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Thoracic Endovascular Aortic Repair (TEVAR) in Proximal (Type A) Aortic Dissection: Ready for a Broader Application?

4

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33	Abbreviations	
34		
35	aTAAD	acute type A aortic dissection
36	СТ	computed tomography
37	cTAAD	subacute or chronic type A aortic dissection
38	D	diameter
39	EKG	electrocardiogram
40	EuroSCORE II	European System for Cardiac Operative Risk Evaluation II
41	F	female
42	IRAD	International Registry of Acute Aortic Dissections
43	L	length
44	LCCA	left common carotid artery
45	Μ	male
46	Ν	number
47	NA	not applicable
48	NBS	non-bare stent
49	PA	pseudoaneurysm
50	RCA	right carotid artery
51	RFV	right femoral vein
52	SD	standard deviation
53	SG	stent-graft
54	ТА	transapical
55	TAVR	transcatheter aortic valve replacement
56	TAx	transaxillary
57	TEVAR	thoracic endovascular aortic repair
58	TF	transfemoral
59		
60	Y	

61	ABSTRACT
62	
63	OBJECTIVE: Thoracic endovascular aortic repair (TEVAR) has demonstrated
64	encouraging results and is gaining increasing acceptance as a treatment option for
65	aortic aneurysms and dissections. Yet, its role in managing proximal aortic
66	pathologies is unknown - this is important because in proximal (Stanford type A)
67	aortic dissections, 10-30% are not accepted for surgery, and 30-50% are technically
68	amenable for TEVAR. We describe our case series of type A aortic dissections treated
69	using TEVAR.
70	
71	METHODS: Between year 2009 and 2016, 12 patients with acute, subacute or
72	chronic type A aortic dissection with the proximal entry tear located between the
73	coronaries and brachiocephalic artery were treated with TEVAR at 3 centers. Various
74	stent-graft configurations were used to seal the proximal entry tear in the ascending
75	aorta under rapid pacing.
76	
77	RESULTS: 12 patients (9 male, 3 female), mean age 81±7 years, EuroSCORE II
78	9.1±4.5, underwent TEVAR for the treatment of type A aortic dissection. Procedural
79	success was achieved in 11/12 patients (91.7%). There was one intra-procedural death
80	and one minor stroke. No additional deaths at 30 days. At 36 months, there were 4
81	further deaths (all from non-aortic causes). The mean survival of these 4 deceased
82	was 23 months (range 15-36 months). Follow-up computed tomography demonstrated
83	favorable aortic remodeling.
84	
85	CONCLUSION: TEVAR is feasible and reveals promising early results in selected
86	patients with type A aortic dissection who are poor candidates for surgical repair. The
87	current iteration of stent-graft technology however needs to be adapted to the specific
88	features of the ascending aorta.
89	
90	Abstract word count: 248

91 Central Message

- 92 TEVAR in type A aortic dissection is feasible in selected patients. Favorable
- 93 aortic remodelling occurs in type A (and B) dissections. Thus TEVAR may be an

94 option in patients at high risk for surgery.

95

96 **Perspective Statement**

- 97 TEVAR offers a potential treatment option in a subset of patients with type A
- 98 aortic dissection at high surgical risk but with suitable anatomy. With TEVAR, 30
- day survival is >90% in these high surgical risk patients. With this proof of
- 100 concept study, broader application may be possible with further specific
- 101 technological advances.
- 102

103 Central Picture legend

- 104 Successful interventional treatment of a type A aortic dissection using TEVAR
- 105

106 **INTRODUCTION**

107

108 The surgical mortality and morbidity remains high for proximal (Stanford 109 type A) aortic dissections, particularly in the elderly with significant co-110 morbidities, despite recent strides to improve its surgical technique and 111 management (1, 2). Considering the Western demographics with increasing 112 aging population and variety of co-morbidities which portend inherent increased 113 surgical risks, the concept of endovascular stent-grafting also known as thoracic 114 endovascular aortic repair (TEVAR) (a catheter-based non-surgical technique) in 115 patients with thoracic aortic disease is increasingly attractive, propelled by the 116 desire to minimize surgical risks. TEVAR has been shown to initiate healing and 117 remodelling of the dissected aorta, by excluding and depressurizing the false lumen (3-5). To date, TEVAR strategies appear encouraging in the treatment of 118 119 various aortic pathologies (6-10). The technology has been embraced without 120 level I evidence for the treatment of distal (Stanford type B) aortic dissections, 121 and even used to treat acute proximal (Stanford type A) aortic dissections (11). 122 However, the complexity of the anatomy in the ascending aorta continues to be a 123 major obstacle for the use of endovascular technologies.

124 Acute type A aortic dissection usually requires very urgent surgical repair 125 of the ascending aorta (12-14); selected cases however may qualify for TEVAR as 126 an option in scenarios of unacceptably high surgical risk. According to the 127 International Registry of Acute Aortic Dissections (IRAD), 86% patients qualify 128 for surgical replacement of the ascending aorta, 23% or 12% require additional 129 partial or total arch replacement, respectively (15). Overall, on aggregate 91% of 130 patients in this registry underwent surgical repair under cardiopulmonary 131 bypass with 25% in-hospital mortality (15, 16). A less traumatic repair of type A 132 aortic dissection using TEVAR, where applicable, may potentially lower the 133 procedural/in-hospital mortality risk, particularly as the technology improves. 134 Surgical repair leaves a patent false lumen in both the aortic arch and

descending aorta in 75% patients, those who survive often require distal reinterventions (15, 16). One solution may be a two-stage hybrid procedure,
whereby initially, surgery is performed to replace the ascending aorta together
with aorto-brachiocephalic artery bypass without hypothermic circulatory

139 arrest. This is followed on a second occasion by surgery for left carotid artery 140 bypass and TEVAR to retrogradely place an endovascular stent-graft in the 141 thoracic aorta transfemorally in the same setting. The stent-graft excludes the 142 retrogradely perfused distal false lumen (17). The objective of such an approach 143 is to avoid surgery on the arch, and to complete the repair with an aortic stent-144 graft in a minimally invasive way. Such approach not only minimizes the 145 procedural risks, but also enables careful evaluation of the distal false lumen 146 prior to stent-graft placement. Alternatively, one-stage hybrid procedures 147 combining open (surgical) insertion of an tube-graft in the ascending aorta with head vessel transposition and antegradely placing an endovascular stent-graft 148 149 in the arch and descending aorta are feasible (18), but require the skills of both a 150 cardiac and endovascular specialist and lack the precision of the two-stage hybrid procedure (19). Moreover, there is some resistance to apply such one-151 152 stage hybrid procedures in acute type A aortic dissections, because experts are 153 aware that the fragile outer aortic wall and friable dissecting lamella are prone to 154 injury or perforation by antegrade positioning of the stent-graft under 155 conditions of circulatory arrest. At present there are no dedicated stent-grafts for 156 the ascending aorta, in particular for the repair of aortic dissections; such 157 challenges will certainly be addressed by customized stent-graft technology in 158 the near future. Nevertheless, the concept of a one-stage hybrid repair with 159 antegrade stent-graft placement may become part of a therapeutic 160 armamentarium for complex type A dissections with distal malperfusion; while a 161 multi-stage hybrid repair incorporating retrograde stent-graft placement may 162 become a preferred option in stable situations.

163 From an anatomical perspective, 30–50% patients with type A aortic 164 dissection are amenable to TEVAR (20, 21). Thus in the future more patients may be considered suitable for TEVAR with life-long follow-up. The ultimate goal is a 165 166 fully catheter-based approach to repair the ascending aorta that minimizes 167 procedural risk and initiates healing (as documented in type B dissections where 168 interventional entry closure is associated with thrombosis of the false lumen and 169 favorable aortic remodelling) (3, 4, 6, 7). Such approach is feasible with current 170 technology (22-25). Here, we describe our 12-case series of type A aortic dissections 171 treated using TEVAR.

173 **METHODS**

174

172

175 *Patient selection*

176 Between year 2009 and 2016, 12 patients with type A aortic dissection 177 consisting of an isolated dissection entry in the ascending aorta, referred to the 178 University Heart Centre (Rostock, Germany), CHU (Liege, Belgium) and Royal 179 Brompton Hospital (London, UK) were selected and subjected to TEVAR. These 180 patients were selected for TEVAR because of high co-morbidities and anatomic 181 suitability, e.g. aortic dimensions suitable to accommodate a ready-made 182 commercial stent-graft. All had elevated anesthetic risk score (American Society 183 of Anesthesiologists classification IV or greater), New York Heart Association class III or IV, chronic lung disease and/or renal impairment. Decisions regarding 184 185 treatment required consensus between cardiac surgeons and cardiologists, with 186 the patients giving informed written consent. TEVAR in this setting was 187 approved by the internal review board of each center. All patients had EKG-gated 188 computed tomography (CT) (Figure 1) and echocardiography for the diagnosis 189 and assessment of aortic dissection. Echocardiography allowed assessment of 190 the aortic valve, left ventricular function, presence/absence of tamponade, and 191 interrogation of the supra-aortic vessels.

192

193 TEVAR procedure

194 Procedural planning was based on contrast-enhanced EKG-gated CT 195 (Figure 1), which was evaluated using standard software (TeraRecon, or 196 3Mensio) to select the appropriate stent-graft size; the diameter of the stent-197 graft was chosen according to an estimate of the previous (before dissection) 198 aortic dimension to avoid oversizing. The stent-grafts used were usually ZENITH 199 TX2 (Cook, Bloomington, Indiana), GORE C-Tag (Gore Ltd. London, UK) or Relay 200 NBS (Bolton, Barcelona, Spain). They are made of a self-expanding nitinol stent 201 platform covered with polyester fabric. They are packed and mounted onto a 202 catheter-based delivery system. Figure 2 and Videos 1-3 show a typical TEVAR 203 procedure. With the patient under general anesthesia, a temporary pacing wire 204 was placed in the right ventricle and vascular access for the TEVAR device (22-

24F) obtained via right femoral arterial cut-down. The true lumen of the aorto-205 206 ilio-femoral arterial route was navigated using a soft long hydrophilic guide wire 207 (Terumo, Inc.) protruding ahead of a pigtail catheter to reach the left ventricle 208 under fluoroscopy and ultrasound guidance (transesophageal 209 echocardiography) (26). Once the pigtail is in the left ventricle, the soft 210 hydrophilic guide wire was exchanged to a stiff 270cm length guide wire through 211 the pigtail catheter. The stiff guide wire has a soft spiral tip that sits within the 212 left ventricular cavity. The stent-graft was then delivered along the stiff guide 213 wire to its intended position, where its distal landing zone is between 'distal to 214 the coronary ostia' and 'proximal to the brachiocephalic artery' in the ascending 215 aorta. In this position, the distal tip of the delivery system may cross the aortic 216 valve. For stent-graft deployment, rapid right ventricular pacing at 180 bpm was used to reduce the systolic blood pressure to \leq 50mmHg in order to avoid 217 218 displacement (windsock effect) of the stent-graft during its deployment, thus 219 enabling its precise placement. At the end of the procedure, the temporary 220 pacing wire was removed, the femoral artery access site closed, and the patient 221 extubated and transferred to the coronary care unit. Procedural success was 222 defined as successful placement of the stent-graft in its intended position with 223 sealing of the entry tear.

- 224
- 225 Follow-up

Overall aortic and true lumen diameters were assessed at the level of
sinotubular junction, ostium of the brachiocephalic artery and left subclavian
artery. Follow-up CT scans were performed approximately at 6 months and then
annually post TEVAR.

230

231 Statistical analysis

232 Descriptive statistic was used to characterize patients, procedural data233 and individual survival.

234

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235 RESULTS
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236

A total of 12 patients with proximal (type A) aortic dissection were

238 selected for TEVAR (Table 1). There were 10 DeBakey type II and 2 DeBakey 239 type I dissections. The mean age \pm standard deviation (SD) was 81 \pm 7 years; 240 male:female ratio was 9:3. All patients were of advanced age with chronic lung 241 disease, coronary artery disease and/or renal impairment. The mean EuroSCORE 242 II was 9.1 ± 4.5 (SD). The median time from onset of symptoms/diagnosis to 243 TEVAR was 24 days. There were 6 cases of acute (≤ 14 days after symptom onset) 244 and 6 cases of subacute (15 days to 3 months) or chronic (>3 months) aortic 245 dissections. The false lumen of the dissection expanded significantly causing 246 various complications, including dyspnoea, hoarseness or laryngeal nerve dysfunction. There was a history of chest and back pain in all cases. There were 247 no significant aortic insufficiency, no clinically apparent distal malperfusion 248 249 syndromes and no distal interventions required.

250 Procedural success was 91.7% (11/12) (Table 1). There was one death 251 due to cardiac tamponade from wire induced perforation of the left ventricle. All 252 remaining 11 patients were discharged alive within 2 weeks of TEVAR. The 253 mean procedural time was 86 ± 33 (SD) minutes. Stent-grafts were deployed 254 under rapid right ventricular pacing which achieved a mean systolic pressure of 255 34 ± 15 (SD) mmHg. The mean follow-up time was 21.1 ± 11.8 (SD) months 256 (range 0 - 36 months) post TEVAR. There were 4 deaths, one each at 15, 19, 23 257 and 36 months (Table 1). All appeared to have died from natural causes. The 258 mean survival in those who died during follow-up was 23 months.

Follow-up CT scans revealed thrombosis or remodelling of the stent-graft excluded false lumen. The diameter of the aorta at the sinotubular junction was not enlarged and remained similar to the normal aorta post TEVAR (Figure 3).

262

263 **DISCUSSION**

264

There is general consensus that proximal aortic dissection or any major
pathology involving the ascending aorta should be subjected to surgical repair.
However, 10-30% of patients with acute type A aortic dissection are considered
too high risk for surgical repair and would therefore receive only medical
therapy with associated high mortality of ≈60% in the intermediate term (13, 15,
27, 28). Surgical mortality is 10-25% (16, 27) depending on the complexity of the

operation and the clinical status of the patient. In our hands, the procedural
mortality of TEVAR was 8%, which compares favorably with the published early
endovascular mortality of 11% [18, 19]. Interestingly, the most anticipated
complications such as major stroke did not occur. One minor stroke (transient)
and one death from guidewire perforation of the left ventricle leading to fatal
tamponade occurred.

We estimate the number of TEVAR procedures performed in this study relative to all emergency surgeries for type A aortic dissection in our 3 centers to be $\approx 2\%$, or 6-17% of inoperable type A cases (assuming the incidence of type A dissection ≈ 40 cases/year for a typical aortic center; therefore across 3 centers spanning the study period of 6 years, the total number of cases = 40 cases x 6 years x 3 centers = 720 cases; generally 10-30% of type A dissections are inoperable (15)).

284 Successful sealing of the false lumen entry with no development of 285 proximal type I endoleak were achieved. During follow-up, no cases of endoleak 286 were identified which is encouraging and different to $\approx 10\%$ incidence reported 287 elsewhere (29). In selecting the appropriate size of stent-grafts, we chose the 288 diameter of the stent-grafts according to an estimate of the previous (before 289 dissection) aortic dimension to avoid oversizing. The goal was to re-shape the 290 dissected ascending aorta, cover the entry tear and depressurize the false lumen 291 (30, 31); there is a fine balance between fixation to the aortic wall and the degree 292 of intimal injury caused by the self-expanding stent-graft. However, once 293 precisely deployed, the process of remodelling of the false lumen appears similar 294 between the proximal and distal dissection, and takes place usually within one 295 year, similar to that reported elsewhere (32, 33). Most of our cases were 296 DeBakey type II dissections, and even in the 2 cases of type I dissection (patients 297 2 and 3 in Table 1), favorable aortic remodelling of the descending aorta were 298 observed. It seems that the therapeutic concept of closing the entry and 299 depressurizing the false lumen in type B (distal) aortic dissection holds true also 300 in type A (ascending) aortic dissection (3, 4). As long as the false lumen is 301 thrombosed and depressurized, survival even with type A aortic dissection can 302 be improved by TEVAR (34). In addition, the patients' exposure to unacceptable 303 risk of surgery is minimized.

304 Current literature (mostly single or small case series) underlines the 305 feasibility of proximal endovascular procedures over more than 10 years, 306 performed by surgeons and interventionalists (Table 2). Our series over 6 years 307 with a mean follow-up of >20 months (range up to 3 years) underlines the fact 308 that in the setting of significant co-morbidities representing high surgical risk 309 but with suitable anatomy, TEVAR can be a viable alternative to surgical repair. 310 In other words, it is feasible to avoid high risk/ complex surgery, and apply a less 311 traumatic intervention to obtain a similar or better short-term outcome in a 312 subset of elderly patients with significant co-morbidities. The advantages of 313 TEVAR includes the avoidance of thoracotomy, cardiopulmonary bypass, 314 selective head perfusion and associated surgical risks in an elderly population, 315 often in a critical condition (35). If the high initial $\approx 60\%$ mortality of type A 316 aortic dissection can be successfully lowered by TEVAR, such less traumatic 317 strategy may potentially become an option in a broader spectrum of patients 318 (27). As TEVAR is a fairly expensive procedure, its associated lower 319 risks/complications and shorter lengths of hospital stay compared to surgery 320 may potentially demonstrate its advantages in terms of cost and clinical outcome 321 over surgery.

322 TEVAR will not be feasible in every patient; the most suitable anatomy is 323 where the entry tear of the dissection is located in the middle portion of the 324 ascending aorta. Entry tears close to the coronaries or aortic valve lack a suitable 325 length of landing zone. Entry tears close to the brachiocephalic artery would 326 require complex branching/fenestration strategies. Currently, only a limited 327 number of choices are available regarding the type of stent-graft and delivery 328 system because relatively large diameter and short length stent-grafts are 329 required. Existing delivery systems need to be modified for ascending aorta 330 intervention: a long nosecone can either damage the aortic valve or increase the 331 chance of left ventricular perforation by the stiff guidewire, as occurred in one of 332 our patients.

On a technical note, with the use of rapid ventricular pacing no
misplacement of these short stent-grafts was seen. Pacing is probably the most
efficient method to avoid windsock effect of the left ventricle, enabling precise
stent-graft placement. Transoesophageal echocardiography is also useful in

337 guiding stent- graft positioning and assessing sealing of the entry tear (26). 338 It should be emphasized that a multidisciplinary team, consisting of 339 cardiac and vascular surgeons and cardiologists, should select suitable patients 340 for the procedure, similar to transcatheter aortic valve replacement (TAVR). 341 Looking forward, we believe TEVAR in the ascending aorta is a definitive 342 solution for patients not accepted for surgery, or a bridging solution in case of 343 unclear neurological diagnosis (e.g., major stroke) to buy time for reconstructive 344 surgery. Selection process in patients not suitable for surgical valve replacement 345 may even be conceivable for combined TAVR- TEVAR technology, in an attempt to treat variants of aortic dissection including those with compromised aortic 346 347 valve function. It should also be emphasised that in the acute setting there is a 348 need to identify and transfer type A aortic dissection patients to a specialized 349 unit as quickly as possible; once the dissection has produced coronary 350 obstruction (usually the right coronary artery) with ensuing (right) ventricular 351 infarction and heart failure, it is a difficult situation to retrieve by either 352 endografting or conventional surgery.

353

354 Study limitations

355 While we could demonstrate proof of concept and feasibility of TEVAR in 356 the ascending aorta with encouraging results, we did not examine possible 357 detrimental effects such as stiffening of the ascending aorta by the stent, lowered 358 vascular compliance, negative effects on the aortic valve function or 359 hypertension in this observational study. Our sample size is relatively small, but it 360 represents one of the biggest case-series in the field and supports the feasibility of 361 TEVAR in practical terms. We have no control group (medically treated or surgery) 362 for comparison, but historical data suggests that surgery confers 25% peri-operative 363 mortality, and medical treatment is associated with 60% mortality (13, 15, 27, 28). A 364 propensity-matched comparison prior to any randomized study would probably 365 be the next step to strengthen the data on TEVAR in the proximal aorta; current 366 technology is unlikely to allow a broader application yet.

367

368 Conclusion

369

TEVAR is feasible and reveals promising early results in selected patients

- 370 with proximal (type A) aortic dissection who are poor candidates for surgical repair.
- The current iteration of stent-graft technology however needs to be adapted to the
- 372 specific features of the ascending aorta before TEVAR as a concept emerges for
- broader applications in the proximal aorta.
- 374

375 **Figure legends**

376

377 Figure 1. From top to bottom: 2-dimensional transverse and coronal section of a 378 localized proximal aortic dissection (type A) with a large entry between the 379 aortic valve and brachiocephalic artery; 3-dimensional reconstruction prior to 380 treatment. 381 382 Figure 2. TEVAR procedural sequence for placing a covered stent-graft to treat a 383 type A aortic dissection. (A) Aortogram and set-up showing a right ventricular 384 pacing wire and transesophageal echo probe; (B) Stent-graft deployment during 385 rapid pacing; (C) Completed deployment of the stent-graft; (D) Aortogram 386 demonstrating procedural success. 387 388 Figure 3. From top to bottom: 2-dimensional transverse and coronal sections of a proximal (type A) aortic dissection before (left) and after stent-grafting (right), 389 390 demonstrating TEVAR reconstruction and remodelling of the aorta. The bottom 391 panels demonstrate the successful intervention in 3-dimensional reconstruction. 392 Video 1. Before TEVAR. Digital subtraction angiogram showing a large tear in the 393 394 ascending aorta, and a marker pigtail, pacing wire and transoesophageal 395 echocardiography probe in place. 396 397 Video 2. During TEVAR. Fluoroscopic display of the launch of a self-expanding 398 Viabahn stent-graft in the brachiocephalic artery followed by a self-expanding C-399 Tag stent-graft covering the ascending aorta under rapid pacing. 400 Video 3. After TEVAR. Completion angiogram after placement of the Viabahn 401 402 stent-graft in the brachiocephalic artery and C-Tag stent-graft in the ascending 403 aorta; the entry to the dissection is sealed and flow is preserved to the 404 brachiocephalic and coronary arteries.

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Patient	Age & Sex	Diagnosis	Euroscore II	SG	SG size DxL (mm)	Procedure duration (min)	Procedural complications	Follow-up (months)	
1	74 M	cTAAD (DeBakey II)	6.9	Cook	34x77	90	None	32	
2	75 M	aTAAD (DeBakey I)	8.1	Bolton NBS	34x60	140	None	29	
3	87 M	aTAAD (DeBakey I)	13.4	Bolton NBS	34x60	79	None	35	
4	89 M	cTAAD (DeBakey II)	15.0	Cook	36x77	149	None	15,†	
5	90 M	cTAAD (DeBakey II)	19.3	Cook	36x77	70	Ventricular rupture, tamponade	0,†	
6	69 M	aTAAD (DeBakey II)	3.9	Cook	34x77	61	None	36, †	
7	75 M	cTAAD (DeBakey II)	4.9	Bolton NBS	34x60	70	None	24	
8	87 M	cTAAD (DeBakey II)	9.4	Cook	36x77	49	Minor Stroke	15	
9	87 F	aTAAD (DeBakey II, post TAVR)	7	Optimed	32x50	120	None	23, †	
10	83 M	cTAAD (DeBakey II)	6.9	Cook	34x77	89	None	19, †	
11	75 F	aTAAD (DeBakey II)	5.9	Cook	34x77	60	none	5	
12	75 F	aTAAD (DeBakey II)	8.9	Gore + Viabahn in innominate artery	34x100	60	none	0	

Abbreviations: aTAAD (acute type A aortic dissection), cTAAD (subacute or chronic type A aortic dissection), D (diameter), F (female), L (length), M (male), NBS (non-bare stent), SG (stent-graft), TAVR (transcatheter aortic valve replacement). † indicates deceased.

Re-intervention	ı	ı	I	Open surgery	I	-	-	Balloon dilatation for endoleak	1	I	I	ı	ı	-	-	Angioplasty and stenting of left renal artery	-	-	Re-stenting for endoleak	Stent extension, open surgery
Complications	I	ı	I	Aortic regurgitation	I	Endoleak	-	ı	1	Tachycardia	I	Aortic regurgitation	I	-	-	Arrhythmia		-	Endoleak Stroke	Contained aortic rupture, endoleak, stent migration,
Cause of death	Cardiac arrest	1	ı	ı	ı	-	Cardiac arrest stent migration	Gastrointestinal bleeding	-	-	-	ı	-	-		·	1	-	ı	Coronary obstruction, aortic rupture, tamponade
earry mortality n (%)	1 (100)	0	0	0	0	0	1 (100)	1 (10)	0	0	0	0	0	0	0	0	0	0	0	3 (17)
Follow-up months	1	12	none	21	1	none	none	35.5	none	32	15 (4-39)	1	none	9	none	22 (12-31)	none	none	33 (3-57)	12 median
Access route	RFV Trans-septal	Aorta	I	ТЕ	ΤF	TF	TF	2 LCCA, 8 TF	TF	Aorta	ТF	ТА	ТА	TA	TA	ı	RCA	TF	4 TA, 1 TF, 1 LCCA	5 ТА, 4 ТАх, 9 ТF
Stent-graft	Lecteba	Covered Z-stent	Gore	Gianturco Z-stent	Jotec	Medtronic	Braile Biomed	Various	Medtronic	Cook	Cook	Cook	Cook	Cook	Jotec	Cook	Gore	Medtronic	Cook Amplatzer	Cook, Gore, Medtronic
Acute n (%)	1 (100)	1 (100)	1 (100)	0	1 (100)	1 (100)	0	6 (60)	1 (100)	1 (100)	4 (100)	0	1 (100)	1 (100)	1 (100)	5 (33)	1 (100)	1 (100)	3 (50)	9 (50)
z	1	1	1	1	T	1	1	10	1	1	4	1	1	1	1	15	1	1	6	18
Year	2000	2003	2004	2004	2006	2007	2007	2011	2012	2012	2013	2013	2013	2013	2013	2013	2014	2014	2015	2015
Author	Dorros et al	Wang et al	lhnken et al	Zhang et al	Zimpfer et al	Senay et al	Palma et al	Ye et al	Metcalfe e al	Gustavo et al	Ronchey et al	Bahaeddin et al	Eric et al	Kölbel et al	Frederic et al	Lu et al	Yuuya et al	Kimberly et al	Vallabhajosyula et al	Roselli et al

Table 2

Abbreviations: LCCA (left common carotid artery), N (number), PA (pseudoaneurysm), RCA (right carotid artery), RFV (right femoral vein), TA (transapical), TAx (transaxillary), TF (transfemoral).

























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