OUTCOMES FOLLOWING TRANSCATHETER AORTIC VALVE IMPLANTATION COMPARING EDWARDS SAPIEN WITH MEDTRONIC COREVALVE REVALVING SYSTEM DEVICES: RESULTS FROM THE MILAN REGISTRY

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Background To assess clinical outcomes of transcatheter aortic valve implantation comparing Medtronic CoreValve ReValving System® with Edwards SAPIEN XT.

Methods All consecutive patients in our center with aortic stenosis treated with transfemoral Medtronic CoreValve ReValving System® (MCV) from November 2009 to September 2011 (learning curve patients excluded) or Edwards SAPIEN XT (ESV) from April 2010 to September 2011 when the device became available were included.

Result In total, there were 192 patients in this analysis. The overall mean age was 79.4±6.1 years, logistic EuroSCORE 21.1±15.9% and STS-PROM score 8.8±2.6%. The MCV group consisted of a greater proportion of males (60.3% vs 43.7%; p=0.026) with a corresponding larger aortic annulus size (24.7±2.0 vs 25.4±1.7; p<0.001). The median clinical follow-up length was 171 (IQR 54–357) days. 30-day all cause mortality was 4.0%, myocardial infarction rate 1.0%.

Conclusions In our single center experience, transcatheter aortic valve implantation was a relatively safe and effective procedure utilising both commercially available devices. However, there was an increased incidence of arrhythmia and pacemaker implantation in the MCV group.

SERIAL CHANGE IN HEALTH RELATED QUALITY OF LIFE OVER 1 YEAR FOLLOWING TRANSCATHETER AORTIC VALVE IMPLANTATION: PREDICTORS OF HEALTH OUTCOMES

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Introduction Severe aortic stenosis (AS) reduces the length and quality of a patient’s life. Transcatheter Aortic Valve Implantation (TAVI) is superior to standard medical therapy and non-inferior to Surgical Aortic Valve Replacement (SAVR) for 1-year mortality. HRQOL is an important outcome measure, for which there is limited evidence in TAVI populations. This study aimed to assess serial changes in patient health related quality of life (HRQOL) over time and identify predictors of patient health.

Method 102 (age 80±0.6 years, female 51%) attending for TAVI consented to participate. Two HRQOL questionnaires, the SF12v2 with physical and mental component summaries (PCS and MCS) and the EQ-5D (with Visual Analogue Scale, VAS) were completed at baseline, 30 days, 6 and 12 months. A SF-6D utility measure was calculated from the SF12 survey.

Results HRQOL significantly improved over 12 months (PCS, p<0.002; EQ-5D, p=0.02; VAS, p=0.01; SF-6D p=0.048). The greatest change occurred from baseline to 30 days (p<0.001) with further significant increases to 6 months (p<0.01). An insignificant decline occurred between 6 and 12 months (p>0.05), Abstract 041 table 1. Female gender (p<0.05), age ≥80 years (p<0.05), previous MI (p<0.04), CKD (p<0.05), pulmonary hypertension (p=0.04) and lower operator experience (p<0.001) were independent predictors of a smaller improvement in HRQOL, Abstract 041 figure 1.