
Date of surgery dd/mm/yyyy

Pre-operative cardiac status

Pre-operative cardiac status

Myocardial infarction	<input type="radio"/> No	<input type="radio"/> Yes
Myocardial infarction: type	<input type="radio"/> NSTEMI	<input type="radio"/> STEMI
Myocardial infarction: when	<input type="radio"/> <= 6 hours	<input type="radio"/> 31-90 days
	<input type="radio"/> 6-24 hours	<input type="radio"/> >90 days
	<input type="radio"/> 1-30 days	
Date of last MI (if known)	<input type="text"/> dd/mm/yyyy	
Angina: CCS classification	<input type="radio"/> 0	<input type="radio"/> 3
	<input type="radio"/> 1	<input type="radio"/> 4
	<input type="radio"/> 2	
Treatment of angina: iv GTN	<input type="radio"/> No	<input type="radio"/> Yes
Treatment of angina: iv heparin	<input type="radio"/> No	<input type="radio"/> Yes
Treatment of angina: full dose heparinoids	<input type="radio"/> No	<input type="radio"/> Yes
History of congestive heart failure	<input type="radio"/> No	<input type="radio"/> Yes
CHF at current admission	<input type="radio"/> No	<input type="radio"/> Yes
Dyspnoea: NYHA classification	<input type="radio"/> 1	<input type="radio"/> 3
	<input type="radio"/> 2	<input type="radio"/> 4
Cardiogenic shock	<input type="radio"/> No	<input type="radio"/> Yes
Resuscitation within 1 hour of operation	<input type="radio"/> No	<input type="radio"/> Yes
Critical pre-operative state	<input type="radio"/> No	<input type="radio"/> Yes

Pre-operative cardiac status - arrhythmia

Arrhythmia	<input type="radio"/> No	<input type="radio"/> Yes
Arrhythmia: type	<input type="radio"/> Sinus rhythm	<input type="radio"/> Ventricular
	<input type="radio"/> Atrial	<input type="radio"/> Other abnormal rhythm
	<input type="radio"/> Heart block/pacing	
Atrial arrhythmia: type	<input type="radio"/> Paroxysmal	<input type="radio"/> Permanent
	<input type="radio"/> Persistent	
Permanent pacemaker <i>in situ</i>	<input type="radio"/> No	<input type="radio"/> Yes

New Zealand Ministry of Health
NZ Adult Cardiac Surgical Database
 Baseline section; Page 5; Version 1.0 (13 Dec 2013)

Unique patient identifier	<input type="text"/>
Date of surgery	<input type="text"/> dd/mm/yyyy
Medication at the time of surgery	
Inotropes	<input type="radio"/> No <input type="radio"/> Yes
iv nitrates	<input type="radio"/> No <input type="radio"/> Yes
Anticoagulation therapy	<input type="radio"/> No <input type="radio"/> Yes
Steroids	<input type="radio"/> No <input type="radio"/> Yes
Thrombolysis (this admission)	<input type="radio"/> No <input type="radio"/> Yes
Thrombolysis: interval	<input type="text"/> hours
Aspirin within 7 days of surgery	<input type="radio"/> No <input type="radio"/> Yes
Aspirin: when	<input type="radio"/> ≤2 days <input type="radio"/> 3-7 days
Clopidogrel within 7 days of surgery	<input type="radio"/> No <input type="radio"/> Yes
Clopidogrel: when	<input type="radio"/> ≤2 days <input type="radio"/> 3-7 days
IIb / IIIa blockade within 7 days of surgery	<input type="radio"/> No <input type="radio"/> Yes
IIb / IIIa blockade: when	<input type="radio"/> ≤2 days <input type="radio"/> 3-7 days
Aggrostat within 7 days of surgery	<input type="radio"/> No <input type="radio"/> Yes
Aggrostat: when	<input type="radio"/> ≤2 days <input type="radio"/> 3-7 days
Other antiplatelet therapy within 7 days of surgery	<input type="radio"/> No <input type="radio"/> Yes
Other antiplatelet: when	<input type="radio"/> ≤2 days <input type="radio"/> 3-7 days



New Zealand Ministry of Health
NZ Adult Cardiac Surgical Database
Baseline section; Page 6; Version 1.0 (13 Dec 2013)



Unique patient identifier

Date of surgery dd/mm/yyyy

Previous interventions (surgical or percutaneous)

Previous cardiothoracic intervention	<input type="radio"/> No		<input type="radio"/> Yes	
Previous surgery	<input type="radio"/> No		<input type="radio"/> Yes	
Type of previous surgery	<input type="checkbox"/> CABG	<input type="checkbox"/> Off-pump CABG	<input type="checkbox"/> Valve	<input type="checkbox"/> Congenital cardiac
				<input type="checkbox"/> Aortic surgery (ascending/arch)
				<input type="checkbox"/> Other cardiac
Number of prior cardiac operations requiring cardiopulmonary bypass	<input type="radio"/> 0	<input type="radio"/> 1	<input type="radio"/> 2	<input type="radio"/> 3
		<input type="radio"/> 4	<input type="radio"/> 5	<input type="radio"/> 6
			<input type="radio"/> 7	<input type="radio"/> 8
				<input type="radio"/> 9
Number of prior cardiac operations without cardiopulmonary bypass	<input type="radio"/> 0	<input type="radio"/> 1	<input type="radio"/> 2	<input type="radio"/> 3
		<input type="radio"/> 4	<input type="radio"/> 5	<input type="radio"/> 6
			<input type="radio"/> 7	<input type="radio"/> 8
				<input type="radio"/> 9
Previous percutaneous intervention	<input type="radio"/> No		<input type="radio"/> Yes	
PTCA/stent	<input type="radio"/> No		<input type="radio"/> Yes	
PTCA/stent: which admission	<input type="radio"/> Prior admission		<input type="radio"/> This admission	
PTCA/stent: interval (same admission)	<input type="text"/>	hours		
Other percutaneous interventions	<input type="checkbox"/> Non-surgical balloon valvuloplasty <input type="checkbox"/> ASD device closure <input type="checkbox"/> VSD device closure <input type="checkbox"/> Percutaneous SVT/VT ablation			

Appendix



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New Zealand Ministry of Health
NZ Adult Cardiac Surgical Database
 Baseline section; Page 7; Version 1.0 (13 Dec 2013)

Unique patient identifier	<input type="text"/>
Date of surgery	<input type="text"/> dd/mm/yyyy
Haemodynamic data	
Cardiac catheterisation	<input type="radio"/> No <input type="radio"/> Yes
Date of cardiac catheterisation	<input type="text"/> dd/mm/yyyy
LVEF method	<input type="radio"/> Not measured <input type="radio"/> LV gram <input type="radio"/> Radionuclide <input type="radio"/> Echo <input type="radio"/> MRI
EF	<input type="text"/> %
EF estimate	<input type="radio"/> Normal <input type="radio"/> Mild <input type="radio"/> Moderate <input type="radio"/> Severe
Left main stenosis >50%	<input type="radio"/> No <input type="radio"/> Yes
Number of diseased coronary systems	<input type="radio"/> None <input type="radio"/> One <input type="radio"/> Two <input type="radio"/> Three
PA systolic	<input type="text"/> mm Hg
Pulmonary hypertension	<input type="radio"/> No <input type="radio"/> Moderate <input type="radio"/> Severe





New Zealand Ministry of Health
NZ Adult Cardiac Surgical Database
Baseline section; Page 8; Version 1.0 (13 Dec 2013)



Unique patient identifier
Date of surgery dd/mm/yyyy

Operation status / category

Surgery data

Consultant surgeon

Operating surgeon Consultant
 Senior registrar Overseas fellow
 Trainee Oversight

Operative urgency / status Elective Emergency
 Urgent Salvage

Direct transfer from cath lab to theatre No Yes

Coronary artery bypass No Yes

Valve surgery No Yes

Valve type Aortic Tricuspid
 Mitral Pulmonary

Redo valve No Yes

Reason for repeat valve placement Prosthetic / homograft valve failure
 Thrombosis Haemolysis
 Dehiscence Prior valve repair
 Embolism Other reason
 Infection

Aortic procedure No Yes

Other cardiac procedures No Yes

Other non-cardiac procedures No Yes

Aortic procedure

Aortic aneurysm repair (type) No repair
 Ascending Descending
 Arch Thoracic/abdominal

Aortic dissection repair (type) No repair Descending
 Ascending

Aortic dissection: when Acute Non-acute

Acute traumatic aortic transection No Yes

New Zealand Ministry of Health
NZ Adult Cardiac Surgical Database
 Baseline section; Page 9; Version 1.0 (13 Dec 2013)

Unique patient identifier
 Date of surgery dd/mm/yyyy

Other cardiac surgery

Atrial arrhythmia surgery	<input type="radio"/> No <input type="radio"/> Yes	
Atrial arrhythmia surgery: lesion set	<input type="radio"/> Cox Maze III <input type="radio"/> Pulmonary vein isolation <input type="radio"/> Radial <input type="radio"/> Left atrial only <input type="radio"/> Mini-Maze <input type="radio"/> Right atrial only <input type="radio"/> Left atrial reduction <input type="radio"/> Other	
Atrial arrhythmia surgery: energy source	<input type="radio"/> Cut & sew <input type="radio"/> Microwave <input type="radio"/> Unipolar RF <input type="radio"/> Laser <input type="radio"/> Bipolar RF <input type="radio"/> Ultrasound <input type="radio"/> Cryoablation <input type="radio"/> Other	
Type of other cardiac surgery	<input type="checkbox"/> AF ablation surgery <input type="checkbox"/> LV rupture <input type="checkbox"/> ASD <input type="checkbox"/> Pericardiectomy <input type="checkbox"/> Atrial myxoma <input type="checkbox"/> Peripheral vascular <input type="checkbox"/> Cardiac transplant <input type="checkbox"/> Permanent LV epicardial lead <input type="checkbox"/> Cardiac trauma <input type="checkbox"/> Primary VAD <input type="checkbox"/> Cardiac trauma - iatrogenic <input type="checkbox"/> Pulm. thromboendarterectomy <input type="checkbox"/> Cardiac tumour <input type="checkbox"/> Pulmonary embolectomy <input type="checkbox"/> Epicardial pacemaker <input type="checkbox"/> Pulmonary transplant <input type="checkbox"/> Left ventricular reconstruction <input type="checkbox"/> VSD (acquired) <input type="checkbox"/> LV aneurysm <input type="checkbox"/> Other congenital <input type="checkbox"/> LVOT myectomy of HOCM <input type="checkbox"/> Other	

Other non-cardiac surgery

Carotid endarterectomy	<input type="radio"/> No <input type="radio"/> Yes	
Lung resection	<input type="radio"/> No <input type="radio"/> Yes	
Other vascular surgery	<input type="radio"/> No <input type="radio"/> Yes	
Other thoracic surgery	<input type="radio"/> No <input type="radio"/> Yes	
Other surgery	<input type="radio"/> No <input type="radio"/> Yes	



New Zealand Ministry of Health
NZ Adult Cardiac Surgical Database
Baseline section; Page 10; Version 1.0 (13 Dec 2013)



Unique patient identifier
Date of surgery dd/mm/yyyy

CPB and support

Minimally invasive

Minimally invasive techniques attempted	<input type="radio"/> No	<input type="radio"/> Yes
Minimally invasive techniques indication	<input type="radio"/> Choice <input type="radio"/> Contraindication	<input type="radio"/> Catheter
Performed off pump	<input type="radio"/> No	<input type="radio"/> Yes
Robotically assisted	<input type="radio"/> No	<input type="radio"/> Yes


CPB and mechanical support

Cardiopulmonary bypass used	<input type="radio"/> No	<input type="radio"/> Yes
Cardioplegia used	<input type="radio"/> No	<input type="radio"/> Yes
Cumulative cross clamp time	<input type="text"/>	min
Cumulative cardiopulmonary bypass time	<input type="text"/>	min
IABP	<input type="radio"/> No	<input type="radio"/> Yes
IABP: when inserted	<input type="radio"/> Pre-op <input type="radio"/> Intra-op	<input type="radio"/> Post-op
IABP: indication	<input type="radio"/> Haemodynamic instability <input type="radio"/> PTCA support <input type="radio"/> Unstable angina	<input type="radio"/> CPB wean <input type="radio"/> Prophylactic
Rota-pump	<input type="radio"/> No	<input type="radio"/> Yes
Rota-pump: when inserted	<input type="radio"/> Pre-op <input type="radio"/> Intra-op	<input type="radio"/> Post-op
Rota-pump: indication	<input type="radio"/> Haemodynamic instability <input type="radio"/> PTCA support <input type="radio"/> Unstable angina	<input type="radio"/> CPB wean <input type="radio"/> Prophylactic
Other mechanical support	<input type="radio"/> No	<input type="radio"/> Yes
Other mechanical support: when inserted	<input type="radio"/> Pre-op <input type="radio"/> Intra-op	<input type="radio"/> Post-op
Other mechanical support: indication	<input type="radio"/> Haemodynamic instability <input type="radio"/> PTCA support <input type="radio"/> Unstable angina	<input type="radio"/> CPB wean <input type="radio"/> Prophylactic


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Unique patient identifier	<input style="width: 100%;" type="text"/>	
Date of surgery	<input style="width: 100%;" type="text"/>	dd/mm/yyyy
CPB and support continued ...		
Other support		
Intra-operative TOE	<input type="radio"/> No	<input type="radio"/> Yes
Intra-operative TOE: type	<input type="radio"/> Non-elective	<input type="radio"/> Elective
Intra-operative antifibrinolytic use	<input type="radio"/> No	<input type="radio"/> Yes
Intra-operative antifibrinolytic use: type	<input type="radio"/> Trasylol	<input type="radio"/> Other
	<input type="radio"/> Tranexamic acid	



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Unique patient identifier
Date of surgery dd/mm/yyyy

Coronary bypass

Intra-operative decision to graft coronary artery No Yes

IMA used No Yes

Which IMA used Left Right

Number of distal arterial grafts integer: 0-9

Number of IMA distal anastomoses integer: 0-6

Number of RA conduits harvested integer: 0-2

Number of radial distal anastomoses integer: 0-6

Number of vein distal anastomoses integer: 0-9

Number of GEPA distal anastomoses integer: 0-6

Were arterial T or Y grafts used No Yes

Total number of distal anastomoses integer: 0-30

Appendix

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Unique patient identifier
 Date of surgery dd/mm/yyyy

Aortic valve surgery

- Aortic valve procedure**
- Replacement
 - Repair / reconstruction without annuloplasty
 - Root reconstruction with valve conduit (Bentall procedure)
 - Root reconstruction with valve sparing (David procedure)
 - Resuspension aortic valve
 - Resection sub-aortic stenosis
 - Repair paravalvular leak
 - Valvotomy
 - Ross procedure
 - Inspection only
 - Decalcification of valve only

- Implant - type**
- None
 - Mechanical
 - Bioprosthesis
 - Autograft
 - Homograft / allograft
 - Ring / band

Implant - manufacturer's model number select from table

Implant - serial number select from table

Implant - size mm

- Explant - type**
- None
 - Mechanical
 - Bioprosthesis
 - Autograft
 - Homograft / allograft
 - Ring / band

Explant - manufacturer's model number select from table

Explant - serial number select from table

Explant - size mm

- Aortic stenosis**
- No
 - Yes

- Aortic regurgitation / insufficiency**
- None
 - Trivial
 - Mild
 - Moderate
 - Severe

- Aortic pathology / aetiology**
- Rheumatic
 - Congenital
 - Ischaemic
 - Idiopathic calcific
 - Myxomatous degen
 - Failed prior repair
 - Prosthetic valve failure
 - Peri-prosthetic leak
 - Prosthetic valve thrombosis
 - Active infection
 - Previous infection
 - Marfans
 - Annuloaortic ectasia
 - Other degenerative disease
 - Dissection
 - Tumour
 - Trauma
 - Iatrogenic
 - Other



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Unique patient identifier
Date of surgery dd/mm/yyyy

Mitral valve surgery

Mitral valve procedure

- Annuloplasty only
- Replacement
- Repair/reconstruction with annuloplasty
- Repair/reconstruction without annuloplasty
- Commissurotomy with annuloplasty ring
- Commissurotomy without annuloplasty ring
- Repair paravalvular leak
- Inspection only
- Decalcification of valve only

Implant - type

- None
- Mechanical
- Bioprosthesis
- Autograft
- Homograft/allograft
- Ring/band

Implant - manufacturer's model number select from table

Implant - serial number select from table

Implant - size mm

Explant - type

- None
- Mechanical
- Bioprosthesis
- Autograft
- Homograft/allograft
- Ring/band

Explant - manufacturer's model number select from table

Explant - serial number select from table

Explant - size mm

Mitral stenosis No Yes

Mitral regurgitation/insufficiency

- None
- Trivial
- Mild
- Moderate
- Severe

Mitral pathology/aetiology

- Functional or isolated annular dilataion
- Rheumatic
- Congenital
- Ischaemic
- Idiopathic calcific
- Myxomatous degen
- Failed prior repair
- Prosthetic valve failure
- Peri-prosthetic leak
- Prosthetic valve thrombosis
- Active infection
- Previous infection
- Marfans
- Other degenerative disease
- Tumour
- Trauma
- Iatrogenic
- Other

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Unique patient identifier
 Date of surgery dd/mm/yyyy

Tricuspid valve surgery

Tricuspid valve procedure	<input type="radio"/> Annuloplasty only <input type="radio"/> Replacement <input type="radio"/> Repair / reconstruction with annuloplasty <input type="radio"/> Repair / reconstruction without annuloplasty <input type="radio"/> Commissurotomy with annuloplasty ring <input type="radio"/> Commissurotomy without annuloplasty ring <input type="radio"/> Repair paravalvular leak <input type="radio"/> Valvectomy (no replacement) <input type="radio"/> Inspection only	
Implant - type	<input type="radio"/> None <input type="radio"/> Mechanical <input type="radio"/> Bioprosthesis	<input type="radio"/> Autograft <input type="radio"/> Homograft / allograft <input type="radio"/> Ring / band
Implant - manufacturer's model number	<input type="text"/> select from table	
Implant - serial number	<input type="text"/> select from table	
Implant - size	<input type="text"/> mm	
Explant - type	<input type="radio"/> None <input type="radio"/> Mechanical <input type="radio"/> Bioprosthesis	<input type="radio"/> Autograft <input type="radio"/> Homograft / allograft <input type="radio"/> Ring / band
Explant - manufacturer's model number	<input type="text"/> select from table	
Explant - serial number	<input type="text"/> select from table	
Explant - size	<input type="text"/> mm	
Tricuspid stenosis	<input type="radio"/> No <input type="radio"/> Yes	
Tricuspid regurgitation / insufficiency	<input type="radio"/> None <input type="radio"/> Trivial <input type="radio"/> Moderate <input type="radio"/> Mild <input type="radio"/> Severe	
Tricuspid pathology / aetiology	<input type="radio"/> Rheumatic <input type="radio"/> Active infection <input type="radio"/> Congenital <input type="radio"/> Previous infection <input type="radio"/> Ischaemic <input type="radio"/> Marfans <input type="radio"/> Idiopathic calcific <input type="radio"/> Other degenerative disease <input type="radio"/> Myxomatous degen <input type="radio"/> Tumour <input type="radio"/> Failed prior repair <input type="radio"/> Trauma <input type="radio"/> Prosthetic valve failure <input type="radio"/> Iatrogenic <input type="radio"/> Peri-prosthetic leak <input type="radio"/> Functional <input type="radio"/> Prosthetic valve thrombosis <input type="radio"/> Other	



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Unique patient identifier
Date of surgery dd/mm/yyyy

Pulmonary valve surgery

Pulmonary valve procedure	<input type="radio"/> Replacement <input type="radio"/> Repair / reconstruction without annuloplasty <input type="radio"/> Commissurotomy without annuloplasty ring <input type="radio"/> Repair paravalvular leak	
Implant - type	<input type="radio"/> None <input type="radio"/> Mechanical <input type="radio"/> Bioprosthesis	
	<input type="radio"/> Autograft <input type="radio"/> Homograft / allograft <input type="radio"/> Ring / band	
Implant - manufacturer's model number	<input type="text"/>	select from table
Implant - serial number	<input type="text"/>	select from table
Implant - size	<input type="text"/>	mm
Explant - type	<input type="radio"/> None <input type="radio"/> Mechanical <input type="radio"/> Bioprosthesis	
	<input type="radio"/> Autograft <input type="radio"/> Homograft / allograft <input type="radio"/> Ring / band	
Explant - manufacturer's model number	<input type="text"/>	select from table
Explant - serial number	<input type="text"/>	select from table
Explant - size	<input type="text"/>	mm
Pulmonary stenosis	<input type="radio"/> No <input type="radio"/> Yes	
Pulmonary regurgitation / insufficiency	<input type="radio"/> None <input type="radio"/> Trivial <input type="radio"/> Mild	
	<input type="radio"/> Moderate <input type="radio"/> Severe	
Pulmonary pathology / aetiology	<input type="radio"/> Rheumatic <input type="radio"/> Congenital <input type="radio"/> Ischaemic <input type="radio"/> Idiopathic calcific <input type="radio"/> Myxomatous degen <input type="radio"/> Failed prior repair <input type="radio"/> Prosthetic valve failure <input type="radio"/> Peri-prosthetic leak <input type="radio"/> Prosthetic valve thrombosis	
	<input type="radio"/> Active infection <input type="radio"/> Previous infection <input type="radio"/> Marfans <input type="radio"/> Other degenerative disease <input type="radio"/> Tumour <input type="radio"/> Trauma <input type="radio"/> Iatrogenic <input type="radio"/> Functional <input type="radio"/> Other	

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Unique patient identifier	<input type="text"/>	
Date of surgery	<input type="text"/>	dd/mm/yyyy
Post-operative data		
RBC blood bank products	<input type="radio"/> No <input type="radio"/> Yes	
Non-RBC blood bank products	<input type="radio"/> No <input type="radio"/> Yes	
Peri-operative transfusion: bank RBC	<input type="text"/>	units
Peri-operative transfusion: platelets	<input type="text"/>	units
Peri-operative transfusion: Novo 7	<input type="text"/>	units
Peri-operative transfusion: FFP	<input type="text"/>	units
Peri-operative transfusion: Cryo	<input type="text"/>	units
ICU admission: date and time	<input type="text"/>	dd/mm/yyyy
Extubation: date and time	<input type="text"/>	dd/mm/yyyy
ICU discharge: date and time	<input type="text"/>	dd/mm/yyyy
Readmitted to ICU	<input type="radio"/> No <input type="radio"/> Yes	
Reintubated	<input type="radio"/> No <input type="radio"/> Yes	
Reintubation: date and time	<input type="text"/>	dd/mm/yyyy
Reextubation: date and time	<input type="text"/>	dd/mm/yyyy
ICC loss (first 4 hours post surgery)	<input type="text"/>	dd/mm/yyyy
Returned to theatre		
Return to theatre	<input type="radio"/> No <input type="radio"/> Yes	
Reason for re-operation	<input type="checkbox"/> Valve dysfunction	<input type="checkbox"/> Sternal infection
	<input type="checkbox"/> Bleeding /tamponade	<input type="checkbox"/> Other cardiac
	<input type="checkbox"/> Graft occlusion	<input type="checkbox"/> Other non-cardiac



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Unique patient identifier
Date of surgery dd/mm/yyyy

Complications

Renal and neurological complications

New renal failure	<input type="radio"/> No	<input type="radio"/> Yes
Haemofiltration	<input type="radio"/> No	<input type="radio"/> Yes
Highest post-op creatinine	<input type="text"/>	µmol l ⁻¹
Perioperative cardiogenic shock	<input type="radio"/> No	<input type="radio"/> Yes
New neurological status	<input type="radio"/> No	<input type="radio"/> Yes
Stroke permanent	<input type="radio"/> No	<input type="radio"/> Yes
Stroke transient	<input type="radio"/> No	<input type="radio"/> Yes
New continuous coma (≥24 hours)	<input type="radio"/> No	<input type="radio"/> Yes

Cardiac complications

Perioperative AMI	<input type="radio"/> No	<input type="radio"/> Yes
Cardiac inotrope use: >4 hours post-operatively	<input type="radio"/> No	<input type="radio"/> Yes
Cardiac inotrope use: low cardiac output syndrome	<input type="radio"/> No	<input type="radio"/> Yes
Cardiac inotrope use: low SVR syndrome	<input type="radio"/> No	<input type="radio"/> Yes
New cardiac arrhythmia	<input type="radio"/> No	<input type="radio"/> Yes
New heart block (requiring PPM)	<input type="radio"/> No	<input type="radio"/> Yes
New other brady arrhythmia (requiring PPM)	<input type="radio"/> No	<input type="radio"/> Yes
Cardiac arrest	<input type="radio"/> No	<input type="radio"/> Yes
New atrial arrhythmia (requiring Rx)	<input type="radio"/> No	<input type="radio"/> Yes
New ventricular tachycardia	<input type="radio"/> No	<input type="radio"/> Yes

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Unique patient identifier
 Date of surgery dd/mm/yyyy

Complications continued ...

Pulmonary, infection, vascular and other complications

Prolonged ventilation >24 hours	<input type="radio"/> No	<input type="radio"/> Yes
Pulmonary embolism	<input type="radio"/> No	<input type="radio"/> Yes
Pneumonia	<input type="radio"/> No	<input type="radio"/> Yes
Reintubation and ventilation	<input type="radio"/> No	<input type="radio"/> Yes
Deep sternal wound infection	<input type="radio"/> No	<input type="radio"/> Yes
Deep thoracotomy wound infection	<input type="radio"/> No	<input type="radio"/> Yes
Septicaemia	<input type="radio"/> No	<input type="radio"/> Yes
Aortic dissection (complication)	<input type="radio"/> No	<input type="radio"/> Yes
Acute limb ischaemia	<input type="radio"/> No	<input type="radio"/> Yes
Anti-coagulant complication	<input type="radio"/> No	<input type="radio"/> Yes
GIT complication	<input type="radio"/> No	<input type="radio"/> Yes
Multi-system failure	<input type="radio"/> No	<input type="radio"/> Yes



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Unique patient identifier

Date of surgery dd/mm/yyyy

Discharge / mortality

Discharge

Home
 Hospital in the home
 Rehabilitation unit/hospital
 Local or referring hospital
 Hospital mortality

Date of discharge dd/mm/yyyy

Mortality post discharge No Yes

Mortality date dd/mm/yyyy

Mortality location

Operating room
 Hospital
 Home
 Other facility

Mortality: primary cause

Cardiac
 Neurological
 Renal
 Vascular
 Infection
 Respiratory failure
 Multisystem failure
 Pulmonary embolism
 Aortic dissection
 Valvular
 Other
 Unknown

Mortality: subsequent cause

Cardiac
 Neurological
 Renal
 Vascular
 Infection
 Respiratory failure
 Multisystem failure
 Pulmonary embolism
 Aortic dissection
 Valvular
 Other
 Unknown

Cognisant patient withdraws from treatment No Yes

Readmission

Readmitted ≤30 days from surgery No Yes

Reason for readmission

Anticoagulant complication
 Arrhythmia
 Congestive heart failure
 Valve dysfunction
 Pericardial effusion
 Cardiac tamponade
 Deep sternal infection
 Other incisional complication
 Respiratory complication including pneumonia
 Myocardial infarction
 Recurrent angina
 Other complication related to cardiac surgery
 Other readmission unrelated to cardiac surgery

