Procedural Results and Clinical Outcomes of Transcatheter Aortic Valve Implantation in Switzerland: An Observational Cohort Study of Sapien 3 Versus Sapien XT Transcatheter Heart Valves

BINDER, RK, et al.

Abstract

BACKGROUND: New generation transcatheter heart valves (THV) may improve clinical outcomes of transcatheter aortic valve implantation. METHODS AND RESULTS: In a nationwide, prospective, multicenter cohort study (Swiss Transcatheter Aortic Valve Implantation Registry, NCT01368250), outcomes of consecutive transfemoral transcatheter aortic valve implantation patients treated with the Sapien 3 THV (S3) versus the Sapien XT THV (XT) were investigated. An overall of 153 consecutive S3 patients were compared with 445 consecutive XT patients. Postprocedural mean transprosthetic gradient (6.5±3.0 versus 7.8±6.3 mmHg, P=0.17) did not differ between S3 and XT patients, respectively. The rate of more than mild paravalvular regurgitation (1.3% versus 5.3%, P=0.04) and of vascular (5.3% versus 16.9%, P

Reference


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Procedural Results and Clinical Outcomes of Transcatheter Aortic Valve Implantation in Switzerland

An Observational Cohort Study of Sapien 3 Versus Sapien XT Transcatheter Heart Valves

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Methods and Results—In a nationwide, prospective, multicenter cohort study (Swiss Transcatheter Aortic Valve Implantation Registry, NCT01368250), outcomes of consecutive transfemoral transcatheter aortic valve implantation patients treated with the Sapien 3 THV (S3) versus the Sapien XT THV (XT) were investigated. An overall of 153 consecutive S3 patients were compared with 445 consecutive XT patients. Postprocedural mean transprosthetic gradient (6.5±3.0 versus 7.8±6.3 mm Hg, P=0.17) did not differ between S3 and XT patients, respectively. The rate of more than mild paravalvular regurgitation (1.3% versus 5.3%, P=0.04) and of vascular (5.3% versus 16.9%, P<0.01) complications were significantly lower in S3 patients. A higher profile delivery system was observed in patients receiving the S3 valve (17.0% versus 11.0%, P=0.01). There were no significant differences for disabling stroke (S3 1.3% versus XT 3.1%, P=0.29) and all-cause mortality (S3 3.3% versus XT 4.5%, P=0.27).

Conclusions—The use of the new generation S3 balloon-expandable THV reduced the risk of more than mild paravalvular regurgitation and vascular complications but was associated with an increased permanent pacemaker rate compared with the XT. Transcatheter aortic valve implantation using the newest generation balloon-expandable THV is associated with a low risk of stroke and favorable clinical outcomes.

Clinical Trial Registration—URL: http://www.clinicaltrials.gov. Unique identifier: NCT01368250.

(Scirc Cardiovasc Interv. 2015;8:e002653. DOI: 10.1161/CIRCINTERVENTIONS.115.002653.)

Key Words: aortic valve stenosis ■ bleeding ■ transcatheter aortic valve replacement ■ transcatheter heart valve ■ transcutaneous aortic valve implantation ■ vascular complications

Since the first transcatheter aortic valve implantation (TAVI) in 2002 and the establishment of the retrograde transfemoral approach in 2005, the procedure has undergone further refinements. Lower profile delivery systems, multi-modality imaging for patient screening and device deployment, transcatheter heart valve (THV) sizing algorithms, and modifications of prosthesis design and delivery systems have reduced the rate of vascular complications and paravalvular regurgitation (PAR) and increased the safety and efficacy of TAVI. Although the procedure was initially restricted to inoperable patients, it is currently approved for operable patients at high surgical risk. Recently, a randomized trial

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A list of collaborators and Swiss TAVI Investigators is available in the Data Supplement.

*Drs Binder and Stortecky contributed equally to this work.

The Data Supplement is available at http://circinterventions.ahajournals.org/lookup/suppl/doi:10.1161/CIRCINTERVENTIONS.115.002653/-/DC1. Correspondence to Peter Wenaweser, MD, Department of Cardiology, Bern University Hospital, CH-3010 Bern, Switzerland. E-mail peter.wenaweser@insel.ch

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WHAT IS KNOWN

- Transcatheter aortic valve implantation (TAVI) with the Sapien XT transcatheter heart valve (THV) is a valuable alternative to surgical aortic valve replacement in selected patients.
- However, TAVI is associated with vascular and bleeding complications, paravalvular regurgitation, and atriointimal conduction disturbances.

WHAT THE STUDY ADDS

- In this preliminary comparison, the use of the new generation Sapien 3 THV was associated with a lower incidence of vascular complications and less paravalvular regurgitation compared with TAVI with the Sapien XT THV.
- The rate of new pacemaker implantation was higher after TAVI with the Sapien 3 THV than after TAVI with the Sapien XT THV.

has indicated superiority of TAVI over surgical aortic valve replacement for 1-year survival in patients with symptomatic severe aortic stenosis and a mean Society of Thoracic Surgeons Predicted Risk of Mortality (STS PROM) of 7.3±3.0%, indicating intermediate surgical risk.

In 2014, the newest generation balloon-expandable THV (Sapien 3, S3; Figure 1) received regulatory approval and was introduced in Switzerland and subsequently replaced its predecessor the Sapien XT (XT, Figure 2) THV as the default balloon-expandable THV for TAVI. The S3 may be delivered via a lower profile delivery system and incorporates a sealing cuff intended to reduce PAR. Despite positive results during the first-in-human S3 experience and subsequent small series, it is not established whether the new features of the S3 will translate into improved procedural and clinical outcomes compared with the XT. We therefore analyzed and compared all patients who underwent transfemoral TAVI with the S3 or the XT in the prospective, nationwide Swiss TAVI registry in Switzerland (ClinicalTrials.gov NCT01368250).

Methods

The Swiss TAVI registry is—a national, prospective cohort study of all TAVI procedures performed in Switzerland aiming for consecutive patient enrollment and with data monitoring as well as end point adjudication by a dedicated clinical events committee according to the recommendations of the Valve Academic Research Consortium.18 The Swiss TAVI registry was designed to provide short-term clinical outcomes and long-term clinical data of TAVI patients treated with CE-approved devices. The study protocol was approved by the local cantonal ethics committee and institutional review boards at each participating center, and all patients provided written informed consent. The Swiss TAVI registry is performed under the lead of the Swiss Cardiovascular Center Bern at Bern University Hospital in cooperation with the Clinical Trials Unit Bern responsible for data management and independent statistical analysis.

For this analysis, all patients of the Swiss TAVI registry who underwent transfemoral TAVI with either the XT or the S3 THV were analyzed (inclusion period: XT, February 2011 to January 2014; S3, February 2014 to June 2014). The grade of PAR was assessed by transthoracic echocardiography before hospital discharge by highly experienced echocardiographers according to Valve Academic Research Consortium-2 guidelines.18 Prespecified end points were more than mild PAR, vascular complications, major bleeding, new permanent pacemaker implantation (PPM), disabling stroke, and mortality after 30-day of follow-up.

Statistical Analysis

Continuous data are reported as means±standard deviation (SD), and categorical variables are reported as number of patients (% of patients). Events are reported as counts of first occurrence per (sub) type of event (% of all patients). Event probabilities at 30 days were compared for patients treated with the XT versus the S3 bioprosthesis using logistic regressions. Reported are crude odds ratios (95% confidence interval) with P values from Wald χ² tests corrected for random effects of the hospital identifier using mixed effects logistic regressions or exact logistic regression odds ratios with P values from exact tests in case of zero events. Reported are adjusted odds ratio (95% confidence interval), with the 2 valves compared using mixed effects logistic regressions, including (1) adjustment for TAVI procedure date (ie, to account for a potential learning effect of time), (2) random effect of hospital identifier, and (3) adjustment for baseline characteristics using inverse probability of treatment weights (ie, to account for potential disbalances between the 2 valve types concerning the patient population treated). The estimates of adjusted odds ratio from 20 data sets after multiple imputation of missing values were combined using Rubin’s rule and presented with adjusted P values (P_adj).

Inverse probability of treatment weights for S3 versus XT THV was calculated within each of the 20 data sets using the following baseline variables: age, sex, body mass index, diabetes mellitus, dyslipidemia, hypertension, previous pacemaker, history of myocardial infarction, cardiac surgery, cerebrovascular event, peripheral vascular disease, chronic obstructive pulmonary disease, coronary artery disease, left ventricular ejection fraction, aortic valve area, mean aortic valve gradient, moderate or severe mitral regurgitation, New York Heart Association class III or IV, Canadian Cardiovascular Society angina class none or I/II or III/IV, logistic EuroSCORE, STS PROM score, and valve size. No adjusted analyses were performed

Figure 1. Aortic root angiogram after Sapien 3 transcatheter heart valve implantation. The Sapien 3 transcatheter heart valve comprises a balloon expandable, cobalt chromium frame, a trileaflet bovine pericardial tissue valve, and a polyethylene terephthalate (PET) skirt. The outer PET cuff was designed to improve paravalvular sealing.
Significant differences in the occurrence of PAR (Figure 3) were observed between S3 and XT patients. In more than half of S3 patients, no PAR was detected (57.3%), although this was observed in only one third of XT patients (31.9%, \( P<0.01 \)). Mild PAR was also less frequent in S3 compared with XT patients (S3 41.3% versus XT 62.9%, \( P<0.01 \)). Furthermore, the rate of more than mild PAR was significantly lower in S3 compared with XT patients (S3 1.3% versus XT 5.3%, \( P=0.04 \)).

At 30-day (Table 3) follow-up, mortality did not differ between S3 and XT patients (S3 3.3% versus XT 4.5%, \( P=0.52, \ P_{\text{adj}}=0.27 \)). Major disabling stroke was low in both groups (S3 1.3% versus 3.1%, \( P=0.24, \ P_{\text{adj}}=0.29 \)). The rate of PPM implantation was higher in S3 patients (S3 17.0% versus XT 11.0%, \( P=0.06, \ P_{\text{adj}}=0.01 \)). Major bleeding occurred twice as often in XT patients than in S3 patients (S3 3.9% versus XT 8.3%, \( P=0.11, \ P_{\text{adj}}=0.81 \)) albeit not significantly different, but the rate of vascular complications (major and minor) was significantly higher in XT patients (S3 5.2% versus XT 16.9%, \( P<0.01, \ P_{\text{adj}}<0.01 \)).

**Discussion**

This study sought to investigate differences in procedural and clinical outcomes of patients undergoing transfemoral TAVI with the S3 versus the XT THV. Analysis of our nationwide, prospective Swiss TAVI registry showed that TAVI with the S3 significantly reduced PAR and vascular complications in comparison to TAVI with the XT.

The success of TAVI depends on the risk of perioperative complications, the predictability of the procedure, and device durability. Within the last decade, multimodality imaging for patient screening, patient selection, and device deployment and iterations to the bioprostheses and refinement of delivery systems have contributed to the successful global spread of TAVI as an alternative to surgical aortic valve replacement. Minimizing the rate of periprocedural complications is mandatory to broaden the indication of TAVI from prohibitive or high surgical risk to intermediate\(^{13,19}\) and low surgical risk\(^{19}\) patients. Considering the S3 as a step into this direction has to be based on firm scientific evidence. Important complications of TAVI that need to be reduced are stroke, PAR, vascular and bleeding complications, and atrioventricular block.

**Paravalvular Regurgitation**

PAR is frequently observed after TAVI\(^{20}\) and is associated with worse survival in patients with moderate to severe PAR.\(^{21}\) Whether mild PAR is an independent mortality predictor, as suggested by a previous study,\(^{22}\) is a matter of controversy. Important predictors for PAR include severe leaflet, annulus and left ventricular outflow tract calcifications, THV undersizing, and THV malpositioning. New THV designs with peri-prosthetic sealing cuffs (eg, the S3) may contribute to a reduction in PAR. In our study, more than mild PAR was less frequently observed after TAVI with the S3 compared with the XT. This may be attributed to the external skirt of the S3. However, improved sizing algorithms and a broader landing zone of the elongated S3 stent frame may also have contributed to the difference. As more than mild PAR is associated with higher mortality,\(^{21}\) this difference may translate into improved TAVI
outcomes. However, as the rate of more than mild PAR was low in our cohort, it did not impact short-term survival.

**Stroke**

Compared with medical management, TAVI is associated with an increased stroke risk. Furthermore, in the Placement of Aortic Transcatheter Valve (PARTNER) trial, patients undergoing TAVI had a higher 30-day rate of any cerebrovascular event compared with patients randomized to surgical aortic valve replacement. Subsequent studies with newer generation devices and large registries have further calmed the debate about TAVI associated stroke risk. In the French Aortic National CoreValve and Edwards Registry (FRANCE II) study, stroke rates were 2.3%, and in the United Kingdom Transcatheter Aortic Valve Implantation (UK TAVI) registry, the rate was 4.1%. In our study, the 30-day disabling stroke rate with the S3 in an all-comer population was as low as 1.3%, which was numerically lower than that for the XT. If and how the incidence of stroke can be further reduced is a matter of debate. Cerebral protection devices were designed to capture or deflect debris during TAVI, which would have otherwise embolized to the brain. However, there is currently no evidence that supports the routine use of these devices. The clinical significance of a reduction in subclinical lesions on brain scanning post TAVI, which has been shown with the Claret device (Claret Embolic Protection and TAVI [CLEAN-TAVI] trial, NCT01833052, presented at Transcatheter Cardiovascular Therapeutics Congress 2014), was not established. Future clinical trials are needed to prove whether these devices effectively reduce the risk of stroke during TAVI. In our study population, a cerebral protection device was rarely used and not documented in the files.

**Vascular Complications**

Major vascular complications during TAVI are independent predictors of mortality. With the first generation balloon-expandable THV, major vascular complications occurred in

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**Table 1. Baseline Characteristics**

<table>
<thead>
<tr>
<th></th>
<th>Sapien 3, N=153</th>
<th>Sapien XT, N=445</th>
<th>P Value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Age, years</td>
<td>82.2±6.05</td>
<td>82.2±6.75</td>
<td>0.94</td>
</tr>
<tr>
<td>Female gender, n (%)</td>
<td>72 (47.1%)</td>
<td>249 (55.8%)</td>
<td>0.07</td>
</tr>
<tr>
<td>Body mass index, kg/m²</td>
<td>26.9±5.56</td>
<td>26.7±4.95</td>
<td>0.75</td>
</tr>
<tr>
<td>Cardiac risk factors</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Diabetes mellitus, n (%)</td>
<td>39 (25.5%)</td>
<td>112 (25.1%)</td>
<td>0.92</td>
</tr>
<tr>
<td>Dyslipidemia, n (%)</td>
<td>65 (42.5%)</td>
<td>236 (52.9%)</td>
<td>0.03</td>
</tr>
<tr>
<td>Hypertension, n (%)</td>
<td>117 (76.5%)</td>
<td>353 (79.1%)</td>
<td>0.49</td>
</tr>
<tr>
<td>Past medical history</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Previous pacemaker implantation, n (%)</td>
<td>15 (9.8%)</td>
<td>35 (7.8%)</td>
<td>0.49</td>
</tr>
<tr>
<td>Previous myocardial infarction, n (%)</td>
<td>24 (15.7%)</td>
<td>67 (15.0%)</td>
<td>0.89</td>
</tr>
<tr>
<td>Previous cardiac surgery, n (%)</td>
<td>17 (11.1%)</td>
<td>59 (13.2%)</td>
<td>0.57</td>
</tr>
<tr>
<td>Previous cerebrovascular accident, n (%)</td>
<td>20 (13.1%)</td>
<td>51 (11.4%)</td>
<td>0.56</td>
</tr>
<tr>
<td>Clinical features</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Peripheral vascular disease, n (%)</td>
<td>24 (15.7%)</td>
<td>65 (14.6%)</td>
<td>0.79</td>
</tr>
<tr>
<td>Chronic obstructive pulmonary disease, n (%)</td>
<td>22 (14.4%)</td>
<td>52 (11.7%)</td>
<td>0.39</td>
</tr>
<tr>
<td>Coronary artery disease, n (%)</td>
<td>86 (56.2%)</td>
<td>242 (54.3%)</td>
<td>0.71</td>
</tr>
<tr>
<td>Left ventricular ejection fraction, %</td>
<td>56.6±14.67</td>
<td>56.2±13.51</td>
<td>0.78</td>
</tr>
<tr>
<td>Aortic valve area, cm²</td>
<td>0.71±0.23</td>
<td>0.71±0.22</td>
<td>0.88</td>
</tr>
<tr>
<td>Mean transaortic gradient, mm Hg</td>
<td>47.18±22.04</td>
<td>43.74±17.27</td>
<td>0.06</td>
</tr>
<tr>
<td>Mitral regurgitation grade moderate or severe</td>
<td>21 (14.2%)</td>
<td>86 (20.5%)</td>
<td>0.11</td>
</tr>
<tr>
<td>New York Heart Association (NYHA) Class</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>NYHA I or II, n (%)</td>
<td>48 (32.9%)</td>
<td>150 (33.7%)</td>
<td>0.92</td>
</tr>
<tr>
<td>NYHA III or IV, n (%)</td>
<td>98 (67.1%)</td>
<td>295 (66.3%)</td>
<td>0.92</td>
</tr>
<tr>
<td>Canadian Cardiovascular Society Angina Class</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>No angina, n (%)</td>
<td>n=152,</td>
<td>n=446,</td>
<td>0.15</td>
</tr>
<tr>
<td>CCS I or II, n (%)</td>
<td>125 (82.2%)</td>
<td>333 (74.7%)</td>
<td>0.06</td>
</tr>
<tr>
<td>CCS III or IV, n (%)</td>
<td>19 (12.5%)</td>
<td>75 (16.8%)</td>
<td>0.25</td>
</tr>
<tr>
<td>Risk assessment</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Log. EuroScore, %</td>
<td>23.7±15.95</td>
<td>21.0±15.99</td>
<td>0.16</td>
</tr>
<tr>
<td>STS score, %</td>
<td>7.15±5.60</td>
<td>8.52±7.98</td>
<td>0.07</td>
</tr>
</tbody>
</table>

Dyslipidemia was more prevalent in the Sapien XT group. All other baseline characteristics did not differ significantly between groups. CCS indicates Canadian Cardiovascular Society; and STS, Society of Thoracic Surgeons.
16.2% of patients in the PARTNER IB trial.11 Meanwhile, down-sizing of access sheath diameters4 allowing fully percutaneous procedures28 has resulted in decreased vascular complications. In our study, major and minor vascular complications were significantly lower in S3 compared with XT patients. This parallels a study that showed decreased vascular complications with lower-profile compared with large-profile sheaths.4 On a large scale, the reduction of major vascular complications with the S3 delivery system is expected to impact prognosis and speed up postprocedural patient mobilization, allowing earlier ambulation and discharge.

Bleeding
Major bleeding and blood transfusions after TAVI are associated with worse prognosis.29,30 The source of bleeding may be procedure-related (eg, access site, ventricular or aortic perforation) or technically unrelated to TAVI but triggered by periprocedural antithrombotic medication (eg, gastrointestinal). The access site is the most common source of procedure-related bleeding. In this study, major bleeding occurred twice as often in patients receiving the XT than in patients treated with the S3 THV; however, the difference did not reach statistical significance. A lower rate of bleeding with the S3 may be attributed to the lower profile of the introducer sheath and delivery system. This observation parallels a study that compared TAVI outcomes with different sheath sizes4 and may translate into improved outcomes.

Permanent Pacemaker Implantation
Atrioventricular conduction disturbances necessitating PPM implantation are frequently observed after TAVI11 and

### Table 2. Procedural Characteristics

<table>
<thead>
<tr>
<th></th>
<th>Sapien 3, N=153</th>
<th>Sapien XT, N=445</th>
<th>P Value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Procedure time, min</td>
<td>71.7±30.54</td>
<td>71.8±30.58</td>
<td>0.98</td>
</tr>
<tr>
<td>Amount of contrast, mL</td>
<td>158.0±87.39</td>
<td>201.1±95.37</td>
<td>&lt;0.01</td>
</tr>
<tr>
<td>General anesthesia, n (%)</td>
<td>46 (30.1%)</td>
<td>172 (38.7%)</td>
<td>0.06</td>
</tr>
<tr>
<td>Lenght of hospital stay, days</td>
<td>9.0±5.72</td>
<td>9.5±5.31</td>
<td>0.38</td>
</tr>
<tr>
<td>Type of access</td>
<td></td>
<td></td>
<td>0.82</td>
</tr>
<tr>
<td>Percutaneous, n (%)</td>
<td>133 (86.9%)</td>
<td>390 (87.6%)</td>
<td>0.89</td>
</tr>
<tr>
<td>Surgical, n (%)</td>
<td>20 (13.1%)</td>
<td>55 (12.4%)</td>
<td>0.89</td>
</tr>
<tr>
<td>Procedure location</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Catheterization laboratory, n (%)</td>
<td>102 (66.7%)</td>
<td>343 (77.1%)</td>
<td>0.01</td>
</tr>
<tr>
<td>Operating room, n (%)</td>
<td>1 (0.7%)</td>
<td>0 (0.0%)</td>
<td>0.26</td>
</tr>
<tr>
<td>Hybrid room, n (%)</td>
<td>50 (32.7%)</td>
<td>102 (22.9%)</td>
<td>0.02</td>
</tr>
<tr>
<td>Concomitant procedure</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Percutaneous coronary intervention, n (%)</td>
<td>8 (5.3%)</td>
<td>45 (10.1%)</td>
<td>0.07</td>
</tr>
<tr>
<td>Carotid stenting, n (%)</td>
<td>0 (0.0%)</td>
<td>1 (0.2%)</td>
<td>1.00</td>
</tr>
<tr>
<td>Iliofemoral stenting, n (%)</td>
<td>5 (3.3%)</td>
<td>17 (3.8%)</td>
<td>1.00</td>
</tr>
<tr>
<td>Device features</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Valve size</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>23 mm</td>
<td>42 (27.5%)</td>
<td>108 (24.3%)</td>
<td>0.45</td>
</tr>
<tr>
<td>26 mm</td>
<td>72 (47.1%)</td>
<td>257 (57.8%)</td>
<td>0.02</td>
</tr>
<tr>
<td>29 mm</td>
<td>39 (25.5%)</td>
<td>80 (18.0%)</td>
<td>0.05</td>
</tr>
<tr>
<td>Prior balloon aortic valvuloplasty, n (%)</td>
<td>143 (93.5%)</td>
<td>410 (92.1%)</td>
<td>0.72</td>
</tr>
<tr>
<td>Mean transprosthetic gradient, mm Hg</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>For 23 mm valve size</td>
<td>11.6±5.98</td>
<td>9.96±4.77</td>
<td>0.08</td>
</tr>
<tr>
<td>For 26 mm valve size</td>
<td>9.0±3.66</td>
<td>8.18±5.61</td>
<td>0.25</td>
</tr>
<tr>
<td>For 29 mm valve size</td>
<td>8.49±3.42</td>
<td>7.42±4.59</td>
<td>0.23</td>
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<tr>
<td>Aortic valve area, mm</td>
<td></td>
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<td></td>
</tr>
<tr>
<td>For 23 mm valve size</td>
<td>1.43±0.33</td>
<td>1.51±0.43</td>
<td>0.38</td>
</tr>
<tr>
<td>For 26 mm valve size</td>
<td>1.73±0.37</td>
<td>1.89±0.58</td>
<td>0.09</td>
</tr>
<tr>
<td>For 29 mm valve size</td>
<td>1.93±0.50</td>
<td>2.24±0.81</td>
<td>0.15</td>
</tr>
<tr>
<td>Aortic regurgitation post-TAVI</td>
<td>n=150</td>
<td>n=439</td>
<td></td>
</tr>
<tr>
<td>Grade 0, n (%)</td>
<td>86 (57.3%)</td>
<td>140 (31.9%)</td>
<td>&lt;0.01</td>
</tr>
<tr>
<td>Grade 1, n (%)</td>
<td>62 (41.3%)</td>
<td>276 (62.9%)</td>
<td>&lt;0.01</td>
</tr>
<tr>
<td>Grade 2, n (%)</td>
<td>2 (1.3%)</td>
<td>20 (4.6%)</td>
<td>0.08</td>
</tr>
<tr>
<td>Grade 3, n (%)</td>
<td>0 (0.0%)</td>
<td>3 (0.7%)</td>
<td>0.57</td>
</tr>
</tbody>
</table>
mostly depend on the THV type implanted. Although PPM rates of 20% to 30% with the self-expanding CoreValve\textsuperscript{13,31} and almost 30% with the Lotus THV\textsuperscript{32} have been observed, the rate of higher degree atrioventricular block is lower for balloon-expandable THVs.\textsuperscript{33} Additional factors that predict PPM implantation after TAVI include preexisting right bundle branch block\textsuperscript{34} or atrioventricular block, as well as THV implant depth\textsuperscript{35} and annulus oversizing.\textsuperscript{34} In our study, there were more new PPM implants in patients treated with the S3. This could be explained by the longer stent frame of the S3, which may protrude more into the left ventricular outflow tract, thereby compressing the interventricular septum. An inflammatory response to the external sealing skirt may be postulated, but is unlikely. Whether prudent higher THV implantations (80% aortal, 20% ventricular) may reduce the risk of conduction disturbances needs further investigation. Although the initial manufacturer recommendation was to place the middle marker of the deployment balloon in the annular plane, current clinical practice demonstrates that a high implant in experienced hands can be safely performed and may reduce atrioventricular conduction disturbances.\textsuperscript{36} Overall, there seems to be no prognostic impact of a new PPM after TAVI.\textsuperscript{34,37}

**Limitations**

The grade of PAR in this study was defined by experienced on-site echocardiographers and reported according to Valve Academic Research Consortium-2 criteria.\textsuperscript{14} The grading of PAR

Table 3. 30-Days Clinical Outcomes

<table>
<thead>
<tr>
<th></th>
<th>Sapien 3, N=153</th>
<th>Sapien XT, N=445</th>
<th>Odds Ratio OR (95% CI)</th>
<th>P Value</th>
<th>Adjusted OR (95% CI)</th>
<th>Adjusted P Value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Mortality, n(%)</td>
<td>5 (3.3)</td>
<td>20 (4.5)</td>
<td>0.72 (0.26–1.95)</td>
<td>0.56</td>
<td>0.63 (0.27–1.43)</td>
<td>0.27</td>
</tr>
<tr>
<td>Cardiovascular mortality, n (%)</td>
<td>4 (2.6)</td>
<td>19 (4.3)</td>
<td>0.60 (0.20–1.80)</td>
<td>0.36</td>
<td>0.77 (0.32–1.81)</td>
<td>0.55</td>
</tr>
<tr>
<td>Cerebrovascular accident, n (%)</td>
<td>2 (1.3)</td>
<td>18 (4.0)</td>
<td>0.31 (0.07–1.37)</td>
<td>0.12</td>
<td>0.35 (0.10–1.15)</td>
<td>0.08</td>
</tr>
<tr>
<td>Disabling stroke, n (%)</td>
<td>2 (1.3)</td>
<td>14 (3.1)</td>
<td>0.41 (0.09–1.81)</td>
<td>0.24</td>
<td>0.44 (0.10–1.99)</td>
<td>0.29</td>
</tr>
<tr>
<td>Nondisabling stroke, n (%)</td>
<td>0 (0.0)</td>
<td>2 (0.4)</td>
<td>1.20 (0.03–15.51)</td>
<td>1.00</td>
<td></td>
<td></td>
</tr>
<tr>
<td>TIA, n (%)</td>
<td>0 (0.0)</td>
<td>2 (0.4)</td>
<td>1.20 (0.03–15.51)</td>
<td>1.00</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Myocardial infarction, n (%)</td>
<td>2 (1.3)</td>
<td>0 (0.0)</td>
<td>7.06 (0.55–∞)</td>
<td>0.13</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Periprocedural myocardial infarction, n (%)</td>
<td>2 (1.3)</td>
<td>0 (0.0)</td>
<td>7.06 (0.55–∞)</td>
<td>0.13</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Spontaneous myocardial infarction, n (%)</td>
<td>0 (0.0)</td>
<td>0 (0.0)</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Acute kidney injury, n (%)</td>
<td>7 (4.6)</td>
<td>26 (5.8)</td>
<td>0.83 (0.35–1.98)</td>
<td>0.89</td>
<td>1.62 (0.54–4.86)</td>
<td>0.39</td>
</tr>
<tr>
<td>Stage 1, n (%)</td>
<td>1 (0.7)</td>
<td>13 (2.9)</td>
<td>0.26 (0.03–2.08)</td>
<td>0.21</td>
<td>0.80 (0.18–3.58)</td>
<td>0.77</td>
</tr>
<tr>
<td>Stage 2, n (%)</td>
<td>2 (1.3)</td>
<td>3 (0.7)</td>
<td>1.95 (0.32–11.79)</td>
<td>0.47</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Stage 3, n (%)</td>
<td>4 (2.6)</td>
<td>10 (2.2)</td>
<td>1.17 (0.36–3.78)</td>
<td>0.80</td>
<td>2.79 (0.56–13.94)</td>
<td>0.21</td>
</tr>
<tr>
<td>Bleeding, n (%)</td>
<td>14 (9.2)</td>
<td>66 (14.8)</td>
<td>0.50 (0.26–0.99)</td>
<td>0.05</td>
<td>0.76 (0.24–2.40)</td>
<td>0.64</td>
</tr>
<tr>
<td>Life threatening bleeding, n (%)</td>
<td>6 (3.9)</td>
<td>24 (5.4)</td>
<td>0.64 (0.24–1.68)</td>
<td>0.36</td>
<td>1.16 (0.56–2.40)</td>
<td>0.68</td>
</tr>
<tr>
<td>Major bleeding, n (%)</td>
<td>6 (3.9)</td>
<td>37 (8.3)</td>
<td>0.48 (0.19–1.18)</td>
<td>0.11</td>
<td>0.84 (0.21–3.45)</td>
<td>0.81</td>
</tr>
<tr>
<td>Minor bleeding, n (%)</td>
<td>2 (1.3)</td>
<td>5 (1.1)</td>
<td>0.93 (0.13–6.59)</td>
<td>0.71</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Vascular access site and access-related complications, n (%)</td>
<td>8 (5.2)</td>
<td>75 (16.9)</td>
<td>0.25 (0.11–0.57)</td>
<td>&lt;0.01</td>
<td>0.31 (0.17–0.59)</td>
<td>&lt;0.01</td>
</tr>
<tr>
<td>Major vascular complications, n (%)</td>
<td>5 (3.3)</td>
<td>41 (9.2)</td>
<td>0.31 (0.11–0.85)</td>
<td>0.02</td>
<td>0.53 (0.27–1.04)</td>
<td>0.07</td>
</tr>
<tr>
<td>Minor vascular complications, n (%)</td>
<td>2 (1.3)</td>
<td>34 (7.6)</td>
<td>0.16 (0.03–0.74)</td>
<td>0.02</td>
<td>0.09 (0.04–0.19)</td>
<td>&lt;0.01</td>
</tr>
<tr>
<td>Repeat unplanned intervention, n (%)</td>
<td>2 (1.3)</td>
<td>2 (0.4)</td>
<td>2.93 (0.41–21.01)</td>
<td>0.28</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Valve in valve treatment, n (%)</td>
<td>0 (0.0)</td>
<td>0 (0.0)</td>
<td>2.91 (0.00–113.43)</td>
<td>1.00</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Permanent pacemaker implantation, n (%)</td>
<td>26 (17.0)</td>
<td>49 (11.0)</td>
<td>1.68 (0.99–2.84)</td>
<td>0.06</td>
<td>1.89 (1.16–3.08)</td>
<td>0.01</td>
</tr>
</tbody>
</table>

Depicted are number of first events with % of all patients at 30 days since procedure. Odds ratios (OR) from mixed effects logistic regressions accounting for random hospital identifier effects or exact logistic regressions in case of zero events (95% confidence interval [CI]). Adjusted odds ratios: see Methods for details. TIA indicates transient ischemic attack.
after TAVI may be difficult and substantial inter- and intra-
observer variability may occur. The lack of a core laboratory
may lead to heterogeneity in the assessment of this parameter.
However, all sites contributed patients to both groups, which
reduces center-specific assessments as a confounder, and out-
come assessments were corrected using random effects of
the site.

As the S3 replaced the XT as default balloon-expandable
THV, both groups were treated consecutively. A learn-
ing curve may be postulated explaining improved outcomes
with the S3. However, all participating centers have started
and gained extensive experience with TAVI before the SWISS
TAVI registry was initiated. Furthermore, the introduction of
a new device implicated a new learning curve for the S3, which
would be in favor of the XT. Therefore, we do not anticipate
that a learning curve explains the observations of this trial.

Assessments of clinical outcomes were not corrected for
multiple testing, which may lead to the reporting of spuri-
ous significant effects. The reporting in this study followed
the Valve Academic Research Consortium-2 criteria and were
predefined. The reduction in vascular access site–related com-
lications does withstand correction for multiple testing by the
Bonferroni method (0.05 divided by 9 main outcomes: 0.005).
Otherwise, further assessments of clinical outcomes compar-
ing S3 versus XT is encouraged using a larger sample size
of patients and longer follow-up. Because of the prospective
design of this nationwide multicenter registry, data collection
was restricted to variables defined at the launch of the registry.
Therefore, no information on specific sizing algorithms and
prosthesis implant depth are available.

Conclusions
The use of the new generation S3 balloon-expandable THV
is associated with a significant reduction of more than mild PAR
and vascular complications when compared with the XT. In
temporary clinical practice, TAVI using the newest genera-
tion balloon-expandable THV is associated with a low risk of
stroke and overall favorable clinical outcomes.

Disclosures
Dr Binder serves as consultant to Edwards Lifesciences and pro-
tor to Boston Scientific. Dr Jeger serves as a consultant to St Jude
Medical and has received reimbursement for travel expenses from
Medtronic, Boston Scientific, and Edwards Lifesciences. Dr Tueller
received speaker fees from Edwards Lifesciences and travel ex-
penses from Medtronic. Dr Toggweiler received speakers fees from
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Edwards Lifesciences and St Jude Medical. Dr Huber is a proctor for
Edwards Lifesciences and Consultant for Medtronic. Dr Windecker
has received research contracts to the institution from Abbott, Boston
Scientific, Biosensors, Cordis, Medtronic, and St Jude. Dr Wenaweser
serves as proctor for Medtronic, Edwards Lifesciences, and Boston
Scientific and has received an unrestricted grant from Medtronic to
the institution (University of Bern). All the other authors have no con-
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CIRCINTERVENTIONS.111.966531.
Procedural Results and Clinical Outcomes of Transcatheter Aortic Valve Implantation in Switzerland: An Observational Cohort Study of Sapien 3 Versus Sapien XT Transcatheter Heart Valves

Ronald K Binder, Stefan Stortecky, Dik Heg, David Tueller, Raban Jeger, Stefan Toggweiler, Giovanni Pedrazzini, Franz W Amann, Enrico Ferrari, Stephane Noble, Fabian Nietlispach, Francesco Maisano, Lorenz Räber, Marco Roffi, Jürg Grünenfelder, Peter Jüni, Christoph Huber, Stephan Windecker and Peter Wenaweser

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Supplemental Material

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