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1 Comparison of fenestrated stentgrafts and open repair  
2 for juxtarenal aortic aneurysms using a propensity score  
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4 Alia Bizos<sup>a</sup>, Aurélien Hostalrich<sup>a</sup>, Xavier Chaufour<sup>a</sup>, Pascal Desgranges<sup>b</sup>, Richard Layese<sup>b</sup>,  
5 Frédéric Cochenec<sup>b</sup>, Florence Canoui-Poitrine<sup>b</sup>

6 <sup>a</sup> Department of Vascular Surgery, Rangueil University Hospital, Toulouse, France

7 <sup>b</sup> Department of Vascular Surgery, Henri-Mondor University Hospital (Assistance Publique-  
8 Hopitaux de Paris – APHP), Créteil, France

9

10 ***Corresponding author.*** Alia Bizos

- 11 • [bizos.a@chu-toulouse.fr](mailto:bizos.a@chu-toulouse.fr)  
12 • Service de chirurgie vasculaire, CHU Rangueil, 1 avenue du Professeur Jean Poulhès, 31400,  
13 Toulouse  
14 • Home telephone number : +33687217991  
15 • Business telephone number : +33561323907  
16 • Fax number : +33561323917

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21

22

## 23 ABSTRACT

24 **Objectives:** The purpose of this study was to compare post-operative morbi-mortality and  
25 medium-term follow-up of fenestrated stentgrafting (FEVAR) and open repair (OR) for  
26 patients with juxtarenal aortic aneurysms (JRAAs).

27 **Materials and Methods:** All consecutive patients who underwent custom-made FEVAR or  
28 open repair for complex abdominal aortic aneurysm (AAA) between 2005 and 2017 in two  
29 tertiary centers were scrutinized. Patients with JRAA constituted the study group. Suprarenal  
30 and thoracoabdominal aortic aneurysms were excluded. The groups were made comparable  
31 through the use of a propensity score matching.

32 **Results:** 277 patients with JRAAs were included, 102 (36.8%) in the FEVAR group and 175  
33 (63.2%) in the OR group respectively. After propensity score matching, 54 FEVAR patients  
34 (52.9%) and 103 OR patients (58.9%) were included for analysis. In-hospital mortality rates  
35 were 1.9% (n=1) in the FEVAR group vs. 6.9% (n=7) in the OR group ( $p=.483$ ).  
36 Postoperative complications were less common in the FEVAR group (14.8% vs. 30.7%;  
37  $p=.033$ ). Mean follow-up was 42.1 months in the FEVAR group and 40 months in the OR  
38 group. Overall mortality rates at 12 and 36 months were 11.5% and 24.5% in the FEVAR  
39 group vs. 9.1 % ( $p=.691$ ) and 11.6% ( $p=.067$ ) in the OR group. Late reinterventions were  
40 more frequent in the FEVAR group (11.3% vs. 2.9%;  $p=.047$ ). However, freedom from  
41 reintervention rates were not significantly different at 12 months (FEVAR: 86% vs. OR: 90%;  
42  $p=.560$ ) and 36 months (FEVAR: 86% vs. OR: 88.4%,  $p=.690$ ). In the FEVAR group,  
43 persistent endoleak during follow-up was identified in 11.3% of cases.

44 **Conclusion:** In the present study, there was no statistical difference in terms of mortality, in-  
45 hospital, at 12 or 36 months, between FEVAR and OR groups for JRAA. FEVAR for JRAA  
46 was associated with a significant reduction of overall postoperative major complications  
47 compared with OR. There were significantly more late reinterventions in the FEVAR group.

## 48 INTRODUCTION

49 Fenestrated endovascular aortic repair (FEVAR) is currently the most commonly used  
50 alternative to open repair (OR) to treat complex abdominal aortic aneurysms (AAA). Recent  
51 systematic reviews derived from high volume centers series have provided excellent short-  
52 term and encouraging mid-term results(1)(2)(3). Long-term results remain poorly known.  
53 Expert centers have reported low rates of aneurysm-related mortality during follow-up, at the  
54 cost of substantial reintervention rates, mainly for endoleaks or branch instability(4). Data  
55 comparing FEVAR and OR are scarce. In the absence of randomized study, they are based on  
56 few meta-analyses(1)(2) and retrospective studies(5)(6)(7) leading to conflicting  
57 conclusions. These studies included heterogeneous patients with heterogeneous anatomies.  
58 There is a need for subgroup analyses reporting specific results for juxtarenal (JRAA) and  
59 suprarenal (SRAA) aortic aneurysms. Although these two entities are often mixed and  
60 reported as “pararenal AAA” or “complex AAA”, SRAA have been associated with increased  
61 mortality and morbidity, especially when treated by OR(8)(9)(10). OR for JRAA has shown  
62 acceptable operative risk with durable results in terms of both graft integrity and preservation  
63 of renal function(11).

64 The purpose of this study was to compare postoperative morbi-mortality, the short-term,  
65 defined as in-hospital, and mid-term, defined as after hospital stay, results of FEVAR and OR  
66 for JRAAs in order to provide the best patient-specific therapy option.

## 67 METHODS

### 68 **Definitions**

69 In this study, juxtarenal and suprarenal aortic aneurysms were defined according criteria used  
70 for open surgery.

71 Juxtarenal AAAs were defined as requiring suprarenal, supramesenteric or supraceliac aortic  
72 cross clamping and infrarenal proximal aortic suture without separate renal revascularisation.

73 Suprarenal AAA were defined as requiring suprarenal, supramesenteric or supraceliac aortic  
74 cross clamping and a suprarenal proximal aortic suture and/or a bevelled proximal aortic  
75 suture encompassing at least one renal artery and/or a separate renal or splanchnic  
76 revascularization.

### 77 **Study population**

78 This retrospective study was conducted in two tertiary centers (Rangueil Hospital, Toulouse,  
79 France and Henri Mondor Hospital, Créteil, France) between January 2005 and December  
80 2017. Both centers frequently perform both FEVAR and OR of complex AAA. Indications  
81 for FEVAR or OR were discussed during multidisciplinary meetings and based on clinical  
82 condition, comorbidities, anatomic criteria and patients' preferences.

83 All consecutive patients who underwent FEVAR or open repair for complex AAA between  
84 2005 and 2017 were scrutinized. Only patients with JRAA treated by custom-made FEVAR  
85 and OR were selected.

86 Selection of JRAA treated by FEVAR was based on preoperative computed tomography  
87 angiography (CTA) analysis. CTA of patients with complex AAA who underwent FEVAR  
88 were reviewed by two senior surgeons (XC, FC) having an expertise in both endovascular and  
89 open repairs. Patients deemed to fulfil the anatomic criteria of JRAA were included. Patients  
90 who were thought to fulfil the definition of SRAA were excluded. In case of disagreement,  
91 decision to include patients was made after discussion between the two senior surgeons.

92 For selection of JRAA treated by OR, operating reports were reviewed. Patients fulfilling the  
93 definition of JRAA were included.

94 Suprarenal or thoracoabdominal aneurysms, infrarenal AAA requiring renal artery  
95 reconstruction for occlusive lesions, inflammatory or infectious aneurysms, symptomatic or  
96 ruptured aneurysms were excluded.

97 The follow-up was the usual protocol of our institutions. In the FEVAR group, it consisted in  
98 CTA within one month, at six months and every year. In the OR group, it consisted in CTA or  
99 duplex ultrasound (DU) at one month, DU at six months and every year. A CTA was  
100 performed if abnormalities were detected on DU. The 1 and 6-month control is to confirm the  
101 technical success of the surgery, to look for an eventual technical complication and to verify  
102 the good healing of the patient. The 2019 ESVS clinical practice guidelines(12) recommend  
103 every 5-year surveillance to detect para-anastomotic aneurysm, recurrent aortic aneurysm and  
104 peripheral aneurysm. This recommendation is based on an old study from 1970 to 1976  
105 published in 1985. We believe a more robust surveillance is needed to ensure the absence of  
106 complications and to provide regular follow-up for our patients with yearly DU, with yearly  
107 DU, which is non-invasive, non-irradiating and inexpensive. In Henri Mondor Hospital,  
108 preoperative, intraoperative and postoperative data were collected prospectively in a  
109 dedicated software (Safir, Opale, Paris). In Ranguel Hospital, data were collected  
110 retrospectively from patients' files. For survival assessment, telephone interviews were  
111 conducted.

112 This study was declared compliant with the reference methodology provided by the CNIL  
113 (Commission Nationale de l'Informatique et des Libertés), declaration number: 2209803 v 0.  
114 Individual consent for this retrospective analysis was not required by the ethics committee.  
115 Risks and benefits of FEVAR and OR were explained to all patients before they gave their  
116 consent to the procedure.

## 117 **Operative techniques**

### 118 Fenestrated Endovascular Aneurysm Repair

119 Patients were treated using various but only custom-made stent grafts (Zenith (Cook Medical,  
120 Bloomington, Ind), Ventana (Endologix, Inc, Irvine, CA, USA) or Anaconda (Vascutek,  
121 Inchinnan, UK), depending on availability, planning centers recommendations and surgeon's  
122 preference. In one centre, most of procedures were performed in a dedicated hybrid room  
123 using fusion techniques whereas in the other centre, most of patients were treated in an  
124 operating theatre equipped with a C-arm. Femoral access was obtained via a percutaneous  
125 approach or a surgical cutdown, depending on surgeon's preference. During the study period,  
126 cone beam CT scans were not performed at the end of the procedure. Technical success was  
127 defined according to the Society of Vascular Surgery (SVS) criteria(13) as successful  
128 insertion and deployment of the device in the absence of surgical conversion, death, type I or  
129 III endoleaks, obstruction of graft limbs or target vessels within the first 24 hours after  
130 surgery.

### 131 Open Repair

132 Surgical approach was a left lumbotomy or a median laparotomy. The level of aortic cross  
133 clamping depended on anatomic characteristics and surgeon's preferences. A suprarenal  
134 aortic cross clamping was performed whenever possible. In some patients with thrombus at  
135 the visceral level or heavily calcified visceral aortas, a supramesenteric or a supraceliac aortic  
136 cross clamping was preferred. All proximal anastomoses were performed below the renal  
137 arteries in an end-to-end fashion. Distal reconstructions were performed according to  
138 standards in vascular surgery. Technical success was achieved when the aortic reconstruction  
139 was patent and patients were alive within the first 24 hours.

### 140 **Endpoints**

141 The primary endpoint was overall survival. Secondary endpoints included moderate to severe  
142 complications occurring during hospital stay, length of hospital stay, reinterventions and  
143 target vessel patency. In the FEVAR group, endoleaks and aortic rupture were also reported.

144 Acute kidney injury (AKI) was defined as an increase in serum creatinine to  $\geq 1.5$  times the  
145 baseline level during the first 7 days, as defined by the clinical practice guidelines for acute  
146 kidney injury of the KDIGO (Kidney Disease Improving Global Outcomes) program(14). The  
147 KDIGO classification combines the RIFLE and AKIN classifications(15). Severe AKI is  
148 defined as stages 2 and 3 of the KDIGO classification. Since mild AKI is regressive and do  
149 not lead to complications(11), that was the case in our population. Only moderate to severe  
150 grade complications, as defined by the SVS criteria(13), were considered for analysis.  
151 Moderate complication indicates the need for significant intervention, prolongation of  
152 hospitalization more than 24 hours, and at most, minor permanent disability that does not  
153 preclude normal daily activity. A severe complication necessitates major surgical or medical  
154 intervention, may be associated with prolonged convalescence, is usually accompanied by  
155 prolonged or permanent disability, and may result in death. This was determined by analyzing  
156 all patients' complications, then classified them in minor and major (moderate to severe)  
157 complications (**Table 1**). These included severe AKI.

158 The limit for early or late reinterventions was the date of discharge from hospital.

### 159 **Statistical analysis**

160 Patient characteristics were expressed in n (%) and mean (standard deviation) or median (Q1-  
161 Q3) according to the variable (categorical or continuous).

162 A propensity score was built by a logistic regression model to account for the indication bias.

163 Explanatory variable was surgical strategy (FEVAR vs. OR). Independant variables were age,  
164 sex, smoking, diabetes, obesity, coronary artery disease, heart/renal/respiratory failure,  
165 American Society of Anesthesiologists (ASA) score, aortic surgery history, aneurysm  
166 diameter and hospital centre. Characteristics with a p value  $<0.20$  in univariate analysis  
167 (Table 2) were included in the model to build the propensity score.



168 One patient in the FEVAR group was matched for 1 to 4 patients in the OR group based on  
169 the value of the propensity score.

170 Characteristics of patients in the unmatched and matched OR groups were compared to  
171 FEVAR. For an unmatched comparison, exact Pearson or Fischer tests and T-test or  
172 Wilcoxon test were used according to the nature and distribution of the variables. The  
173 comparison of the matched groups was done using a mixed logistic regression model that  
174 considered the paired nature of the data.

175 The analysis of the outcomes was conducted on paired groups. A descriptive analysis was  
176 carried out. Survival analysis was performed using Kaplan-Meier survival curves for death  
177 and reintervention outcomes. For the latter outcomes, a mixed proportional Hazard Cox  
178 Model was used to compare the FEVAR and OR groups. For other outcomes, a mixed logistic  
179 regression model was used.

180 Missing data on outcomes were not imputed. Missing data are displayed in each table.

181 Two-tailed tests were used. A 0.05 threshold was used for significance. The analysis was  
182 done using Stata SE v15.0 (College Station, TX, USA).

## 183 RESULTS

### 184 Demographics

185 Between January 1<sup>st</sup> 2005 and December 31th 2017, 548 patients were treated for complex  
186 AAA of which 277 were treated for asymptomatic JRAA, either by FEVAR (n=102) or OR  
187 (n=175) (**Figure 1**). Before propensity score matching, ten baseline characteristics were  
188 significantly different between the two groups (**Table 2**).

189 After propensity score matching, 157 patients were included in the analysis. There were no  
190 significant differences in baseline characteristics between the two groups (**Table 2**).

### 191 Intraoperative data

192 Technical success (FEVAR: 94.4%, OR: 100%;  $p=.992$ ) and median procedure length  
193 (FEVAR: 180 min, OR: 185 min;  $p=.076$ ) were no statistically different.

194 Intraoperative details of matched patients who underwent FEVAR are given in **Table 3**.

195 There was no open conversion. There were 139 fenestrations (average of 2.6 fenestrations per  
196 patients) and 35 unstented scallops.

197 Intraoperative data of matched patients who underwent OR are given in **Table 4**.

### 198 **Early post-operative results**

199 Group comparison for early postoperative outcomes are detailed in **Table 5**.

200 There were four 30-day mortality events in the OR group, one in the FEVAR group ( $p=.510$ ).

201 In-hospital mortality rates were similar in both groups (FEVAR: 1.9%; OR: 6.9%;  $p=.483$ ). In  
202 the FEVAR group, one patient died from pneumonia. In the OR group, one patient died  
203 during the procedure from haemorrhagic shock, four patients died from multi-visceral failure,  
204 one patient died from colic ischemia. For another patient, cause of death remained unclear.

205 The overall complication rate was significantly lower in the FEVAR group ( $p=.033$ ).

206 The AKI and severe AKI rates were similar in both groups ( $p=.232$  and  $p=.09$ , respectively).

207 Twelve patients (FEVAR group:  $n=2$ ; OR group:  $n=10$ ) required transient dialysis after the  
208 procedure during the hospital stay, with no statistical difference between groups. No patient  
209 required permanent dialysis.

210 Two FEVAR patients occluded a renal artery stent during the postoperative course. One  
211 patient had a control CT scan eleven days after the surgery, that made discover the renal stent  
212 occlusion, he was asymptomatic so no salvage was attempted. For the second one we have  
213 unfortunately no data about how it was discovered or why was there no salvage attempted.

214 One of them had a persistent stage 1 AKI at discharge, the other one required temporary  
215 dialysis and was discharge with persistent stage 2 AKI.

216 Two early-reinterventions for two FEVAR patients and twelve early-reinterventions for eight  
217 OR patients ( $p=.345$ ) were performed.

218 In the FEVAR group, both patients required an early-reintervention for an acute limb  
219 ischemia. There was no significant statistical difference in terms of in-hospital mortality  
220 ( $p=.883$ ), overall complication ( $p=.203$ ), early reintervention ( $p>.99$ ) and endoleak at  
221 discharge ( $p>.99$ ) between the different type of endografts used (Supplemental files: **Table 7**).

222 In the OR group, reinterventions consisted in: haemostasis for haemorrhage ( $n=4$ ), acute limb  
223 ischemia ( $n=4$ ) and bowel resection ( $n=4$ ). In one of the patients reoperated for acute limb  
224 ischemia, a right ilio-renal bypass was performed during the same procedure, we do not have  
225 any information about why this bypass was performed.

## 226 **Midterm results**

227 Mid-term results are detailed in **Table 6**.

228 There was no statistical difference about overall survival rates in the OR group at 12 months  
229 (FEVAR: 88.5%, 95% confidence interval (CI): 76.2-94.7; OR: 90.9%; 95%CI: 83.2-95.2;  
230  $p=.691$ ) and 36 months (FEVAR: 75.5%, 95%CI: 60.7-85.3; OR: 88.4%, 95%CI: 79.9-93.4;  
231  $p=.067$ ) (**Figure 2**).

232 During follow-up, eighteen deaths occurred in the FEVAR group from cancer ( $n=8$ ), stent  
233 graft infection ( $n=2$ ), cardiac disease ( $n=1$ ), respiratory failure ( $n=1$ ), stroke ( $n=1$ ) and  
234 unknown cause ( $n=5$ ). Twenty deaths occurred in the OR group from unknown cause ( $n=9$ ),  
235 cancer ( $n=4$ ), stroke ( $n=1$ ), end-stage Parkinson's disease ( $n=1$ ), aortic arch rupture ( $n=1$ ),  
236 secondary to heart surgery ( $n=1$ ), endocarditis ( $n=1$ ) and critical ischemia of the lower limbs  
237 ( $n=1$ ).

238 No statistical difference in terms of renal function decline could be identified (**Table 6**).

239 Freedom from reintervention rates were not significantly different at 12 months (FEVAR:  
240 86%, 95% CI:72.8-93.1; OR: 90%, 95% CI: 82.1-94.5;  $p=.560$ ) and 36 months (FEVAR:  
241 86%, 95% CI: 72.8-93.1; OR: 88.4%, 95% CI: 79.7-93.5;  $p=.690$ ) (**Figure 3**).

242 Apart from early reinterventions mentioned above, ten late reinterventions were performed in  
243 six patients of the FEVAR group and three in three patients of the OR group.

244 In the FEVAR group, late reinterventions consisted in endovascular treatment (n=2) or open  
245 ligation of the inferior mesenteric artery (n=1) for type II endoleaks, drainage of access site  
246 complications (n=4), bypasses for graft limb thrombosis (n=2) and peripheral stenting for  
247 acute limb ischemia (n=1). There was no significant statistical difference in terms of late  
248 reintervention between the different type of endografts used ( $p=.645$ ) (Supplemental files:  
249 **Table 8**).

250 In the OR group, two late endovascular reinterventions were required, one to treat a growing  
251 iliac aneurysm due to a dissection on the recipient artery from the suture line and the other to  
252 treat an ectatic primary iliac artery responsible of distal embolisms. One late open  
253 reintervention was required to treat a false distal aneurysm.

254 In the matched FEVAR group, no aortic rupture occurred during follow-up. However, two  
255 aortic ruptures were recorded in the unmatched FEVAR group. Six patients presented with  
256 persistent endoleak (n= 2 type Ia, n= 4 type II). Two of them required reintervention, as  
257 mentioned above. Two patients with type Ia endoleaks died during follow-up, one from a  
258 stent graft infection, the other one from unknown cause after he was lost from follow-up.

259 In the OR group, no renal artery occlusion was reported but most of DU did not focus on  
260 renal artery patency.

## 261 DISCUSSION

262 In this retrospective comparative cohort study with propensity score matching, there was no  
263 statistical difference in terms of mortality, between FEVAR and OR groups for JRAA,

264 although a trend towards higher mortality rates was observed at 36 months in the FEVAR  
265 group. FEVAR for JRAA was associated with a significant reduction of all major  
266 postoperative complications compared with OR, but no significant difference was found when  
267 considering individual complications. Only the length of stay was statistically different, which  
268 was longer in the OR group ( $p=.012$ ). There was no statistical difference between groups in  
269 terms of early-reintervention rates, but significantly more late-reinterventions were observed  
270 in the FEVAR group ( $p=.047$ ). Eleven-point three percentage of the FEVAR patients had a  
271 persistent endoleak during the follow-up.

272 To the best of our knowledge, it is the first study to compare FEVAR and OR specifically for  
273 juxtarenal AAA, that is suprarenal and type IV TAAAs excluded. Previous comparative  
274 studies have reported on so called “complex” AAA or “pararenal” AAA, which typically  
275 refers to a mix population of JRAA, SRAA, type IV thoracoabdominal aortic aneurysms and  
276 sometimes infrarenal AAA with occlusive lesions of renal arteries. Without subgroup  
277 analysis, results of these studies can hardly be extrapolated to JRAA. Recent data showed that  
278 OR of JRAA provides excellent short-term and long-term results, with low rates of graft-  
279 related complications during follow-up(11)(16). In contrast, there is a reasonable amount of  
280 data in the literature suggesting that OR of SRAA is associated with increased risks of  
281 mortality and morbidity compared to OR of JRAA(8)(9)(10). When treated by FEVAR,  
282 differences between JRAA and SRAA in terms of postoperative outcomes are not so clear.  
283 There is a need for comparative studies with a standardized definition of JRAA and SRAA.

284 So far, no randomized study comparing FEVAR and OR for juxta and suprarenal AAA has  
285 been published. It is unlikely that such a trial will be conducted in the coming years since  
286 practice in each centre is based on surgeon skills or preferences, and intensive care unit  
287 expertise. Most of consistent comparative data in the literature comes from meta-  
288 analysis(1)(2)(17)(18), registries(5)(19)(20)(21)(22) and retrospective studies with

289 propensity score matching(6)(7). Again, these studies included heterogeneous patients with  
290 heterogenous anatomies. They provided conflicting results in terms of early mortality. Some  
291 studies reported similar postoperative mortality rates(1)(17)(5)(7) which is consistent with  
292 our results. Others reports suggested that FEVAR is associated with reduced postoperative  
293 mortality rates(2)(18)(19)(20)(22)(6) especially in octogenarians(21). However, almost all  
294 comparative studies reported a reduced complication rate in the FEVAR group, which is in  
295 line with our finding.

296 Comparative data on mid-term results remain scarce in the literature. In the present study,  
297 late-reintervention rates were significantly higher in the FEVAR group. Indeed, with  
298 endovascular treatment of AAA, the disease is not cured but only contained. The tissue  
299 evolves and the aorta may expand, leading to endoleaks and therefore reinterventions. Stent  
300 patency as well as acute limb ischemia due to EVAR/FEVAR embolization may also be a  
301 cause of reinterventions. These results are consistent to those of the most recent meta-analysis  
302 comparing FEVAR and OR for juxta and suprarenal AAA. At a mean follow-up of 31  
303 months, estimated survival was similar for FEVAR and OR but the rate of late reintervention  
304 following FEVAR was higher(1).

305 There is also a lack of comparative data on renal function decline during follow-up. To our  
306 knowledge, only one retrospective study with a propensity-matched comparison found no  
307 difference in terms of renal function decline(7). The contribution of our study to provide more  
308 data on that matter is poor. We did not find any significant difference in terms of renal  
309 function decline but creatine levels at last follow-up were only available in half of the  
310 patients.

311 Our study has the advantage of providing specific data on juxtarenal AAA but several  
312 limitations are worth mentioning. It is a non-randomised study and matching ended up with  
313 limited numbers for group comparison. Propensity score matching mainly selected the most

314 fragile patients in the OR group and the less fragile patients in the FEVAR group. Thus,  
315 results are only applicable to patients at intermediate risk. A significant proportion of patients  
316 were included before 2010, period during which the FEVAR technique in both centers had  
317 not reached a state of maturity. We performed a sensitivity analysis with the inclusion of the  
318 date of surgery in the propensity score. It reduces our numbers and the results were similar  
319 except for length of stay which was no longer statistically different and late reinterventions  
320 which could no longer be compared due to the lack of events in the OR group. Therefore, we  
321 choose not to consider year of surgery in our propensity score. Devices were inserted via  
322 femoral cutdowns in half of the patients whereas most of procedures are currently performed  
323 percutaneously. Ventana and Anaconda devices were used during the study period. Ventana  
324 are not available anymore. We choose to include them anyway in our study. The Cook device  
325 was by far the most commonly used, given limited numbers of Anaconda and Ventana  
326 devices, one can hardly expect that statistical comparison of mortality and complication rates  
327 according to the type of device can lead to a valid and informative conclusion. Furthermore,  
328 we believe such a comparison would be out of the scope of this specific study. We did not  
329 observe any significant difference in the primary or secondary endpoints between the different  
330 endografts used in our population (Supplemental files: *Table 7* and *Table 8*). Most of patients  
331 were implanted two fenestrated-vessel devices. Even for juxtarenal AAA, due to improved  
332 technical skills and better patient selection, there is a current trend in both centers to promote  
333 devices with three or four fenestrations. Indeed, in order to reduce the risk of type Ia endoleak  
334 that may occur during follow-up, there is a current trend to treat JRAA using devices with  
335 three or four fenestrations instead of two. Whether this strategy is truly beneficial for patients  
336 remains controversial. Some studies have shown increased risks when devices with three or  
337 four fenestrations were used(23)(24)(4). Other studies suggested after the learning curve is  
338 reached, procedure complexity does not influence outcomes significantly(25)(26). Thus, the

339 rationale of treating JRAA with three or four fenestrated stent grafts rather than with a durable  
340 open repair is questionable. However, one could argue that with enhancement of anaesthetic  
341 and surgical techniques, OR for juxtarenal AAA also improved over the last decade. For  
342 example, we currently tend to cross-clamp the aorta in an infrarenal position whenever  
343 possible the time to ligate lumbar arteries. The clamp is subsequently moved to a suprarenal  
344 position to perform the proximal anastomosis. This allows to reduce the clamping time of  
345 renal arteries to 10-20 minutes. Finally, because of the retrospective design, there is lack of  
346 data on cause of deaths in both groups and long-term target artery patency in the OR group.

## 347 CONCLUSION

348 In the present study, there was no statistical difference in terms of mortality, between FEVAR  
349 and OR groups for JRAA. FEVAR for JRAA was associated with a significant reduction of  
350 postoperative overall major complications compared with OR. During follow-up, even if there  
351 was a non-significant trend towards higher mortality rates in the FEVAR group, the survival  
352 rates after FEVAR and OR at 12 and 36 months were statistically non-significant. There were  
353 significantly more late reinterventions in the FEVAR group compared with the OR group.  
354 And 11.3% of the FEVAR patients had a persistent endoleak during the follow-up. This is  
355 why open surgery still has its place in the management of JRAA in fit patients.



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## 433 TABLES AND FIGURES LEGENDS

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