

Health NZ



**New Zealand Cardiac Surgery
National Report**

2016

Preface**New Zealand Cardiac Surgical Annual Report 2016**

This report of the New Zealand National Cardiac Surgery Registry (NZCS) covers the period of 1 January 2016 to 31 December 2016.

The report includes all cardiac surgical procedures undertaken at the 5 District Health Boards (DHBs) performing publicly-funded cardiac surgery in New Zealand (Auckland, Waikato, Capital and Coast, Canterbury, Southern).

This is our second annual report of the National Cardiac surgical services in New Zealand. The report has been an accomplishment for all involved to be proud of. It is encouraging to note that data completeness has improved for 2016 across all units. This is a credit to the time and diligence of all members of the cardiac surgical units at the 5 DHBs. With the implementation of mobile devices in 2017 we expect the processes to become further streamlined. I take this opportunity to thank all the effort of the different teams involved in making this possible at both a National and Local level. The data in itself is a testimony to the quality of cardiac surgical service being provided around the country.

The data presents total volume of cardiac surgery procedures in New Zealand. The risk factors associated with cardiac disease and the performance and outcomes of the most common procedures performed by a cardiac surgeons, namely coronary artery bypass surgery (CABG) and aortic valve replacement (AVR).

The report examines the impact of variables (risk factors) on surgical outcomes and compares these with international registries. In doing this we are able to benchmark New Zealand outcomes to internationally accepted standards.

As stated previously, the registry data is in its early years. This being the second year of its compilation and any inference from the data needs to be considered with caution. Bearing this in mind, we are proactively analysing quality measures in the data and hope this will continue to improve quality of care for our patients on an ongoing basis. Areas presently under consideration include minimising infection rates (already low), improve early discharge care strategies, blood and blood product utilization.

The data so far collected when compared with International data, suggests that New Zealand provides a high standard of Cardiac surgical services. For the data to be of any statistical significance we accept a minimum of 3 years of data needs to be available.

I would like to thank the members of the steering and implementation group, the data managers, DHB information services and the company Dendrite Clinical Systems for their effort and persistence.

The registry project acknowledges the Ministry of Health's (MOH) support, guidance commitment and funding which has made this important project a reality. The document would not be complete without acknowledging the work put in by Charlotte Allin.



Harsh Singh
Chairman



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Consumer advocate summary

I am the Consumer Representative who has the honour of advocating on your behalf, expressing your voice to the National Cardiac Surgical Clinical Network Governance Group.

In our first report, published last year, I indicated several conflicting ideas regarding its requirements and value for you, as a consumer. Initially, it was suggested a performance league table be included, however, this is not very useful for consumers. The key reason is because an escalation process is in use. What this means, is that more complicated and seriously ill patients are referred to cardiac surgeons with greater experience.

Due to this treatment methodology, a performance league table would not be the fairest way to compare results. For you, as a consumer, the fundamental service delivery issue remains. That is, being treated by the *right person at the right time, at the right place* to meet your needs.

This year, we publish our second report and the figures are very similar to those of last year. These findings confirm and validate the consistency of the nationwide service. Recorded numbers are still low, which can have a marked effect on final percentages.

However, what astounds me is the higher survival rate for patients who receive this service. When I joined the network, after listening to stories in my community, I had expected much lower survival rates. Now, I know first-hand, that the five centres excel when compared with international statistics.

Thank you for taking the time to read our latest report.

Kevin Murray

Consumer Representative, National Cardiac Surgical Clinical Network Governance Group



Data presentation

- The data has been compiled for the period 01 January-31 December 2016.
- The report 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 standard operations included are coronary bypass grafting (CABG), and aortic valve replacement (AVR) these account for over 65% of the workload of all cardiac surgical units.
- The dataset aims to enable benchmarking with international standards by using identical definitions so a realistic comparison can be made.
- This dataset is a single year's results. It does not take into account retrospective years of surgical performance and as a result fails to recognise surgical experience and the unit's total performance. We suggest caution in interpreting performances of units. Overall with limited data available it is reassuring to see all units and surgeons are performing above the accepted international bench marks.
- The National cardiac surgical network with the MOH help was set up as a working group to review public funded cardiac surgery and equity of care at a national level. The group recognised the lack of a uniform registry to review outcomes and performance. With an MOH initiative in 2008-2009 funding was allocated for development of a national registry.
- Data is analysed independently by Dendrite Clinical Systems a specialist supplier of clinical databases, analysis software, consultancy and publishing services for the international healthcare sector.
- This is the second national report of the registry. It shows an analysis of the performance of the 5 DHB performing cardiac surgery for the one year period commencing 1 January 2016.

Overview of people who had cardiac surgery

In the 12 months period (2016) a total of 2,807 cardiac operations were performed across the 5 District Health Boards (DHB) undertaking Cardiac surgery: Auckland DHB, Waikato DHB, Capital and Coast DHB, Canterbury DHB, Southern DHB. This was 43 extra procedures for the year compared to the previous year.

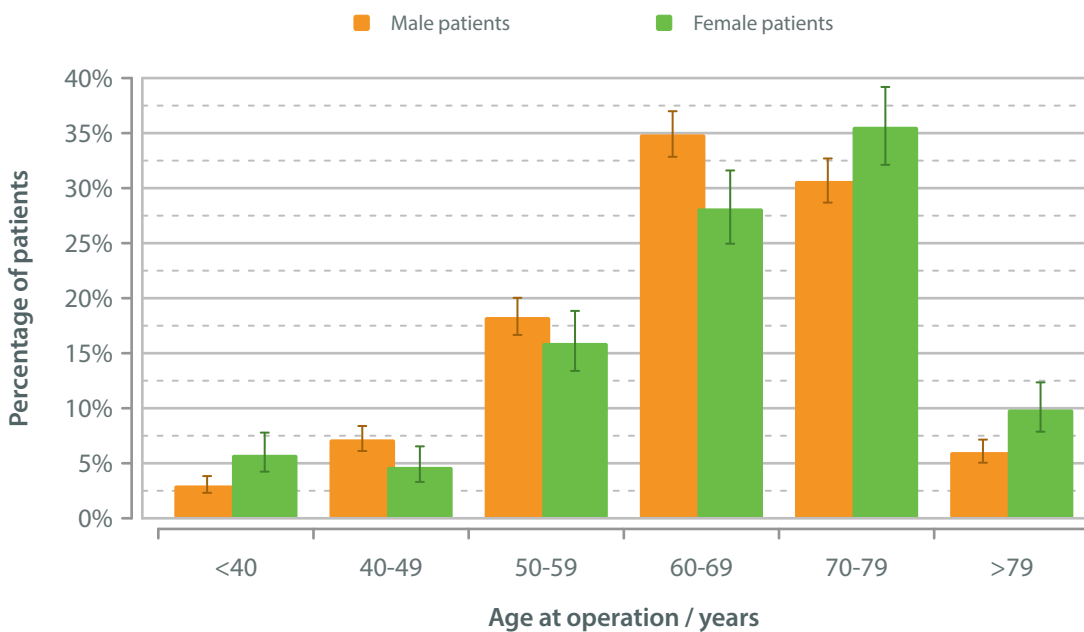
The presentation of over 70% of the patients was after 60 years of age. With men presenting at a slightly younger age than women. Heart disease was seen to be more common in men who accounted for 75% of the total number (Table 1, Fig. 1). This ratio distribution matches international figures and shows a similar pattern from the year before.

Table 1. All cardiac surgery patients in 2016: age and gender

Age at operation / years	Gender		
	Male	Female	All
<40	62	42	104
40-49	149	34	183
50-59	380	116	496
60-69	725	205	930
70-79	637	259	896
>79	125	72	197
Unspecified	1	0	1
All	2,079	728	2,807

Fig. 1

All cardiac surgery patients: Age & gender distributions; calendar year 2016 (n=2,806)





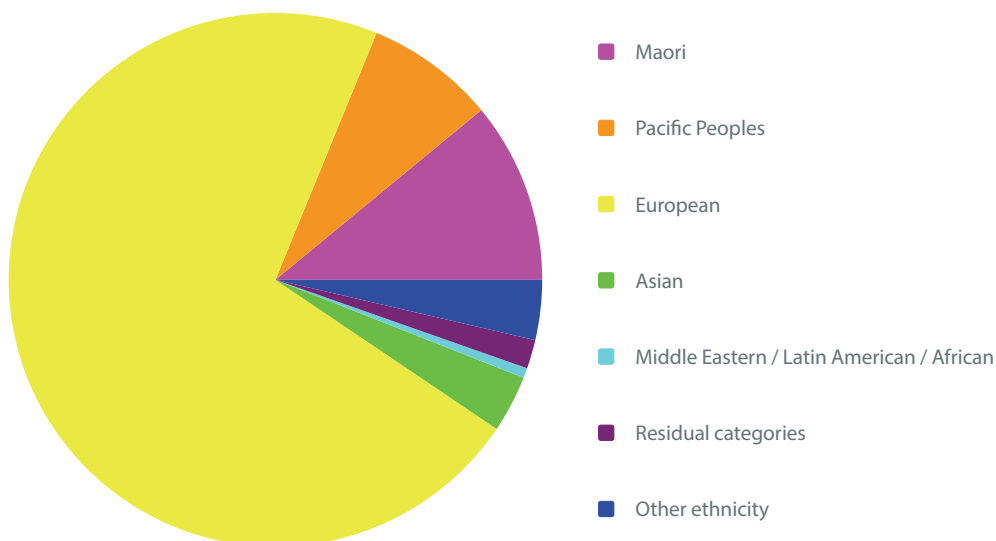
Ethnicity

The ethnicity spectrum remains similar as before. With roughly 20% of the population being of Maori / Pacific Islanders (Census 2013 shows Maori / Pacific account for 22.3% of total population). Considering the high incidence of heart disease in this sub population, effort may need to be put into health care for the sub-group.

Table 2. Ethnicity of patients undergoing cardiac surgery in 2016

	Count	Percentage
Maori	308	11.0%
Pacific Peoples	220	7.8%
European	2,014	71.7%
Asian	97	3.5%
Middle Eastern / Latin American / African	17	0.6%
Residual categories	49	1.7%
Other ethnicity	102	3.6%
Unspecified	0	
All	2,807	

Fig. 2 All cardiac surgery patients: Ethnicity; calendar year 2016 (n=2,807)



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.

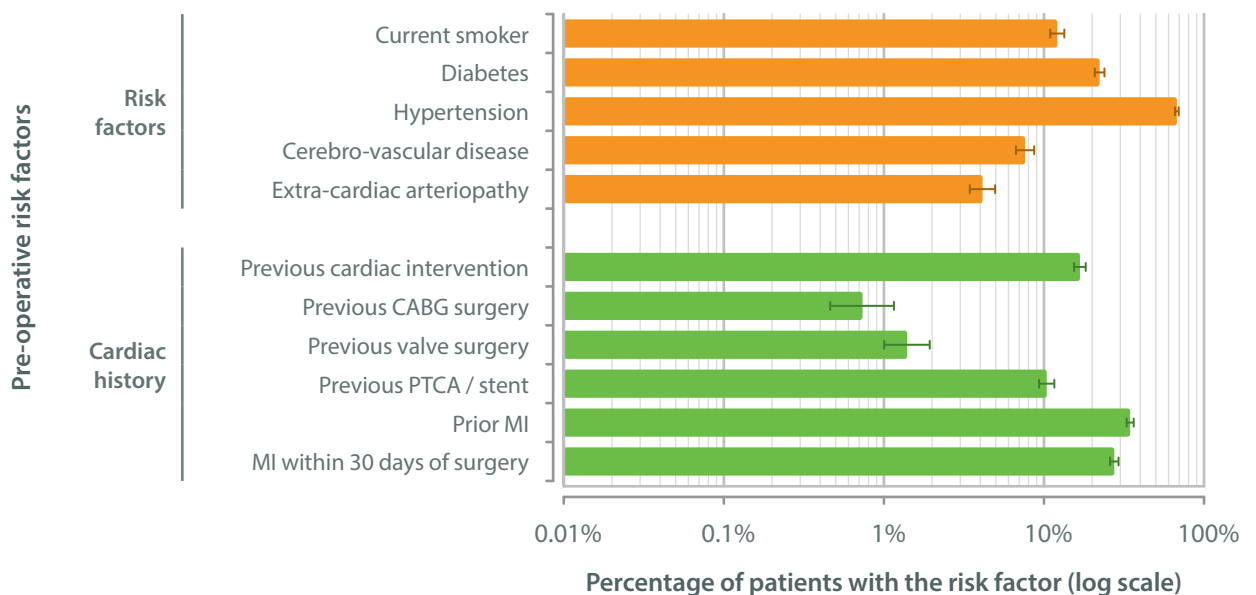
- One in eight patients were still smoking at the time of surgery.
- Approximately one-quarter of the patients had diabetes.
- Over 50% of the patients had high blood pressure.

Table 3. All patients in 2016: Pre-operative risk factors

	Risk factor present			Percentage with the risk factor	
	No	Yes	Unspecified		
Risk factors	Current smoker	2,466	340	1	12.1%
	Diabetes	2,181	625	1	22.3%
	Hypertension	907	1,899	1	67.7%
	Cerebro-vascular disease	2,592	214	1	7.6%
	Extra-cardiac arteriopathy	2,690	116	1	4.1%
Cardiac history	Previous cardiac intervention	2,335	471	1	16.8%
	Previous CABG surgery	2,700	20	87	0.7%
	Previous valve surgery	2,682	38	87	1.4%
	Previous PTCA / stent	2,514	292	1	10.4%
	Prior MI	1,836	970	1	34.6%
	Prior MI within 30 days of surgery	2,033	772	2	27.5%

Fig. 3

**All cardiac surgery patients:
Risk factors; calendar year 2016**





Types of operations performed

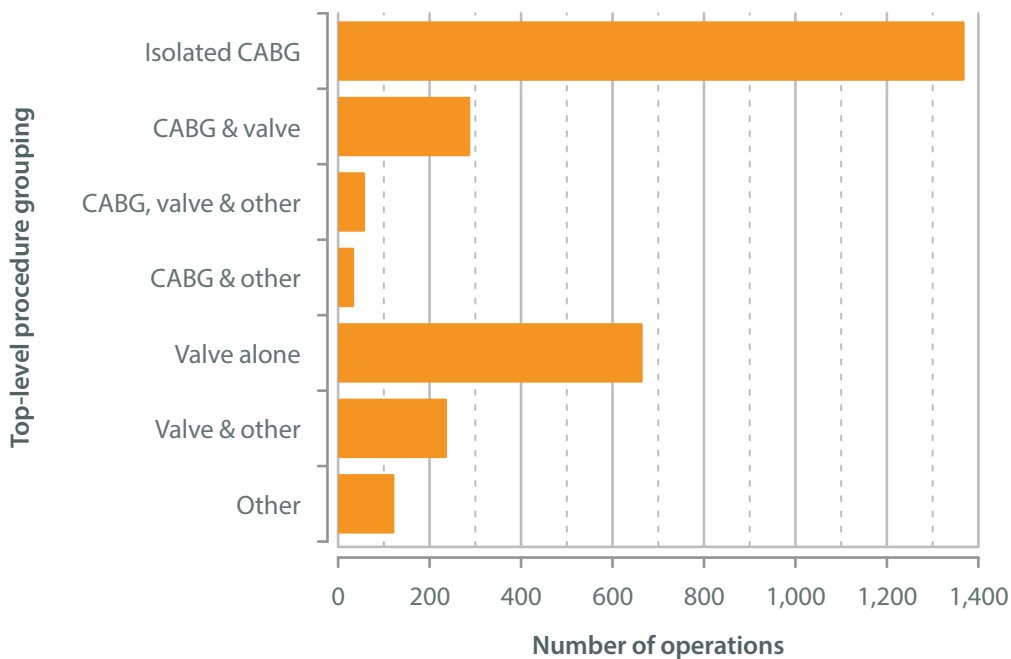
- Isolated coronary artery bypass accounted for 49.3% of the total volume of cases.
- Isolated heart valve operations were 24.5%.
- Combined valve and coronary artery bypass 10.4%.
- Approximately 15% of cases were for Other, less common procedures.

Comparing the two years 2015 / 2016, the distribution of cases performed remains similar. Suggesting no early impact of TAVI on total AVR procedures. The year 2017 has seen an increase in TAVI procedures nationally and will be interesting to note the impact of this in future years.

Table 4. Procedures performed in 2016

	Count	Percentage
CABG	1,370	49.3%
CABG & valve	289	10.4%
CABG, valve & other	59	2.1%
CABG & other	35	1.3%
Valve alone	666	24.0%
Valve & other	238	8.6%
Other	123	4.4%
Unspecified	27	
All	2,807	

Fig. 4 **Operations performed in the calendar year 2016**



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 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.

The operation is the most commonly performed operation by a Cardiac surgeon. In the year 2016 a total of 1,360 patients underwent a publicly-funded isolated CABG operation (49.3%) of the total volume of cardiac surgery. (Table 4). The volumes of the procedure is consistent over the two years audited.

Table 5. First-time isolated CABG in 2016: Body Mass Index

	Count	Percentage
<20.0	22	1.6%
20.0-24.9	266	19.6%
25.0-29.9	545	40.1%
30.0-34.9	354	26.0%
35.0-39.9	121	8.9%
40.0-44.9	40	2.9%
>44.9	9	0.7%
Unspecified	3	0.2%
All	1,360	

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. 5) 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.

BMI classifications

Ministry of Health New Zealand. Body size. Retrieved from: <http://www.health.govt.nz/our-work/populations/maori-health/tatau-kahukura-maori-health-statistics/nga-tauwehe-tuono-me-te-marumaruru-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



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

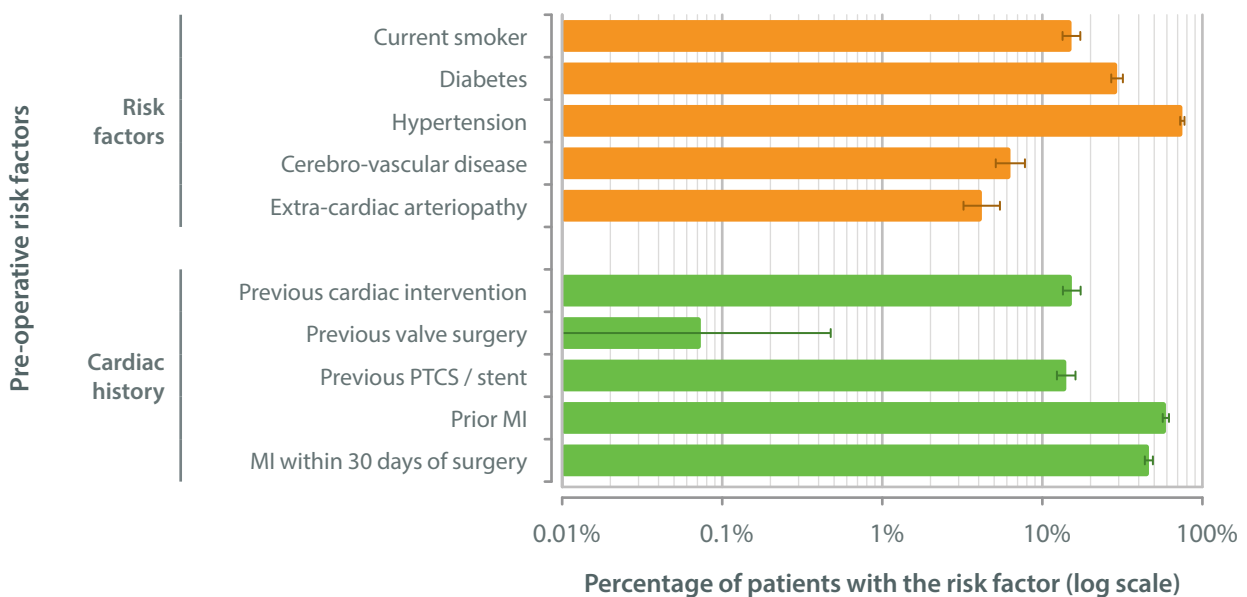
- One in six patients (15.2%) were current smokers.
- One in four (29.3%) were diabetic.
- One in three (38.7%) were obese and (12.7%) had a BMI of greater than 35.
- More than half the patients were obese.
- One in nine patients were morbidly obese.

Table 7. First-time isolated CABG in 2016: Pre-operative risk factors

	Risk factor present			Percentage with the risk factor	
	No	Yes	Unspecified		
Risk factors	Current smoker	1,153	207	0	15.2%
	Diabetes	961	399	0	29.3%
	Hypertension	340	1,020	0	75.0%
	Cerebro-vascular disease	1,274	86	0	6.3%
	Extra-cardiac arteriopathy	1,303	57	0	4.2%
Cardiac history	Previous cardiac intervention	1,152	208	0	15.3%
	Previous valve surgery	1,359	1	0	0.1%
	Previous PTCA / stent	1,168	192	0	14.1%
	Prior MI	555	805	0	59.2%
	Prior MI within 30 days of surgery	729	630	1	46.4%

Fig. 5

First-time isolated CABG: Risk factors; calendar year 2016



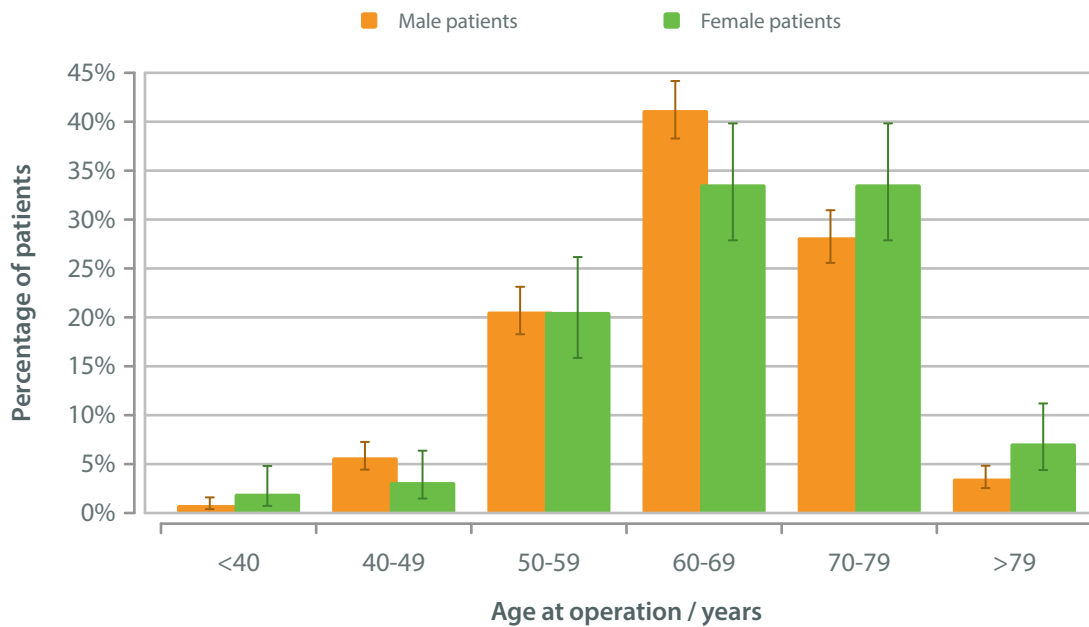
Isolated CABG

Table 8. First-time isolated CABG in 2016: age and gender

Age at operation / years	Gender		
	Male	Female	All
<40	9	5	0
40-49	63	8	71
50-59	228	52	280
60-69	456	85	541
70-79	312	85	397
>79	39	18	57
Unspecified	0	0	0
All	1,107	253	1,360

Fig. 6

First-time isolated CABG: Age & gender distributions; calendar year 2016 (n=1,360)



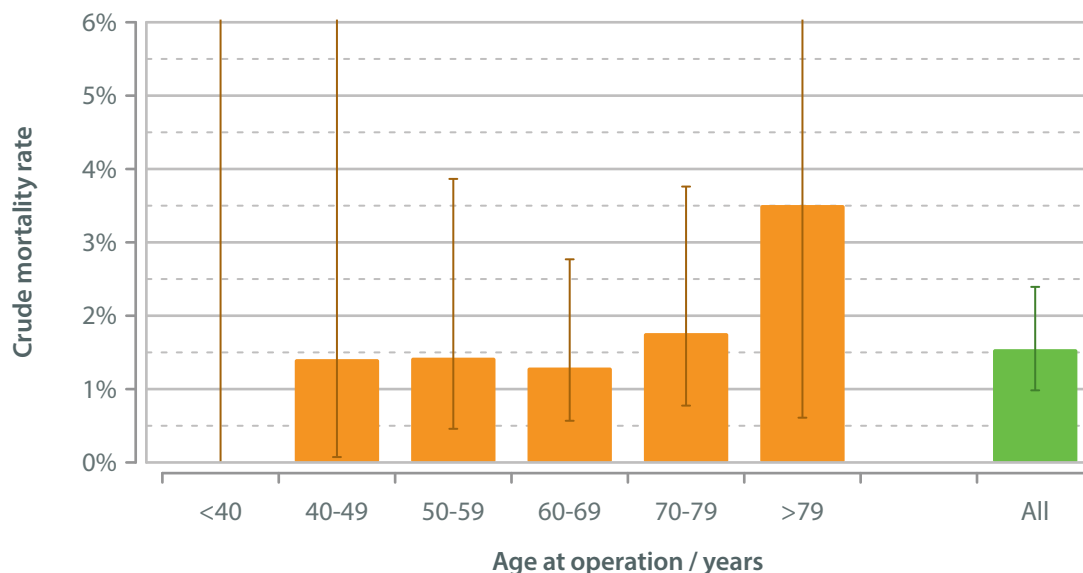


The overall survival results of isolated CABG operations nationwide is well within the International standard benchmark of care at 98.5%¹. This is similar to the year before. The most common presentation of these patients is between 50 to 75 years of age, which accounts for over 85% of the total volume. 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 5: 1.

Table 9. First-time isolated CABG in 2016: age and in-hospital mortality

Age at operation / years	In-hospital mortality			Mortality rate (95% CI)
	No	Yes	All	
<40	14	0	14	0.0% (0.0-19.3%)
40-49	70	1	71	1.4% (0.1-8.7%)
50-59	276	4	280	1.4% (0.5-3.9%)
60-69	534	7	541	1.3% (0.6-2.8%)
70-79	390	7	397	1.8% (0.8-3.8%)
>79	55	2	57	3.5% (0.6-13.2%)
Unspecified	0	0	0	NA
All	1,339	21	1,360	1.5% (1.0-2.4%)

Fig. 7 First-time isolated CABG: In-hospital mortality and age; calendar year 2016 (n=1,360)

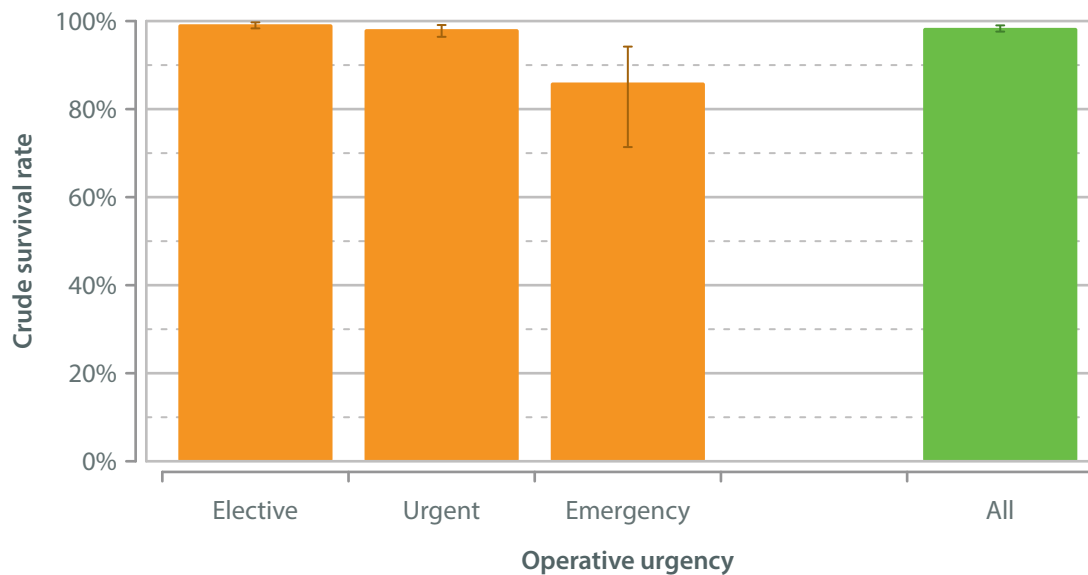


1. <http://anzscts.org/wp-content/uploads/2016/12/ANZSCTS-National-Annual-Report-2015.pdf>

Table 10. First-time isolated CABG in 2016: operative urgency and in-hospital survival

		In-hospital survival			
		Yes	No	All	Survival rate (95% CI)
Operative urgency	Elective	820	6	826	99.3% (98.3-99.7%)
	Urgent	482	9	491	98.2% (96.4-99.1%)
	Emergency / salvage	37	6	43	86.0% (71.4-94.2%)
	All	1,339	21	1,360	98.5% (97.6-99.0%)

Fig. 8 First-time isolated CABG: In-hospital survival rates; calendar year 2016 (n=1,360)



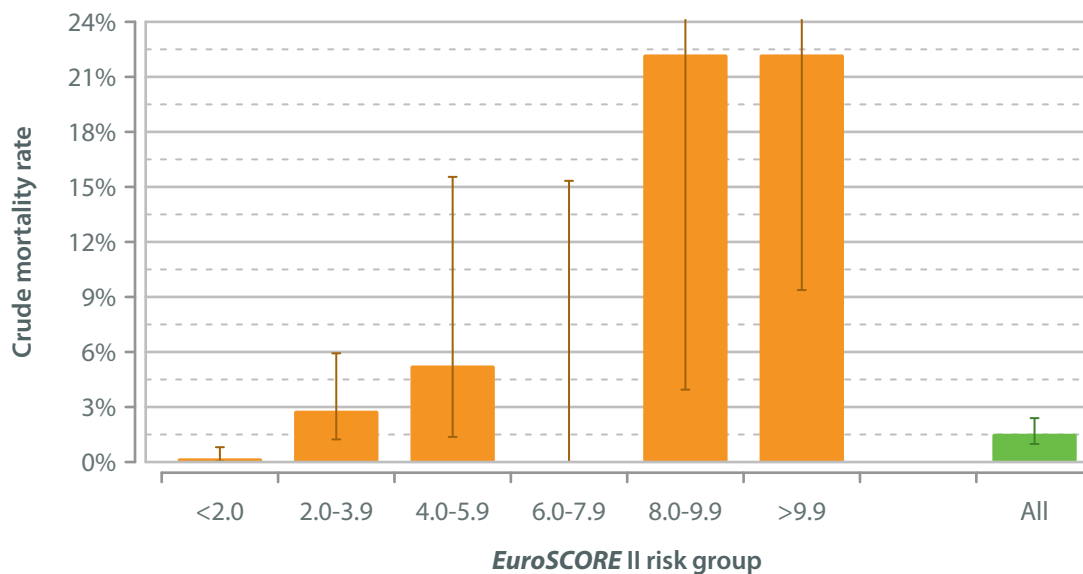


Majority of the deaths were in the high risk group of patients. As expected the salvage and emergency procedures had a higher mortality. The risk of the patient was based on a **EuroSCORE**². This takes into account risk factors associated with coronary artery disease and the higher the score the greater risk of morbidity and mortality post-surgery (e.g., **EuroSCORE** 6.0-7.9). **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.

Table 11. First-time isolated CABG in 2016: **EuroSCORE II** risk score and in-hospital mortality

	In-hospital mortality			
	No	Yes	All	Mortality rate (95% CI)
EuroSCORE II <2.0	991	2	993	0.2% (0.0-0.8%)
2.0-3.9	243	7	250	2.8% (1.2-5.9%)
4.0-5.9	54	3	57	5.3% (1.4-15.5%)
6.0-7.9	18	0	18	0.0% (0.0-15.3%)
8.0-9.9	7	2	9	22.2% (3.9-59.8%)
>9.9	21	6	27	22.2% (9.4-42.7%)
Unspecified	5	1	6	16.7% (0.9-63.5%)
All	1,339	21	1,360	1.5% (1.0-2.4%)

Fig. 9 First-time isolated CABG: In-hospital mortality and pre-operative risk; calendar year 2016



2. 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.

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 2016 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 2016 was 23 hours. Time spent in ICU is determined by how quickly the patients recover which is impacted by core morbidity 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 literature ¹.

Table 12. First-time isolated CABG in 2016: hospital resource utilisation

		No	Yes	Rate
Same day admission		1,305	55	4.0% (3.1-5.3%)
Resource utilisation		Count	Median	Inter-quartile range
	Ventilation time / hours	1,352	6.0	4.0-10.0
	Time on ICU / hours	1,350	23.0	20.0-44.0
	Post-operative stay / days	1,356	6.0	5.0-7.0
	Hospital stay / days	1,357	9.0	7.0-15.0

1. <http://anzscts.org/wp-content/uploads/2016/12/ANZSCTS-National-Annual-Report-2015.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.

Of note was the slight increase in sternal wound infection. Which in terms of numbers is small however to ensure continued improved quality outcomes this has triggered a review by the surgical site infection improvement programme (SSII) and surgical processes have been analysed and continue to be monitored at all sites. A slight increase in return to theatre for bleeding was noticed. It is important that we carefully interpret this data as the numbers are small and not statistically significant. They still compare favourably with international standard benchmarks. 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 2016: complications

		Complication			
		No	Yes	Unspecified	Rate (95% CI)
In-hospital	Deep sternal wound infection	1,350	10	0	0.7% (0.4-1.4%)
	Any return to theatre	1,301	59	0	4.3% (3.3-5.6%)
	Return to theatre for bleeding	1,324	36	0	2.6% (1.9-3.7%)
30-day	Readmission	1,227	133	0	9.8% (8.3-11.5%)
	Deep sternal wound infection	1,344	16	0	1.2% (0.7-1.9%)

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.

Surgical aortic valve replacement 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/TAVR) are performed almost exclusively for aortic stenosis in New Zealand in a much smaller patient population. The outcomes of TAVI/TAVR are not currently discussed in this report. All patients presented here underwent standard open surgical aortic valve replacement performed by a cardiac surgical team.

Table 14. Valve surgery in 2016

	Top-level procedure classification	
	Valve alone	CABG & valve
Aortic valve alone	398	216
Mitral valve alone	162	53
Aortic & mitral valves	36	9
Mitral & tricuspid valves	49	9
Others	20	2
Unspecified	1	0
All	666	289

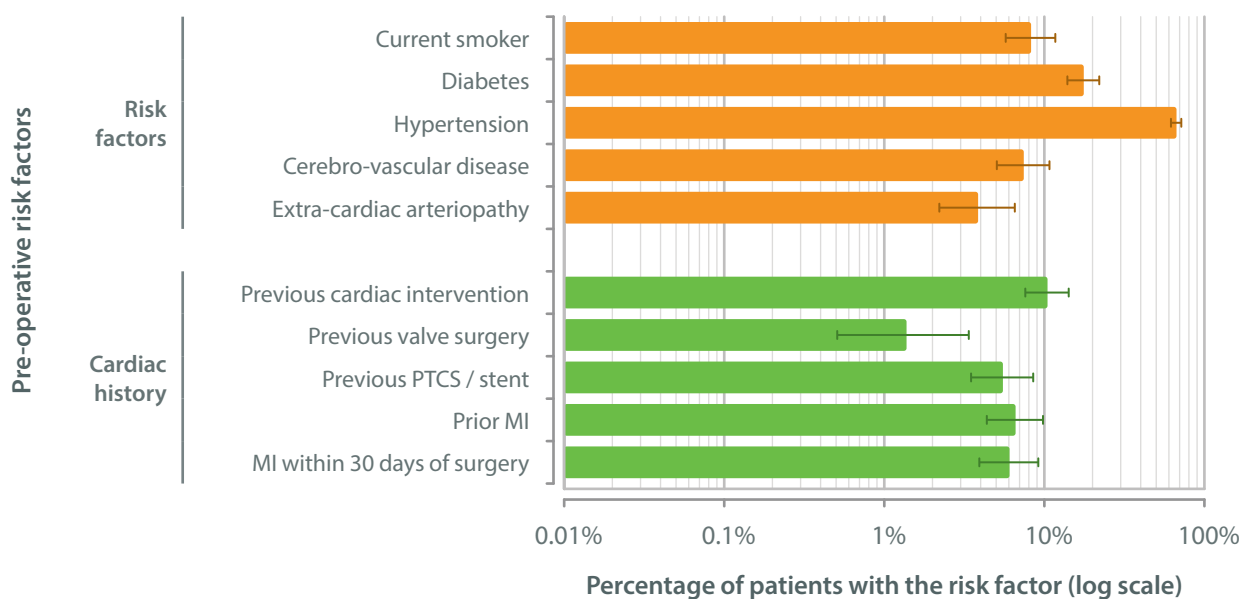


Table 15. First-time isolated AVR in 2016: Pre-operative risk factors

	Risk factor present			Percentage with the risk factor	
	No	Yes	Unspecified		
Risk factors	Current smoker	333	30	0	8.3%
	Diabetes	299	64	0	17.6%
	Hypertension	120	243	0	66.9%
	Cerebro-vascular disease	336	27	0	7.4%
	Extra-cardiac arteriopathy	349	14	0	3.9%
Cardiac history	Previous cardiac intervention	325	38	0	10.5%
	Previous CABG surgery	358	5	0	1.4%
	Previous PTCA / stent	343	20	0	5.5%
	Prior MI	339	24	0	6.6%
	Prior MI within 30 days of surgery	341	22	0	6.1%

Fig. 10

**First-time isolated AVR:
Risk factors; calendar year 2016**



In the New Zealand registry 363 isolated first time AVRs have been performed which is approximately 13% of the overall surgical volume. This is an increase of 35 isolated aortic valve procedures compared to that reported in 2015.

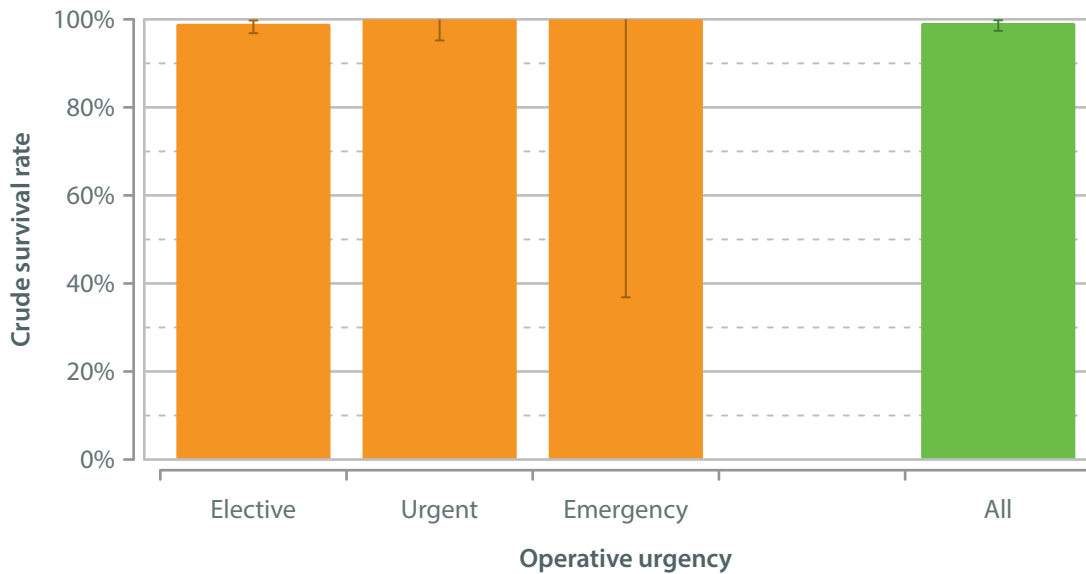
Most patients undergo surgery in a planned or elective fashion with smaller numbers undergoing urgent, emergency or salvage surgery. As expected a significant number of the patients have additional cardiovascular risk factors including 17.6% diabetics, 66.9% with hypertension and 7.4% having had a previous cardiovascular intervention (Table 15).

Aortic valve surgery

Table 16. First-time isolated AVR in 2016: operative urgency and in-hospital survival

	In-hospital survival			
	Yes	No	All	Survival rate (95% CI)
Operative urgency				
Elective	296	3	299	99.0%(96.8-99.7%)
Urgent	61	0	61	100.0% (95.2-100.0%)
Emergency / salvage	3	0	3	100.0% (36.8-100.0%)
All	360	3	363	99.2% (97.4-99.8%)

Fig. 11 First-time isolated AVR: In-hospital survival rates; calendar year 2016 (n=363)



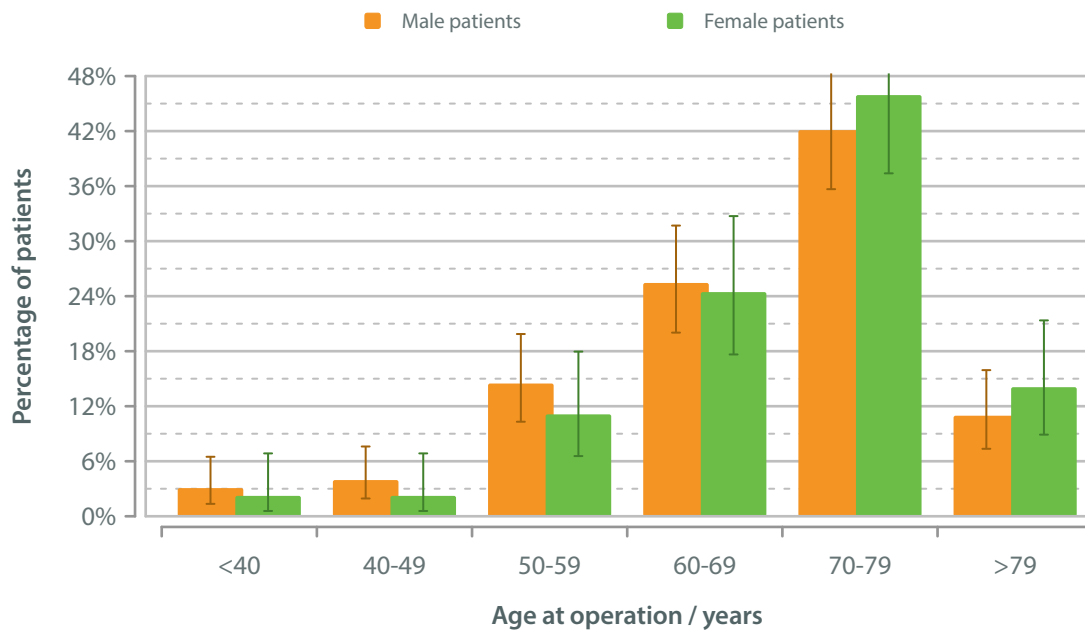


Aortic valve surgery may be required because of either leakage of the valve (aortic regurgitation) or blockage of the valve (aortic stenosis). These conditions can occur for a variety of reasons, the most common being degenerative age related calcification or hardening of the valve. Dysfunction of the 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. 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. 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 2:1.

Table 17. First-time isolated AVR in 2016: age and gender

Age at operation / years	Gender		All
	Male	Female	
<40	7	3	10
40-49	9	3	12
50-59	33	15	48
60-69	58	33	91
70-79	96	62	158
>79	25	19	44
Unspecified	0	0	0
All	228	135	363

Fig. 12 First-time isolated AVR: Age & gender distributions; calendar year 2016 (n=363)



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. In the 2015 report Auckland DHB were unable to generate **EuroSCORE II** data as at that time they did not collect all of the necessary data fields to enable risk modelling to be completed. The 2016 report has **EuroSCORE II** generated for all but 6 of 363 patients and gives a better reflection of risk adjusted outcomes for the entire New Zealand surgical population.

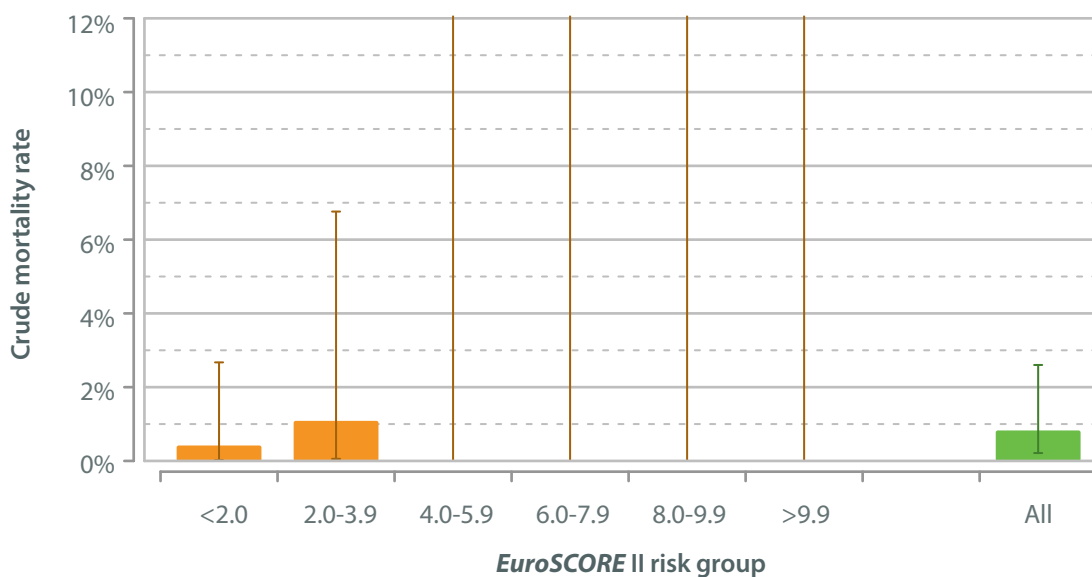
Table 18 shows the distribution of risk profiles in patients undergoing isolated AVR and the observed mortality rate. The overall observed mortality for isolated AVR in New Zealand was extremely low (0.8%), which is favourable when compared to internationally accepted outcomes. The 2015 ANZCTS publication of surgical outcomes (reference in footnote) reported a 1.8% mortality for isolated AVR in a similar cohort of patients.

As can be seen the majority of NZ patients (239) are in the low risk category (**EuroSCORE II** <2% predicted mortality) with an observed mortality of 0.4% in low risk patients.

Table 18. First-time isolated AVR in 2016: **EuroSCORE II** risk score and in-hospital mortality

	In-hospital mortality			
	No	Yes	All	Mortality rate (95% CI)
EuroSCORE II <2.0	238	1	239	0.4% (0.0-2.7%)
2.0-3.9	91	1	92	1.1% (0.1-6.8%)
4.0-5.9	18	0	18	0.0% (0.0-15.3%)
6.0-7.9	1	0	1	0.0% (0.0-95.0%)
8.0-9.9	3	0	3	0.0% (0.0-63.2%)
>9.9	4	0	4	0.0% (0.0-52.7%)
Unspecified	5	1	6	16.7% (0.9-63.5%)
All	360	3	363	0.8% (0.2-2.6%)

Fig. 13 First-time isolated AVR: In-hospital mortality and age;calendar year 2016 (n=357)





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 was 1 death recorded in patients with a **EuroSCORE II** >2 and one deceased patient who did not have sufficient information to calculate **EuroSCORE**.

As the registry grows we will be able to make more accurate assessment of outcomes in these higher risk cohorts.

Major morbidity compares favourably to international reported results². This includes low incidence of deep sternal wound infection rates (NZ 0.6% versus ANZCTS 0.6%), return to theatre for bleeding (NZ 2.6% versus ANZCTS 3.7%).

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 2016: hospital resource utilisation

Resource utilisation	No	Yes	Rate
	Count	Median	Inter-quartile range
Same day admission	346	17	4.7% (2.8-7.5%)
Ventilation time / hours	358	5.0	4.0-9.0
Time on ICU / hours	357	23.0	20.0-34.0
Post-operative stay / days	363	6.0	5.0-8.0
Hospital stay / days	363	8.0	7.0-12.0

Table 20. Isolated aortic valve surgery in 2016: complications

	Complication				
	No	Yes	Unspecified	Rate (95% CI)	
In-hospital	Deep sternal wound infection	360	2	1	0.6% (0.1-2.2%)
	Any return to theatre	342	21	0	5.8% (3.7-8.8%)
	Return to theatre for bleeding	353	10	0	2.8% (1.4-5.2%)
30-day	Readmission	330	33	0	9.1% (6.4-12.6%)
	Deep sternal wound infection	361	2	0	0.6% (0.1-2.2%)

Summary

- The New Zealand Cardiac Surgery Registry has been created to record and provide analysis of publicly funded cardiac surgical procedures.
- In our second year of data collection we have continued to bench mark against the ANZCTS dataset and used **EuroSCORE II** as the risk stratification tool.
- We note that the risk adjusted outcomes and other measures of quality of care (ventilator time, ICU and hospital stay) for all five units for the two most common procedures (CABG and AVR) performed by adult cardiac surgeons compare well with internationally accepted standards.
- The spectrum of different operations performed is not dissimilar to other developed countries.
- Other smaller volume procedure will be able to be analysed once 5 years of data has been compiled and statistically significant volumes are available.
- The New Zealand Cardiac Surgery Registry will allow us in the future to research improved quality measures. We are presently working with the Surgical Site Infection Improvement Programme (SSII) HQSCNZ.
- Plans are underway to include Transcatheter Aortic Valve Implantation (TAVI) data within the cardiac surgical registry.
- Uniformity in Mortality and Morbidity (M and M) audit presentations will allow all cardiac surgical units to conform to a similar structure for such meetings.
- The results of the data are very favourable for the quality of surgery being delivered nationally. The outcome of surgically treated isolated aortic valve replacement of over 99% stands out. We are aware that this can alter as the number of cases increase.
- It is important to note that a cardiac surgical team comprises of a number of Medical staff from different specialities both doctors and nurses, allied workers and others who all play a role in the care of the patient and impact on the overall outcome of the procedure. Whilst the operation is ultimately the largest intervention undertaken it is important to stress that each of the medical professionals involved (cardiologist, surgeon, perfusionist, intensive care specialist, anaesthetist, junior doctor, nurse, social worker, physiotherapist, pharmacist and occupational therapist) play an important role and can impact on the morbidity and mortality associated with the procedure for each individual patient.
- Whilst the registry and our regulatory bodies (NZMC, RACS) have processes in place to identify and further assess under-performing 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 cardiosurgical team.
- The New Zealand Cardiac Surgical Registry 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).



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. **Urgent:** not routine; there is a medical reason for operating this admission.
4. **Emergency:** unscheduled surgery required in next available theatre on same day due to refractory angina or cardiac compromise.
5. **Salvage:** the patient is undergoing CPR *en route* to the operating room, that is, prior to surgical incision.
6. **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 countries².
7. **MI:** myocardial infarction.
8. **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.
9. **PTCA:** percutaneous transluminal coronary angioplasty.

2. 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
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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

Appendix

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 No Yes

Previous surgery No Yes

Type of previous surgery

<input type="checkbox"/> CABG	<input type="checkbox"/> Congenital cardiac
<input type="checkbox"/> Off-pump CABG	<input type="checkbox"/> Aortic surgery (ascending / arch)
<input type="checkbox"/> Valve	<input type="checkbox"/> Other cardiac

Number of prior cardiac operations requiring cardiopulmonary bypass

<input type="radio"/> 0	<input type="radio"/> 1	<input type="radio"/> 4	<input type="radio"/> 7
	<input type="radio"/> 2	<input type="radio"/> 5	<input type="radio"/> 8
	<input type="radio"/> 3	<input type="radio"/> 6	<input type="radio"/> 9

Number of prior cardiac operations without cardiopulmonary bypass

<input type="radio"/> 0	<input type="radio"/> 1	<input type="radio"/> 4	<input type="radio"/> 7
	<input type="radio"/> 2	<input type="radio"/> 5	<input type="radio"/> 8
	<input type="radio"/> 3	<input type="radio"/> 6	<input type="radio"/> 9

Previous percutaneous intervention No Yes

PTCA / stent No Yes

PTCA / stent: which admission Prior admission This admission

PTCA / stent: interval (same admission) 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

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

Unique patient identifier

Date of surgery dd/mm/yyyy

Haemodynamic data

Cardiac catheterisation No Yes

Date of cardiac catheterisation dd/mm/yyyy

LVEF method Not measured Echo
 LV gram MRI
 Radionuclide

EF %

EF estimate Normal Moderate
 Mild Severe

Left main stenosis >50% No Yes

Number of diseased coronary systems None Two
 One Three

PA systolic mm Hg

Pulmonary hypertension No Severe
 Moderate



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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 Trainee Overseas fellow Oversight

Operative urgency / status Elective Urgent Emergency Salvage

Direct transfer from cath lab to theatre No Yes

Coronary artery bypass No Yes

Valve surgery No Yes

Valve type Aortic Mitral Tricuspid Pulmonary

Redo valve No Yes

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

Aortic procedure No Yes

Other cardiac procedures No Yes

Other non-cardiac procedures No Yes

Aortic procedure

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

Aortic dissection repair (type) No repair Ascending Descending

Aortic dissection: when Acute Non-acute

Acute traumatic aortic transection No Yes



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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"/> Radial <input type="radio"/> Mini-Maze <input type="radio"/> Left atrial reduction	<input type="radio"/> Pulmonary vein isolation <input type="radio"/> Left atrial only <input type="radio"/> Right atrial only <input type="radio"/> Other
Atrial arrhythmia surgery: energy source	<input type="radio"/> Cut & sew <input type="radio"/> Unipolar RF <input type="radio"/> Bipolar RF <input type="radio"/> Cryoablation	<input type="radio"/> Microwave <input type="radio"/> Laser <input type="radio"/> Ultrasound <input type="radio"/> Other
Type of other cardiac surgery	<input type="checkbox"/> AF ablation surgery <input type="checkbox"/> ASD <input type="checkbox"/> Atrial myxoma <input type="checkbox"/> Cardiac transplant <input type="checkbox"/> Cardiac trauma <input type="checkbox"/> Cardiac trauma - iatrogenic <input type="checkbox"/> Cardiac tumour <input type="checkbox"/> Epicardial pacemaker <input type="checkbox"/> Left ventricular reconstruction <input type="checkbox"/> LV aneurysm <input type="checkbox"/> LVOT myectomy of HOCM	<input type="checkbox"/> LV rupture <input type="checkbox"/> Pericardiectomy <input type="checkbox"/> Peripheral vascular <input type="checkbox"/> Permanent LV epicardial lead <input type="checkbox"/> Primary VAD <input type="checkbox"/> Pulm. thromboendarterectomy <input type="checkbox"/> Pulmonary embolectomy <input type="checkbox"/> Pulmonary transplant <input type="checkbox"/> VSD (acquired) <input type="checkbox"/> Other congenital <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



New Zealand Ministry of Health
NZ Adult Cardiac Surgical Database
Baseline section; Page 11; Version 1.0 (13 Dec 2013)

Unique patient identifier
Date of surgery 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	



New Zealand Ministry of Health
NZ Adult Cardiac Surgical Database
Baseline section; Page 12; Version 1.0 (13 Dec 2013)



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

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

Unique patient identifier
 Date of surgery dd/mm/yyyy

Aortic valve surgery

Aortic valve procedure	<input type="radio"/> Replacement <input type="radio"/> Repair /reconstruction without annuloplasty <input type="radio"/> Root reconstruction with valve conduit (Bentall procedure) <input type="radio"/> Root reconstruction with valve sparing (David procedure) <input type="radio"/> Resuspension aortic valve <input type="radio"/> Resection sub-aortic stenosis <input type="radio"/> Repair paravalvular leak <input type="radio"/> Valvotomy <input type="radio"/> Ross procedure <input type="radio"/> Inspection only <input type="radio"/> Decalcification of valve 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
Aortic stenosis	<input type="radio"/> No <input type="radio"/> Yes	
Aortic regurgitation / insufficiency	<input type="radio"/> None <input type="radio"/> Trivial <input type="radio"/> Mild	
	<input type="radio"/> Moderate <input type="radio"/> Severe	
Aortic 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"/> Annuloaortic ectasia <input type="radio"/> Other degenerative disease <input type="radio"/> Dissection <input type="radio"/> Tumour <input type="radio"/> Trauma <input type="radio"/> Iatrogenic <input type="radio"/> Other	



New Zealand Ministry of Health
NZ Adult Cardiac Surgical Database
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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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Tricuspid valve surgery

Tricuspid valve procedure

- Annuloplasty only
- Replacement
- Repair / reconstruction with annuloplasty
- Repair / reconstruction without annuloplasty
- Commissurotomy with annuloplasty ring
- Commissurotomy without annuloplasty ring
- Repair paravalvular leak
- Valvectomy (no replacement)
- Inspection 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

Tricuspid stenosis

- No
- Yes

Tricuspid regurgitation / insufficiency

- None
- Trivial
- Mild
- Moderate
- Severe

Tricuspid 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
- Other degenerative disease
- Tumour
- Trauma
- Iatrogenic
- Functional
- Other



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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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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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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

