

Appendix 1, Hypothetical Power Analysis

| RRT in on-pump group | <i>Proportion Diff</i> | <i>Power</i> | <i>N Per Group</i> |
|-------------------------|------------------------|--------------|--------------------|
| 0.12 | -0.025 | 0.8 | 2409 |
| 0.12 | -0.025 | 0.9 | 3224 |
| 0.12 | -0.030 | 0.8 | 1638 |
| 0.12 | -0.030 | 0.9 | 2193 |
| 0.12 | -0.035 | 0.8 | 1178 |
| 0.12 | -0.035 | 0.9 | 1577 |
| 0.12 | -0.040 | 0.8 | 882 |
| 0.12 | -0.040 | 0.9 | 1181 |
| 0.12 | -0.045 | 0.8 | 681 |
| 0.12 | -0.045 | 0.9 | 912 |

| RRT in on-pump group | <i>Proportion Diff</i> | <i>Nominal Power</i> | <i>N Per Group</i> |
|-------------------------|------------------------|----------------------|--------------------|
| 0.13 | -0.025 | 0.8 | 2604 |
| 0.13 | -0.025 | 0.9 | 3485 |
| 0.13 | -0.030 | 0.8 | 1774 |
| 0.13 | -0.030 | 0.9 | 2375 |
| 0.13 | -0.035 | 0.8 | 1279 |
| 0.13 | -0.035 | 0.9 | 1711 |
| 0.13 | -0.040 | 0.8 | 960 |
| 0.13 | -0.040 | 0.9 | 1284 |
| 0.13 | -0.045 | 0.8 | 743 |
| 0.13 | -0.045 | 0.9 | 994 |

| RRT in on-pump group | <i>Proportion Diff</i> | <i>Nominal Power</i> | <i>N Per Group</i> |
|-------------------------|------------------------|----------------------|--------------------|
| 0.14 | -0.025 | 0.8 | 2793 |
| 0.14 | -0.025 | 0.9 | 3739 |
| 0.14 | -0.030 | 0.8 | 1907 |
| 0.14 | -0.030 | 0.9 | 2552 |
| 0.14 | -0.035 | 0.8 | 1377 |
| 0.14 | -0.035 | 0.9 | 1842 |
| 0.14 | -0.040 | 0.8 | 1035 |
| 0.14 | -0.040 | 0.9 | 1385 |
| 0.14 | -0.045 | 0.8 | 803 |
| 0.14 | -0.045 | 0.9 | 1075 |

The sample size was calculated for hypothetical randomized clinical trial with RRT as the primary endpoint and equal allocation of patients within on and off pump group. Sample sizes are summarized according to the combination of following scenarios based on the data in our cohort:

Proportion of RRT in on-pump group: 0.12, 0.13, 0.14

Risk reduction in off-pump group: -0.025 -0.03 -0.035 -0.04 -0.045

Power: 0.8, 0.9.

We provide the Table above as a reference. The number of patients required for an appropriately clinical trial could be substantially reduced by utilizing study entry criteria that enrich the test population for those patients at highest risk, thereby increasing the event rate and decreasing the requisite number of patients.

Appendix 2, Propensity Score Distribution

| | | Propensity score distribution | | | | |
|------------|--------------------|-------------------------------|--------------------|--------------------|--------------------|-------|
| GFR strata | Trt group | min | 20 th % | 50 th % | 80 th % | max |
| 15-29 | On-Pump | 0.0579 | 0.177 | 0.240 | 0.325 | 0.876 |
| | Intent-to-off pump | 0.0917 | 0.209 | 0.292 | 0.408 | 0.814 |
| 30-59 | On-Pump | 0.0477 | 0.154 | 0.195 | 0.257 | 0.796 |
| | Intent-to-off pump | 0.0797 | 0.172 | 0.227 | 0.315 | 0.783 |
| 60-89 | On-Pump | 0.068 | 0.147 | 0.18165 | 0.241 | 0.756 |
| | Intent-to-off pump | 0.078 | 0.163 | 0.21245 | 0.303 | 0.799 |
| ≥90 | On-Pump | 0.0713 | 0.139 | 0.175 | 0.234 | 0.743 |
| | Intent-to-off pump | 0.0717 | 0.156 | 0.207 | 0.302 | 0.816 |

Appendix 3

The propensity models include each of the following variables: age, gender, race (white, black, Asian, and other), body surface area (BSA), left ventricular ejection fraction (EF), past or present smoker, hypertension, hypercholesterolemia, cerebrovascular disease, cerebrovascular accident, peripheral vascular disease, immunosuppressive treatment, chronic lung disease (severe, moderate, mild, none), diabetes (insulin, non-insulin, none), renal function (eGFR90-, eGFR60-89, eGFR 30-59, eGFR15-29), arrhythmia (atrial fibrillation, Sust VT/VF, Heart block), endocarditis (active, treated, none), angina (stable, unstable), history of myocardial infarction(<21 days, \geq 21 says, none), percutaneous coronary intervention \leq 6 hr, pre-operative intra-aortic balloon pump or administration of inotropes, congestive heart failure (NYHA class IV, NYHA class I-III, no heart failure), left main disease, number diseased coronary vessels (0,1,2,3), aortic stenosis , mitral Stenosis , aortic insufficiency(moderate to severe), mitral insufficiency(moderate to severe), tricuspid insufficiency(moderate to severe), preoperative medications (Beta Blockers , Any Anticoagulants, ACE or ARB Inhibitors, steroid, lipid lowering) previous CABG, previous valve, previous PCI, number of previous cardiovascular interventions (0, 1, 2 or more), acuity status (urgent, elective), concomitant CABG, and year of surgery

Appendix 4. Characteristics of Patients at on- pump versus off-pump centers

| Characteristics | Baseline | | |
|------------------------------------|-----------------------------------|-----------------------------------|-------------------------------|
| | On-Pump Centers (n=265,497) | Off-Pump Centers (n=26,904) | Standardized Difference, % |
| Age, median (IQR) | 65 (58, 73) | 66 (58, 74) | 6.2 |
| Female, % | 26.8 | 27.2 | 7.1 |
| BSA, median (IQR) | 2.0 (1.8, 2.2) | 2.0 (1.8, 2.2) | 8.8 |
| Hypertension | 82.3 | 81.7 | 0.2 |
| Diabetes | 38.0 | 37.4 | 4.9 |
| CLD | 21.9 | 20.7 | |
| eGFR | | | |
| ≥90 | 24.1 | 22.5 | 2.6 |
| 60-89 | 51.6 | 52.1 | 1.1 |
| 30-59 | 22.9 | 23.8 | 2.7 |
| 15-29 | 1.4 | 1.6 | 4.0 |
| Prior CV Surgery | 4.7 | 4.6 | 3.9 |
| Prior MI | 43.5 | 42.5 | 0.6 |
| CHF | 12.2 | 13.2 | 0.7 |
| Ejection Fraction, median (IQR) | 55 (45, 60) | 55 (45, 60) | 6.8 |
| Left Main >50% | 30.5 | 29.7 | 5.4 |
| CAD, # vessels | | | |
| 1 | 3.8 | 5.5 | 33.6 |
| 2 | 18.8 | 21.7 | 17.1 |
| 3 | 77.5 | 72.8 | 34.6 |
| Procedure Status | | | |
| Elective | 48.8 | 51.4 | 7.2 |
| Urgent | 51.2 | 48.5 | 7.2 |

Appendix 5. Unadjusted and adjusted risk difference between on-pump and off-pump centers for AVR+CABG procedures

| Event | Unadjusted Risk Difference | | | | Adjusted risk difference | | | |
|---------------------|----------------------------|-------------------------|-----------------------------|---------|--------------------------|-------------------------|-----------------------------|---------|
| | On- Pump Centers | Off- Pump Centers | Risk Difference (95 CI)* | P-value | On- Pump Centers | Off- Pump Centers | Risk Difference (95 CI)* | P-value |
| | % | % | | | % | % | | |
| Death or RRT | 5.1 | 5.4 | -0.30 (-1.22, 0.62) | 0.51 | 5.1 | 8.0 | -2.87 (-6.66, 0.93) | 0.14 |
| Death | 3.8 | 4.0 | -0.18 (-0.98, 0.61) | 0.65 | 3.8 | 5.9 | -2.10 (-5.51, 1.31) | 0.23 |
| RRT | 2.4 | 2.3 | 0.10 (-0.52, 0.71) | 0.76 | 2.4 | 2.8 | -0.39 (-2.29, 1.50) | 0.69 |

*:Number of patients with the outcome per 100 patients treated at centers with a preference for on-pump CABG minus the number of patients with the outcome per 100 patients treated at centers with a preference for off-pump CABG

Table 5, Summary of Previous Studies

| Study | Total | Baseline CKD |
|---|-------|--------------|
| | n | n (%) |
| Ascione et. al (1999) ¹ | 50 | 0(0) |
| Nathoe et. al. (2003) ² | 281 | 0(0) |
| Tang et al (2002) ³ | 40 | 0(0) |
| Legare et al (2003) ⁴ | 300 | 11(3.7) |
| Puskas et. al (2003) ⁵ | 197 | 4(2) |
| Straka et. al (2004) ⁶ | 388 | 3(0.8) |
| Khan et al (2004) ⁷ | 104 | 0(0) |
| Gerola et al. (2004) ⁸ | 160 | 0(0) |
| Wan et al (2004) ⁹ | 37 | 0(0) |
| Staton et. al. (2005) ¹⁰ | 197 | 0(0) |
| Jensen et. al. (2006) ¹¹ | 120 | 0(0) |
| Motallebzadeh et. al.(2006) ¹² | 210 | 0(0) |
| Sajja et. al. (2007) ¹³ | 116 | 116(100) |
| Magee et. al. (2008) ¹⁴ | 3014 | 67(2.2) |
| Paulitsch et. al. (2009) ¹⁵ | 92 | 0(0) |
| Shroyer et al (2009) ¹⁶ | 2203 | 173(7.9) |
| Hueb et. al. (2010) ¹⁷ | 308 | 0(0) |
| Moller et. al. (2010) ¹⁸ | 339 | 13(3.8) |
| | | |
| | | |
| Total | 8156 | 387(4.7) |
| | | |

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