Delirium after surgical and transcatheter aortic valve replacement is associated with increased mortality

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ABSTRACT

Objective: The purpose of this study was to determine the incidence and clinical significance of postoperative delirium (PD) in patients with aortic stenosis undergoing surgical aortic valve replacement (SAVR) or transcatheter aortic valve replacement (TAVR).

Method: Between 2010 and 2013, 427 patients underwent TAVR (n = 168) or SAVR (n = 259) and were screened for PD using the Confusion Assessment Method for the Intensive Care Unit. The incidence of PD in both treatment groups was determined and its association with morbidity and mortality was retrospectively compared.

Results: PD occurred in 135 patients (32%) with a similar incidence between SAVR (33% [86 out of 259]) and TAVR (29% [49 out of 168]) ($P = .40$). TAVR by transfemoral approach had the lowest incidence of PD compared with SAVR (18% vs 33%; $P = .025$) or TAVR when performed by alternative access techniques (18% vs 35%; $P = .02$). Delirium was associated with longer initial intensive care unit stay (70 vs 27 hours), intensive care unit readmission (10% [14 out of 135] vs 2% [6 out of 292]), and longer hospital stay (8 vs 6 days) ($P < .001$ for all). PD was associated with increased mortality at 30 days (7% vs 1%; $P < .001$) and 1 year (21% vs 8%; $P < .001$). After multivariable adjustment, PD remained associated with increased 1-year mortality (hazard ratio, 3.02; 95% confidence interval, 1.75-5.23; $P < .001$). There was no interaction between PD and aortic valve replacement approach with respect to 1-year mortality ($P = .12$). Among propensity-matched patients (n = 170), SAVR-treated patients had a higher incidence of PD than TAVR-treated patients (51% vs 29%; $P = .004$).

Conclusions: PD occurs commonly after SAVR and TAVR and is associated with increased morbidity and mortality. Given the high incidence of PD and its associated adverse outcomes, further studies are needed to minimize PD and potentially improve patient outcomes. (J Thorac Cardiovasc Surg 2016;151:815-23)

Surgical aortic valve replacement and transcatheter aortic valve replacement with and without delirium.

Central Message

Delirium after surgical and transcatheter aortic valve replacement is associated with increased mortality.

Perspective

Postoperative delirium was assessed in patients undergoing surgical and transcatheter aortic valve replacement. The development of postoperative delirium was associated with increased perioperative morbidity as well as both early and 1-year mortality.

See Editorial Commentary page 823.

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Postoperative delirium (PD) is a well-recognized complication after cardiac surgery defined as an acute and fluctuating neurologic disorder that reflects a change from baseline cognition and is characterized by the cardinal features of inattention and disorganized thinking.¹ PD is more common

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Abbreviations and Acronyms

AKI	= acute kidney injury
AS	= aortic stenosis
AVR	= aortic valve replacement
CAM	= confusion assessment method
ICU	= intensive care unit
NYHA	= New York Heart Association
PRBC	= packed red blood cell
PD	= postoperative delirium
PROM	= predicted risk of mortality
RASS	= Richmond Agitation and Sedation Scale
SAVR	= surgical aortic valve replacement
STS	= Society of Thoracic Surgeons
TAVR	= transcatheter aortic valve replacement
TA	= transapical
Tao	= transaortic
TF	= transfemoral

in patients older than age 65 years and has an estimated incidence between 25% and 50% after cardiac surgery, which translates into more than 120,000 patients affected per year in the United States alone.²⁻⁴ Although efforts to detect PD have increased during the past decade, it remains underestimated in many clinical settings. Moreover, as older and frailer patients increasingly undergo major procedures, the incidence of PD is likely to increase substantially during the next decade.⁵

PD after traditional cardiac surgery has been associated with increased morbidity, longer hospital length of stay, and greater mortality.⁶⁻⁸ These negative outcomes have resulted in increased health care costs estimated at \$150 billion annually.⁹ Although the influence of PD on short-term and perioperative outcomes seems intuitive, PD has additionally been linked to late cognitive impairments that persist at intermediate and long-term follow-up.^{4,10} Multiple studies have focused on treating PD, but most have had limited success, resulting in a greater emphasis placed upon the prevention of PD and the identification of risk factors for its development.⁵

Recent advances in the treatment of patients with severe aortic stenosis (AS), specifically in the area of transcatheter aortic valve replacement (TAVR) now allow for aortic valve replacement (AVR) procedures to be performed without the need for cardiopulmonary bypass, requiring fewer transfusions, and in many cases, without even a surgical incision or general anesthesia.^{11,12} As such, TAVR procedures minimize or eliminate many of the traditional factors associated with the development of PD after conventional AVR surgery. Although several large trials have evaluated outcomes after TAVR in comparison to surgical aortic valve replacement (SAVR), there are fewer data available

regarding the use of TAVR and its association with PD and subsequent sequelae.

The purpose of this study was to compare the incidence of PD after TAVR and SAVR, examine the adverse outcomes associated with PD, and explore possible risk factors for the development of PD after either procedure. We hypothesized that the incidence of PD would be lower after TAVR than SAVR, would be associated with increased mortality after either procedure, and would be associated with different risk factors for its development within each treatment population.

METHODS

In this retrospective study from 2010 to 2013, we included all patients with severe AS who underwent an isolated AVR procedure at Barnes-Jewish Hospital in whom PD was assessed with the Confusion Assessment Method for the Intensive Care Unit (CAM-ICU). Severe AS was defined as an indexed aortic valve area $<0.6 \text{ cm}^2/\text{m}^2$, a mean gradient $>40 \text{ mm Hg}$, or a maximum jet velocity $>4.0 \text{ m/sec}$. Patients were offered TAVR in the setting of a clinical trial or if they met TAVR eligibility criteria once TAVR became approved for use by the US Food and Drug Administration. A total of 427 patients were treated with either TAVR ($n = 168$) or SAVR ($n = 259$).

This study was written in compliance with Strengthening the Reporting of Observational studies in Epidemiology guidelines and approved by the local institutional review board from which a waiver of written informed consent was obtained given the retrospective design of the study.

TAVR and SAVR Approaches

The TAVR procedures have previously been described¹¹ and were performed via a transfemoral (TF) approach when femoral access was suitable ($n = 57$). In cases where TAVR-TF was not possible, TAVR was performed either by a direct aortic (TAo) ($n = 37$) or transapical (TA) ($n = 74$) approach. All procedures were performed under general anesthesia and all TF approaches were performed using surgical femoral exposure. SAVR procedures were performed via limited or full sternotomy using cardiopulmonary bypass and myocardial arrest.

Anesthetic Approach for TAVR and SAVR

General anesthesia was typically induced with propofol, fentanyl, and rocuronium. A limited use of benzodiazepines was not uncommon. All patients had endotracheal tubes placed and were mechanically ventilated for the procedure. Anesthesia was maintained with a balanced anesthetic technique, including a potent inhaled agent and titration of opioid analgesics. Transesophageal echocardiography was performed in all cases. Intraoperative data, including vital signs and medications, were automatically archived every minute by an electronic anesthetic record system (Metavision; IMDsoft, Needham, Mass). Volatile anesthetic concentrations were converted into age-adjusted minimum alveolar concentration equivalents,¹³ summed across all volatile anesthetic agents.

Delirium Assessment

After the AVR procedure, all patients were admitted to an intensive care unit (ICU), where clinical nurses trained in the administration of the CAM-ICU assessed patients for PD every 12 hours using the CAM-ICU. The CAM-ICU assessment has been a routine part of the standard postoperative nursing assessment of all cardiac surgical patients since its introduction in 2010. Although the CAM-ICU was developed for nonverbal ICU patients, it has also been shown to be specific and reasonably sensitive in nonintubated ICU patients.^{14,15} The tool is based on operationalized

criteria derived from the Diagnostic and Statistical Manual of Mental Disorders and assesses 4 features: acute-onset and fluctuating course, inattention, disorganized thinking, and altered level of consciousness. The CAM-ICU assessment was positive for delirium when both acute-onset and fluctuating course and inattention, and either disorganized thinking or altered level of consciousness were present (http://www.icudelirium.org/docs/CAM_ICU_training.pdf). Patients were identified as having PD if they had at least 1 positive CAM-ICU assessment during their ICU stay.

Delirium episodes were further classified as hypoactive, hyperactive, or mixed by using the Richmond Agitation and Sedation Scale (RASS) documented at the time of the delirium assessment.^{16,17} The RASS is a 10-point sedation scale that measures level of consciousness. A RASS of 0 reflects an alert and calm patient, whereas scores >0 indicate increasing agitation, and scores <0 indicate deepening sedation. RASS scores were documented every 4 hours in the ICU and the corresponding RASS score within 12 hours of a given CAM-ICU assessment (the 6 hours before and after the assessment) was used to assess delirium phenotype. When patients were CAM-ICU positive, a RASS consistently >0 was indicative of hyperactive delirium, a RASS consistently <0 was indicative of hypoactive delirium, and for patients exhibiting scores both >0 and <0 , the delirium phenotype was considered mixed.

All perioperative data, including Society of Thoracic Surgeons database variables, intraoperative monitoring data, delirium, and RASS assessments were prospectively collected within institutional databases and subsequently retrospectively reviewed for perioperative outcomes.

Statistical Analyses

Descriptive statistics were determined for TAVR and SAVR patients separately. Comparisons between groups were done using Student *t* test for continuous variables and Fisher exact test for categorical variables. Nonnormal and ordinal variables were summarized as median (first quartile, third quartile) and compared via the Kruskal-Wallis test.

Propensity score matching was used to compare PD rates between TAVR and SAVR patients. A logistic regression model was used to create the probability that patient was assigned to TAVR for treatment using the following variables: age, gender, Society of Thoracic Surgeons score, New York Heart Association (NYHA) functional class III or IV (vs I or II), diabetes mellitus, coronary artery disease, peripheral vascular disease, cerebrovascular disease, moderate/severe lung disease, glomerular filtration rate, and hemoglobin. An optimal matching algorithm with caliper set to 0.05 was used to create a 1:1 matched sample. Delirium rates were compared within the matched sample using McNemar test (Figures E1 and E2).

Mortality was evaluated using time to event analysis. Start time was date of the AVR procedure and patients were followed until event occurrence or last available follow-up, to 1 year. Kaplan-Meier curves were created by delirium status and compared using the log-rank test. Additional multivariable Cox proportional hazards regression models were created to evaluate the association between delirium and 1-year mortality while adjusting for AVR approach, age, sex, and STS predicted risk of mortality (STS-PROM) score; the interaction between PD and AVR approach was assessed. These covariates were selected a priori based on their known association with mortality. Within AVR approach, the rates of occurrence for prolonged ventilation, stroke, acute kidney injury (AKI), and ICU readmission were individually compared between delirium groups using Fisher exact test. AKI was defined in accordance with Valve Academic Research Consortium-2 criteria.¹⁸ Thirty-day and 1-year Kaplan-Meier estimates for survival were compared using a *z* test. The Kruskal-Wallis test was used to compare initial ICU and hospital length of stay.

Multivariable logistic regression models were built to explore predictors of delirium within TAVR and SAVR patients separately. Variables for inclusion within the multivariable models were selected based on a univariable cutoff *P* value of $<.1$. The number of variables included in

the models was limited based on the incidence of PD in each cohort. The variables age, STS-PROM, steroid use, TAVR approach, and intraoperative packed red blood cell (PRBC) transfusions, were entered into the TAVR model. For SAVR patients, the variables included age, STS-PROM, previous percutaneous coronary intervention, NYHA functional class $\geq$ III, moderate/severe lung disease, hemoglobin, moderate/severe mitral regurgitation, intraoperative PRBC use, intra-aortic balloon pump, and intubation time. These models evaluated the association between each variable and PD while adjusting for other model variables.

All analyses were conducted using SAS version 9.4 (SAS Institute Inc, Cary, NC).

RESULTS

Demographic and Intraoperative Differences Between TAVR and SAVR

When compared with SAVR, patients undergoing TAVR were significantly older, had greater frequencies of comorbidities of nearly all measured variables, were of worse functional class by American Society of Anesthesiologists score, and had higher predicted risks of mortality by the STS-PROM (Table 1).

Intraoperatively, whereas TAVR procedures were shorter in duration (153 ± 58 minutes vs 253 ± 73 minutes; $P = .001$) these patients received a higher mean minimum alveolar concentration of volatile anesthetic than patients undergoing SAVR (0.95 ± 0.13 vs 0.86 ± 0.11 ; $P < .001$). PRBC transfusions (67% vs 40%; $P < .001$) and tranexamic acid administration (93% vs 2%; $P < .001$) were more frequent during SAVR procedures.

Delirium Incidence in TAVR and SAVR

For the entire study, PD was identified in 135 patients (32%; 95% confidence interval [CI], 27%-36%). Hypoactive delirium was the most common subtype, encountered in 71 patients (53%); 64 patients (47%) presented with a mixed phenotype; and no patient consistently had hyperactive delirium. The incidence of PD after TAVR was not significantly different than after SAVR (29% vs 33%; odds ratio [OR], 0.82; 95% CI, 0.54-1.26; $P = .40$); however within the subset of TAVR, patients undergoing TAVR-TF had an 18% incidence of PD, the lowest incidence of any AVR approach. The incidence of PD after TAVR-TF was significantly lower when compared with SAVR (18% vs 33%; OR, 0.43; 95% CI, 0.21-0.89; $P = .025$) or TAVR when performed using alternative access (ie, TA or TAo) techniques (18% vs 35%; OR, 0.39; 95% CI, 0.18-0.86; $P = .020$). After propensity matching, 29% (25 out of 85) of the TAVR-treated patients and 51% (43 out of 85) of the SAVR-treated patients developed PD ($P = .004$).

Perioperative Complications and Mortality

Table 2 shows univariable associations between delirium and postoperative complications. For both TAVR and SAVR patients, PD was associated with a longer ICU and hospital length of stay. For SAVR specifically, there were additional

TABLE 1. Baseline characteristics of patients undergoing transcatheter aortic valve replacement (TAVR) and surgical aortic valve replacement (SAVR)

Characteristic	TAVR (n = 168)	SAVR (n = 259)	P value
Age (y)	81 ± 8	71 ± 11	<.001
Female	92 (55)	108 (42)	.010
Body mass index	27.5 ± 6.3	31 ± 8	<.001
Society of Thoracic Surgeons predicted risk of mortality score	9.3 (6.1, 13.5)	3.0 (1.5, 6.0)	<.001
American Society of Anesthesiologists class ≥ 4	162 (96)	180 (70)	<.001
New York Heart Association functional class III-IV	147 (88)	139 (54)	<.001
Diabetes mellitus	72 (43)	95 (37)	.220
Hypertension	156 (93)	209 (81)	<.001
Coronary artery disease	143 (85)	114 (44)	<.001
Cerebrovascular disease	52 (31)	57 (22)	.040
Atrial fibrillation	65 (39)	49 (19)	<.001
Chronic lung disease*	59 (35)	39 (15)	<.001
Peripheral vascular disease	113 (67)	71 (27)	<.001
Ejection fraction (%)	53 ± 15	58 ± 14	.002
Mean aortic valve gradient (mm Hg)	44 ± 14	43 ± 14	.700
Preoperative glomerular filtration rate (mL/min)	64 ± 27	75 ± 26	<.001
Hemoglobin (g/dL)	11.3 ± 1.5	12.5 ± 1.9	<.001
Albumin (g/dL)	4.0 ± 0.5	4.1 ± 0.4	.120

Values are presented as mean ± standard deviation, n (%), or median (first quartile, third quartile). TAVR, transcatheter aortic valve replacement; SAVR, surgical aortic valve replacement. *Moderate/severe.

complications associated with PD that included prolonged ventilation, and a greater need for readmission to the ICU. PD after TAVR and SAVR was associated with an increased likelihood that at hospital discharge, patients required transfer to an extended care facility, and were less likely to be discharged to home. These associations between PD and AKI, discharge location, ICU time and hospital length of stay all remained unchanged even after adjustment by STS-PROM score.

In unadjusted analyses, the development of PD after SAVR or TAVR was associated with increased mortality over the first year for either procedure (Figure 1). For TAVR patients without PD, survival was better at 30 days (98% vs 88%; $P = .029$) and tended to be better at 1 year (80% vs 66%; $P = .09$) based on Kaplan-Meier estimates. Similarly for SAVR, Kaplan-Meier estimates of survival were better for patients without PD at both 30 days (100% vs 95%; $P = .04$) and 1 year (97% vs 83%; $P = .001$), respectively. By multivariable analysis, after adjusting for age, sex, STS-PROM score, and AVR type, PD was associated with increased risk of mortality through the first year after AVR for the entire study population ($n = 427$) (hazard ratio [HR], 3.02; 95% CI, 1.75-5.23; $P < .001$) (Table 3). PD additionally remained significantly associated with increased risk of mortality within both TAVR (HR, 2.02; 95% CI, 1.12-4.35) and SAVR groups (HR, 6.28; 95% CI, 2.04-19.31) after similarly adjusting for age, sex, and STS-PROM score. There was also no significant interaction between AVR type (TAVR vs SAVR) and PD with respect to 1-year mortality (interaction $P = .12$), indicating that the relationship between PD and post-AVR mortality was similar for patients treated with TAVR and SAVR.

Predictors of Delirium After TAVR and SAVR

Univariable associations with PD for patients undergoing both TAVR and SAVR are shown in Table 4. The use of a non-TF approach ($P = .013$) and preoperative steroid use (8% vs 24%; $P = .004$) were associated with the development of PD after TAVR, whereas for SAVR, the development of PD was associated with increasing age, elevated STS-PROM score, worse American Society of Anesthesiologists score, NYHA functional class III or IV, and the presence of specific comorbidities such as coronary artery disease and chronic lung disease. PD was seen more frequently in patients with lower preoperative hemoglobin levels and, similar to TAVR, PD was more frequent when PRBC transfusions were required intraoperatively.

TABLE 2. Perioperative morbidity and mortality for patients undergoing transcatheter aortic valve replacement (TAVR) and surgical aortic valve replacement (SAVR)

Event	TAVR			SAVR		
	No delirium (n = 119)	Delirium (n = 49)	P value	No delirium (n = 173)	Delirium (n = 86)	P value
Prolonged ventilation	2 (2)	0 (0)	>.999	10 (6)	18 (21)	<.001
Stroke	1 (1)	1 (2)	.500	1 (1)	1 (1)	>.999
Acute kidney injury	8 (7)	12 (24)	.003	25 (14)	15 (17)	.580
Initial intensive care unit stay (h)	26	50	<.001	31	72	<.001
Readmission to the intensive care unit	4 (3)	5 (10)	.120	2 (1)	9 (10)	.001
Hospital length of stay (d)	5	7	<.001	7	9	<.001
Discharged to home	87 (73)	26 (53)	.018	146 (84)	49 (57)	<.001
30-d survival (%)	98	88	.029	100	95	.040
1-y survival (%)	80	66	.090	97	83	.001

Values are presented as n (%) unless otherwise noted. TAVR, transcatheter aortic valve replacement; SAVR, surgical aortic valve replacement.

FIGURE 1. Time to event curves for 1-year death from any cause. One year time to event curves are shown for patients treated with surgical aortic valve replacement (SAVR) and transcatheter aortic valve replacement (TAVR) with and without delirium. Within each treatment group, patients with and without delirium were compared with the use of the log-rank test. AVR, Aortic valve replacement.

Specifically for SAVR, there were no differences in PD with respect to cardiopulmonary bypass (110 ± 37 vs 108 ± 25 ; $P = .63$) or aortic crossclamp times (76 ± 23 vs 72 ± 18 ; $P = .15$). Procedural time was not a factor for PD for either TAVR or SAVR.

The independent predictors for the development of delirium are shown in [Tables 5](#) and [6](#). Within the SAVR cohort, the independent predictors for the development of PD were age, and the presence of concomitant valvular heart disease, specifically mitral regurgitation. An intubation period <24 hours after SAVR was associated with less PD ([Table 5](#)). For TAVR, no specific comorbidities were identified as independent predictors for PD with only preoperative steroid use and the development of AKI postoperatively remaining statistically significant. The intraoperative associations for TAVR between transfusions (OR, 2.00; $P = .08$) and TAVR approach (OR, 2.00; $P = .11$) did not remain statistically significant within the multivariable model ([Table 6](#)).

TABLE 3. Multivariable analysis for 1-year mortality for patients undergoing transcatheter aortic valve replacement (TAVR) and surgical aortic valve replacement (SAVR)

Variable	Hazard ratio	95% Confidence interval	P value
Delirium (yes)	3.023	1.748-5.228	<.001
TAVR (yes)	2.360	1.269-4.389	.007
Age (per 1 year)	1.006	0.976-1.036	.710
Female (yes)	0.580	0.329-1.023	.060
Society of Thoracic Surgeons predicted risk of mortality score (per 1 unit)	1.099	1.054-1.146	<.001

TAVR, Transcatheter aortic valve replacement.

DISCUSSION

We found that PD is common after TAVR and SAVR, occurring in nearly one-third of patients. Importantly, patients undergoing TAVR-TF had the lowest incidence of PD when compared with SAVR or TAVR performed by any other access route. PD was associated with increased morbidity and resource use after both TAVR and SAVR and a 3-fold increase in mortality during the first year after valve replacement. The predictors of PD after TAVR or SAVR were different and based primarily upon preexisting comorbidities rather than intraoperative characteristics.

This is the largest study of PD performed specifically among AS patients undergoing valve replacement. It also examines a cohort of patients at higher risk for PD in that it includes a substantial number of patients undergoing TAVR who are older and have more comorbidities than patients typically offered traditional cardiac surgery. Our findings from the patients treated with SAVR are consistent with the literature available regarding PD after cardiac surgery with respect to incidence and inciting risk factors. And although the list of possible risk factors for PD after SAVR remains long, the variables identified in this study such as age, a history of coronary artery disease, a need for an intra-aortic balloon pump preoperatively, and greater PRBC transfusions have all previously been cited as candidate risk factors for PD with on-pump surgery.^{3,6,8,10,19} And although there are only a few studies available describing PD after TAVR, the data available thus far are consistent with those from this investigation suggesting that PD is an equally prevalent complication for TAVR, particularly when performed using alternative access techniques.^{20,21} It is possible that despite the fact that TAVR may be a less-invasive procedure than SAVR,

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TABLE 4. Univariable analysis of delirium among patients undergoing transcatheter aortic valve replacement (TAVR) and surgical aortic valve replacement (SAVR)

Characteristic	TAVR			SAVR		
	No delirium (n = 119)	Delirium (n = 49)	P value	No delirium (n = 173)	Delirium (n = 86)	P value
Age (y)	82 ± 8	81 ± 8	.410	69 ± 11	74 ± 10	<.001
Female	68 (57)	24 (49)	.390	69 (40)	39 (45)	.420
Body mass index	28 ± 6	27 ± 5	.360	31 ± 7	31 ± 9	.990
Society of Thoracic Surgeons predicted risk of mortality score	9.5 (5.7, 14.0)	9.2 (6.4, 12.0)	.920	2.2 (1.2, 4.7)	4.2 (2.3, 7.5)	<.001
American Society of Anesthesiologists class ≥ 4	115 (97)	47 (96)	>.999	110 (64)	70 (81)	.004
New York Heart Association functional class III-IV	103 (87)	44 (90)	.800	81 (47)	58 (67)	.002
Diabetes mellitus	50 (42)	22 (45)	.740	59 (34)	36 (42)	.270
Hypertension	111 (93)	45 (92)	.750	138 (80)	71 (83)	.620
Coronary artery disease	100 (84)	43 (88)	.640	70 (40)	44 (51)	.110
Cerebrovascular disease	35 (29)	17 (35)	.580	34 (20)	23 (27)	.210
Atrial fibrillation	42 (35)	23 (47)	.170	29 (17)	20 (23)	.240
Chronic lung disease*	39 (33)	20 (41)	.370	19 (11)	20 (23)	.016
Peripheral vascular disease	80 (67)	33 (67)	>.999	42 (24)	29 (34)	.140
Preoperative glomerular filtration rate (mL/min)	63 ± 21	68 ± 38	.390	76 ± 25	71 ± 27	.130
Chronic steroid use	9 (8)	12 (24)	.004	9 (5)	6 (7)	.580
Hemoglobin (g/dL)	11.4 ± 1.5	11.1 ± 1.5	.230	12.8 ± 1.9	11.9 ± 1.7	<.001
Albumin (g/dL)	4.01 ± 0.4	4 ± 0.5	.900	4.12 ± 0.4	4 ± 0.4	.070
Procedure time (min)	150 ± 60	160 ± 53	.370	251 ± 76	258 ± 66	.480
TAVR approach			.013			
TAVR-transfemoral	47 (39)	10 (20)				
TAVR-transapical/transaortic	72 (61)	39 (80)				
Minimum alveolar concentration volatile anesthetic	0.95 ± 0.13	0.95 ± 0.13	.980	0.87 ± 0.11	0.85 ± 0.10	.150
Packed red blood cell transfusion	42 (35)	25 (51)	.080	120 (69)	74 (86)	.004

Values are presented as mean ± standard deviation, n (%), or median (first quartile, third quartile). TAVR, Transcatheter aortic valve replacement; SAVR, surgical aortic valve replacement. *Moderate/severe.

given the high-risk patient population for whom TAVR is currently restricted to as a procedural option, the incidence of PD following TAVR will remain high. The results from propensity-matched patients within this study suggest that a typical patient undergoing TAVR, with numerous comorbidities, would likely have an even higher likelihood of PD if offered an SAVR approach. The predictors of PD after TAVR remain poorly defined due to the small data sets currently available. What does appear clear based on the results of this investigation is that the association of PD with perioperative morbidity and mortality after TAVR are consistent with those reported after SAVR

affecting patient outcomes within 30 days and extending out to 1 year.^{4,22}

The implications of this study are several and highlight that increased effort may need to be directed toward the identification of PD after any cardiac surgery, including specifically TAVR procedures. This report suggests that diagnosing PD after TAVR procedures may also be difficult given the frequency of the hypoactive phenotype seen in this investigation, which can be easily overlooked but yet associated with even worse outcomes.⁹ This study reinforces that patients with PD represent a subgroup at higher risk for death up to 1 year postoperatively.^{8,22} By

TABLE 5. Multivariate analysis for delirium in patients undergoing surgical aortic valve replacement.

Variable	Odds ratio	95% Confidence interval	P value
Age (per 1 year)	1.043	1.010-1.078	.011
Society of Thoracic Surgeons predicted risk of mortality score (per 1 point)	0.980	0.904-1.064	.630
Previous percutaneous coronary intervention (yes)	1.531	0.761-3.078	.230
New York Heart Association functional class III-IV (yes)	1.217	0.607-2.443	.580
Lung disease moderate/severe (yes)	1.608	0.652-3.970	.300
Preoperative hemoglobin (per 1 unit)	0.841	0.699-1.012	.070
Mitral regurgitation moderate/severe (yes)	2.939	1.176-7.344	.021
Intraoperative packed red blood cells (yes)	1.034	0.435-2.460	.940
Preoperative intra-aortic balloon pump (yes)	3.509	0.878-14.027	.080
Intubation time < 24 h (yes)	0.400	0.161-0.996	.049

TABLE 6. Multivariate Analysis for Delirium in patients undergoing transcatheter aortic valve replacement (TAVR)

Variable	Odds ratio	95% Confidence interval	P value
Age (per 1 year)	0.982	0.940-1.026	.420
Society of Thoracic Surgeons predicted risk of mortality score (per 1 unit)	0.962	0.899-1.029	.260
Preoperative steroids (yes)	3.944	1.371-11.376	.011
TAVR via transapical or transaortic approach	2.002	0.851-4.710	.110
Intraoperative packed red blood cells (yes)	2.000	0.919-4.352	.080
Acute kidney injury present (yes)	3.473	1.191-10.129	.023

TAVR, Transcatheter aortic valve replacement.

identifying these patients in hospital, an opportunity may be provided to increase the awareness of patients and caregivers of delirium and its potential complications. Resources could be targeted to prevent hospital readmissions, and approaches could be explored to mitigate larger issues such as increased mortality and cognitive decline.^{4,22} From a pragmatic standpoint, this may require that delirium assessments be performed even after patients have left the ICU with different instruments such as the 3 dimensional CAM, which may be better suited than the CAM-ICU toward diagnosing PD in the stepdown setting.²³

This study does suggest that maintaining a TF first approach for TAVR has the greatest potential to further decrease the incidence of PD after AVR. Because TAVR delivery systems decrease in size, patients eligible for TAVR will more likely be TF candidates, with only few patients requiring TA or TAO procedures. And although the results of this investigation and those of Eide and colleagues^{12,21} suggest that TAVR-TF has the lowest incidence of PD among all AVR procedures currently performed, smaller TAVR devices now allow for TF procedures to be performed with a totally percutaneous approach, without the need for any incision or general anesthesia. Despite anesthetic use not having been shown to be a factor associated with PD in this study, the general premise that less postoperative pain and providing a lighter depth of anesthesia in older patients will reduce postoperative delirium²³ has implications for all AVR procedures and is currently the focus of an ongoing prospective trial (NCT02241655). These efforts for a minimalist approach to AVR, while not specifically addressing the predisposition or vulnerability of older patients susceptible to developing PD, focus on minimizing the precipitating factors for PD as much as possible.²⁴

Limitations

This study has several limitations. The patients studied in both the SAVR and TAVR cohorts for the development of

PD were quite different with respect to the severity and frequency of their comorbidities, potentially biasing the frequency of PD within each cohort. Assessments for PD were done for patients while in the ICU only, and by the CAM-ICU only. Although the CAM-ICU is highly specific, its sensitivity is less, predisposing to underreporting of PD for both the TAVR and SAVR groups.

The relatively small number of patients with PD, particularly in the TAVR, cohort limited the ability to identify clinically significant, independent predictors for delirium after TAVR. Greater numbers of patients will be required to better delineate independent risk factors for PD for TAVR and these types of patients. Given the association between PD and later cognitive impairment, future studies should incorporate cognitive and frailty assessments with PD because these metrics are particularly valuable to this older population.

Lastly, because this was a retrospective analysis for PD, it is impossible to account for the many confounding variables that exist for the assessment of PD and future prospective studies will be required to further explore the findings of this investigation.

CONCLUSIONS

PD occurs commonly after SAVR and TAVR and is associated with increased perioperative morbidity and resource use. Moreover, the development of PD is associated with increased mortality at both 30 days and over the first year after procedure. Although further studies are needed to confirm and extend the findings of this investigation, addressing PD and any potentially modifiable risk factors for its development has the potential to reduce this specific postoperative complication and may also influence other associated adverse outcomes after AVR therapy.

Conflict of Interest Statement

Dr Damiano reports consulting fees from Atricure, lecture fees and grant support from Edwards and On-X Lifesciences, and grant support from Thrasos Inc. Dr Lindman reports consulting fees from Roche Diagnostics. Dr Lasala reports consulting fees from Boston Scientific; equity ownership in Abiomed and DirectFlow; and lecture fees from Abiomed, St. Jude, Boston Scientific, and Eli Lilly. Dr Melby reports consulting fees from On-X and lecture fees from ClearFlow. All other authors have nothing to disclose with regard to commercial support.

You can watch a Webcast of this AATS meeting presentation by going to: http://webcast.aats.org/2015/Video/Monday/04-27-15_608_1545_Maniar.mp4.

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Key Words: TAVR, delirium, cardiac surgery, aortic stenosis, aortic valve replacement

Discussion

Dr Lars G. Svensson (Cleveland, Ohio). Your work raises more questions than answers concerning the whole issue of neurologic injury after either transcatheter aortic valve repair (TAVR) or aortic valve replacement (AVR). Ideally it would have been interesting to see a comparison using postoperative diffusion-weighted magnetic resonance imaging (MRI) and higher executive function neuropsychologic testing.

I could ask questions specifically related to postoperative testing to diagnose the cause of the symptom of delirium; however, I would rather encourage you to do a prospective study examining MRI and neurocognitive higher function testing.

As we try to understand postoperative consequences of TAVR, it is important to note that MRI brain lesions are associated with a greater risk of recurrent events and later development of Alzheimer disease or dementia. Indeed, Ghanem found that 1 of 8 patients after TAVR at 2 years had deterioration of Repeatable Battery for the Assessment of Neuropsychological Status assessment. Related also to your study, Rodes-Cabau found no difference between a transapical versus transfemoral approach.

You showed differences in delirium rates, yet the key is higher neurocognitive function at 30 days. That would be an important study but should also be supported by MRI evaluation. For example, we don't know what role anesthesia may have played.

In a study from my group that will be shortly published in this journal, we found that both tests are needed because neurocognitive testing may not correlate with any MRI brain lesions, and alternatively, positive brain lesions on MRI may not be detected by neurocognitive testing.

I have 1 question: Will you do a prospective study to better elucidate the neurologic injuries that can be associated with the delirium?

Dr Maniar. The simple answer to your question is yes. You raise several issues in regard to postoperative delirium. It is not clear whether postoperative delirium is for a symptom of something else occurring within a patient. MRI-based assessments are going to be what is needed to determine that.

We are going to see some of these data come out through the SENTINEL trial where embolic filters are placed in patients during TAVR, and we are going to have the imaging available to us. We will have to focus on prospective studies that also include neurocognitive testing. As you know, there is a suggestion that patients with impaired neurocognitive assessments preoperatively

are also then part of the subset that have most of the problems postoperatively. I think it will be 2-fold: looking at true incidence, and helping elaborate on modifiable or identify new risk factors.

Regarding anesthesia, that is really going to be an interesting area. TAVR has now moved into the second generation where now the majority of transfemoral procedures are going to be done percutaneously. There is also a movement afoot by our anesthesia colleagues in a

multicenter trial that we're participating in where sedation is going to be individualized to a given patient looking at monitoring brain function during the anesthetic procedure in an attempt to not overanesthetize some of our patients.

We are going to see better testing, better imaging, and I think more fine-tuning of our anesthesia, and it will be interesting to see the differences that play out here.

Dr Svensson. Thank you.

EDITORIAL COMMENTARY

Delirium after aortic replacement: A transvalvular approach is no panacea

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Postoperative delirium (PD) is an increasingly recognized complication of all surgical procedures, both minimally invasive and other. Surgery for aortic valve replacement is no exception, and Maniar and colleagues¹ provide the largest study to date of PD after transcatheter aortic valve replacement (TAVR). Despite the relatively less invasive nature of TAVR, compared with open surgical aortic valve replacement (SAVR), the overall incidence of PD in this retrospective study was essentially the same (30% for SAVR vs 29% for TAVR; $P = .40$). On the surface, this result may seem disappointing, ie, that a relatively "less invasive" procedure would not lower the incidence of this serious complication. However, the TAVR approaches varied considerably (including transfemoral, transapical, and transaortic), and in a subset analysis of the transfemoral approach, which is arguably the least invasive, the PD incidence was, in fact, lower than that with SAVR (18% vs 33%, respectively; $P = .025$).

However, the lack of difference in the primary comparison, even ignoring this subset analysis, may be due to the multifactorial nature of delirium. The etiology

and pathophysiology of PD is incredibly complex; it may depend as much on nonsurgical factors as on actual surgical trauma and its associated inflammatory response, which is commonly included in discussions of delirium pathophysiology. Similarly, this complexity may partially explain the lack of PD difference in the 2 groups. Inflammation from TAVR is thought to be lower, compared with SAVR,² but it is only one possible target for addressing delirium. Further, the importance of inflammation as an etiologic factor in PD has been questioned recently, given the failure of high-dose steroids to mitigate its occurrence.³

Hilary P. Grocott, MD, FRCPC

Central Message

Delirium remains an important neurologic sequela of aortic valve replacement, despite use of a more minimally invasive transvalvular approach.

See Article page 815.

FIGURE E1. Covariate balance showing the standardized difference between transcatheter aortic valve replacement (*TAVR*) and surgical aortic valve replacement (*SAVR*) before and after propensity score matching.

FIGURE E2. Distribution of propensity scores before matching to evaluate overlap. *STS*, Society of Thoracic Surgeons; *CAD*, coronary artery disease; *PVD*, peripheral vascular disease; *NYHA*, New York Heart Association, *Hgb*, hemoglobin; *GFR* glomerular filtration rate, *CVD*, cerebrovascular disease; *DM*, diabetes mellitus.

PM

TABLE E1. Kaplan-Meier estimates of mortality among patients undergoing transcatheter aortic valve replacement (TAVR) and surgical aortic valve replacement (SAVR)

AVR type	Time from implant	Without delirium		With delirium		<i>P</i> value
		Survival %	95% Confidence interval	Survival %	95% Confidence interval	
TAVR	30 d	98.3	93.4-99.6	87.8	74.8-94.3	.029
	6 mo	89.5	82.2-93.9	79.1	64.7-88.2	.110
	1 y	80.4	71.1-87.0	66.4	50.0-78.5	.090
SAVR	30 d	1.00	1.00-1.00	95.3	88.1-98.2	.041
	6 mo	98.8	95.4-99.7	88.4	79.4-93.6	.003
	1 y	97.3	92.7-99.0	82.8	72.6-89.5	.001

AVR, Aortic valve replacement; TAVR, transcatheter aortic valve replacement; SAVR, surgical aortic valve replacement.