

**Figure 1:** Criteria for subsidy of empagliflozin by special authority. Obtained from the New Zealand Formulary website.<sup>11</sup>

**Initial Application**

*Applications from any relevant practitioner. Approvals valid without further renewal unless notified.*

**Pre-requisites**

Patient has type 2 diabetes

**and**

Patient is Māori or any Pacific ethnicity **or**

Patient has pre-existing cardiovascular or risk equivalent (see note a) **or**

Patient has an absolute 5-year cardiovascular disease risk of 15% or greater according to a validated cardiovascular risk assessment calculator **or**

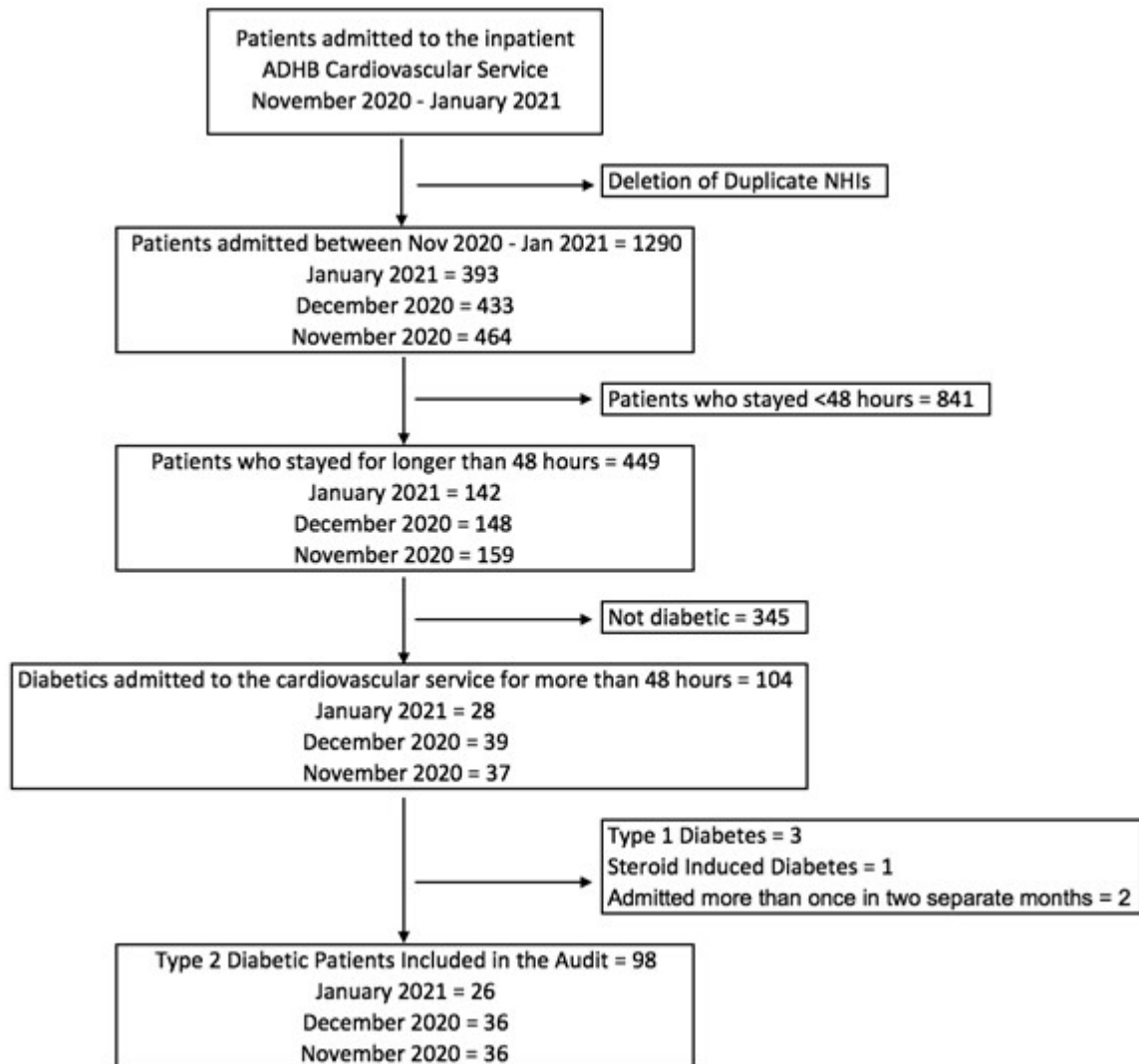
Patient has a high lifetime cardiovascular risk due to being diagnosed with type 2 diabetes during childhood or as a young adult **or**

Patient has diabetic kidney disease (see note b) \*\*

**and**

Target HbA<sub>1c</sub> (of 53mmol/mol or less) has not been achieved despite the regular use of at least one blood-glucose lowering agent (e.g. metformin, vildagliptin or insulin) for at least 3 months

Figure 2: Study flowchart.



**Table 1:** Baseline characteristics.

	Mean $\pm$ SD	Median (IQR)	N=98 (%)
<b>Age [years]</b>	65 $\pm$ 13	64 (56–76)	
<b>Male</b>			65 (66)
<b>Ethnicity</b>			
European			29 (30)
Māori			15 (15)
Asian			24 (24)
Pasifika			29 (30)
Middle Eastern			1 (1.0)
<b>Length of stay [hours]</b>	153 $\pm$ 131	102 (74–181)	
<b>Presentation</b>			
ST-elevation myocardial infarction			12 (12)
Non-ST-elevation myocardial infarction			16 (16)
Unstable angina			3 (3.1)
Heart failure			22 (22)
Arrhythmia			18 (18)
Aortic valve intervention			3 (3.1)
Non-cardiac chest pain			3 (3.1)
Other cardiac			6 (6.1)
Other non-cardiac			6 (6.1)
<b>Body mass index [kg/m<sup>2</sup>]</b>	32 $\pm$ 8.4	30 (25–37)	
<b>Cardiovascular comorbidities</b>			
Hypertension			77 (79)
Heart failure			48 (49)
Atrial arrhythmias			30 (31)
Coronary heart disease			68 (69)
Dyslipidaemia			61 (62)
Stroke			13 (13)
Peripheral vascular disease			5 (5.1)
<b>Smoking status</b>			
Never smoked			45 (46)

**Table 1 continued:** Baseline characteristics.

Ex-smoker			40 (41)
Current smoker			13 (13)
<b>Cardiac medications</b>			
Statins and/or ezetimibe			76 (78)
Alpha-blockers			11 (11)
Calcium channel blockers			25 (26)
ACEi or ARB			80 (82)
Beta-blockers			71 (72)
Diuretics			52 (53)
<b>Left ventricular ejection fraction [%]</b>	45±16	46 (33–58)	
<b>NT-proBNP [pmol/L]</b>	412±593	151 (42–551)	

Abbreviations: ACEi = angiotensin-converting enzyme inhibitor; ARB = angiotensin II receptor blocker; IQR = interquartile range.

**Table 2:** Baseline diabetes characteristics.

	Mean (SD)	Median (IQR)	N=98 (%)
<b>HbA<sub>1c</sub> (mmol/mol)</b>	64±18	59.5 (52–71)	
<60			48 (50)
≥60			48 (50)
<b>Blood pressure</b>			
SBP ≥140mmHg or DBP ≥90mmHg			25 (26)
SBP<140mmHg or DBP <90mmHg			73 (74)
<b>eGFR [mL/min/1.73m<sup>2</sup>]</b>			
≥90			13 (13)
60 to <90			41 (42)
30 to <60			31 (32)
<30			13 (13)
<b>Chronic kidney disease stage</b>			
1			13 (13)
2			41 (42)
3A			20 (20)
3B			11 (11)
4			7 (7.1)
5			6 (6.1)
<b>Urine albumin:creatinine ratio (mg/g)</b>			
<3			34 (35)
≥3 to <30			43 (44)
≥30 to 300			11 (11)
>300			6 (6.1)
Nil			4 (4.1)
<b>Glycaemic medications on admission</b>			
Metformin			49 (50)
Vildagliptin			5 (5.0)
Vildagliptin/metformin combination			16 (16)
Sulphonylurea			18 (18)
Insulin			34 (35)
None			16 (16)

Abbreviations: DBP = diastolic blood pressure; eGFR = estimated glomerular filtration rate; IQR = interquartile range; NTproBNP = N-terminal pro-brain natriuretic peptide; SBP = systolic blood pressure.

**Table 3:** Changes to glycaemic medications during admission.

<b>All patients (n=98)</b>	
Changed	36 (37%)
No change	51 (52%)
No treatment at discharge	11 (11%)
<b>HbA<sub>1c</sub> &gt;60 mmol/mol (n=48)</b>	
Changed	24 (50%)
No change	21 (44%)
No treatment at discharge	3 (6.0%)
<b>HbA<sub>1c</sub> ≤60 mmol/mol (n=48)</b>	
Changed	12 (25%)
No change	8 (17%)
No treatment at discharge	28 (58%)

NB: Two patients had type 2 diabetes as part of their medical history, however, their primary residence was not Auckland, so no HbA<sub>1c</sub> was recorded on their electronic medical records, thus they were not included in the sub-group analysis of HbA<sub>1c</sub> control.

**Table 4:** Eligibility for subsidisation of SGLT-2 trials.

<b>Eligible for subsidisation of empagliflozin under special authority</b>	
Yes	49 (50%)
No due to <3 months glycaemic therapy prior to admission	11 (11%)
No due to HbA <sub>1c</sub> ≤53mmol/mol	23 (23%)
Insufficient information	2 (2.0%)
Excluded as eGFR<30 mL/min/1.73 m <sup>2</sup>	13 (13%)
<b>Eligible for inclusion in EMPA-REG OUTCOME and/or EMPEROR-Reduced</b>	
Yes	34 (35%)
No	64 (65%)

Abbreviations: eGFR = estimated glomerular filtration rate; EMPA-REG OUTCOME = Empagliflozin, Cardiovascular Outcomes, and Mortality in Type 2 Diabetes; EMPEROR-Reduced = Cardiovascular and Renal Outcomes with Empagliflozin in Heart Failure.