A rare case of an acute type B aortic dissection contained infrarenal rupture of the false lumen after prior endovascular abdominal aneurysm repair

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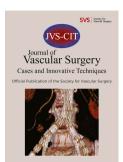
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- 1 A rare case of an acute type B aortic dissection contained infrarenal rupture of the false
- 2 lumen after prior endovascular abdominal aneurysm repair
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the high risk for complications and mortality.

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### **Abstract**

13 New-onset acute type B aortic dissection (ATBD) after prior endovascular aneurysm repair 14 (EVAR) is extremely rare. The extension of aortic dissection can cause destabilization of the 15 previously implanted stent graft, thrombosis of the stent graft, and the rupture of the aneurysmal 16 sac with high mortality rate without therapy. This report describes the case of a 66-year-old 17 patient complaining of sudden abdominal pain radiating to both flanks. A computed tomography 18 angiography (CTA) of the aorta revealed ATBD with infrarenal rupture of the false lumen after 19 endovascular abdominal aneurysm repair five years prior. The patient underwent infrarenal open 20 surgical conversion with suprarenal aortic clamping and implantation of a bifurcated dacron graft. 21 Postoperatively no serious complications occurred of the treatment, except a fascial dehiscence. 22

In these cases, the patients can be treated in an emergency situation with open repair in spite of

1 **Keywords:** ATBD; aortic abdominal rupture; acute type B dissection; endovascular repair; 2 open repair 3 Introduction Acute type B aortic dissection accounts for 25% to 40% of all aortic dissections<sup>1</sup>. Rupture of 4 5 acute type B aortic dissection is uncommon, occurring in fewer than 5% of all acute type B dissections<sup>2</sup>. Acute aortic dissection in combination with infrarenal aortic rupture after EVAR is 6 7 an extremely rare complication. Few case reports have described that treatment of patients with 8 ruptured aortic Standford type B dissection after EVAR are often lethal<sup>3</sup>. In one of two cases, the 9 patient died after the open aortic repair at postoperative day 4, as a result of multisystem organ 10 failure. In the other case, the patient died without therapy after suffering cardiac arrest. Regarding 11 the endo repairs in these cases, there are no information in the literature. 12 This report describes a case of a patient who first received EVAR for treatment of an abdominal 13 aortic aneurysm (AAA) and five years later developed complicated ATBD with infrarenal aortic 14 rupture. In this case, according to Society for Vascular Surgery reporting standards for type B 15 aortic dissection from 2020, the primary entry tear of the hyperacute aortic dissection (< 24 16 hours) is in zone 3 extending to zone 9 (type B<sub>3,9</sub>). Open surgery of the patient was the last 17 option for the treatment in this case. 18 For publishing, data and images were obtained from a patient publication consent form. 19 Case report 20 A 66-year-old-man was seen with normal blood pressure (120/70 mmHg) after rupture of the 21 infrarenal aorta and complained of sudden abdominal pain radiating to both flanks at our 22 emergency department. The patient presented with a history of smoking and arterial hypertension. He had been taking candesartan 16mg as an antihypertensive drug and 23 24 acetylsalicylic acid 100mg per day. The physical examination of patient was unspectacular, with

1	the exception of abdominal pain radiating to both flanks. Blood analysis revealed a high serum
2	creatinine level (3.63 mg/dl). The patient mentioned treatment history of EVAR (AFX,
3	Endologix, Irvine, USA) because of an asymptomatic, abdominal aortic aneurysm (diameter 5.1
4	cm) five years prior. A computed tomography angiography (CTA) scan of the aorta showed an
5	acute aortic dissection type B with main entry distal to the left subclavian artery in combination
6	with the infrarenal rupture of the false lumen, which was not present in the last follow-up
7	contrast-enhanced CT scan of the abdomen and pelvis three years prior. Spontaneous aneurysma
8	sac shrinkage after EVAR was not observed during last follow-up. The false lumen extended
9	distally to the infrarenal aorta (segment 4) beyond the stent graft with retroperitoneal hematoma
10	suggestive for rupture (Fig.1). CTA scan showed the false lumen with major supply for the right
11	kidney, as well as the superior mesenteric artery, while the celiac trunk had dual supply from
12	both lumens. The left kidney was already non-functional in the CTA scan after ATBD.
13	Additionally, the CTA showed an aneurysmal lesion of the right common iliac artery of 3.75cm,
14	and left internal iliac artery aneurysm of 3.1cm.
15	The patient declined surgery due to potential complications, such as dialysis and intestinal
16	ischemia. However, after 12 hours, the patient decided for the surgery after speaking with his
17	family. After receiving informed consent, the patient underwent infrarenal open surgical
18	conversion, with suprarenal aortic clamping of 30 minutes, and implantation of a bifurcated
19	dacron graft (Hemagard, GETINGE, Göteborg, Sweden). The identification of the true lumen of
20	dissection was unproblematic because the aortic neck with the stent graft inside was easy to
21	identify. Proximal aortic anastomosis including the whole aortic wall was infrarenal with end-to-
22	end configuration after removal of the stent graft. Distal anastomoses were performed on the left
23	external iliac artery and right common iliac artery with end-to-end configurations. The non-

1	functional left renal artery after ATBD and left internal iliac artery with aneurysm were ligated.
2	The duration time of surgery was 210 minutes. Estimated blood loss was 200 ml.
3	The postoperative course was significant for a transient elevation of creatinine, and oliguria that
4	improved gradually. Ultrasound was performed, which showed normal perfusion of the right
5	kidney. The relevant postoperative complication was a fascial dehiscence after two weeks, which
6	was closed with Vicryl mesh (Johnson& Johnson, New Brunswick, USA) and treated with
7	vacuum therapy for another two weeks. Afterwards, wound dehiscence was treated
8	conservatively.
9	The postoperative computed tomography scan after three weeks showed unaltered type B
10	dissection of the thoracic aorta, and successful surgical treatment of ATBD with infrarenal false
11	lumen rupture after EVAR, and reperfusion of the right kidney artery (Fig.2).
12	
13	Discussion
14	Antegrade TBAD after endovascular repair of AAA is an extremely rare, and potentially lethal
15	condition. Only a few cases describe aortic dissection post aortic aneurysm repair. <sup>3-7</sup>
16	Nomura <sup>3</sup> et al. summarized in 2018 case reports of similar antegrade complicated TBAD. They
17	found that timing of aortic dissection after abdominal aorta repair varied from 11 weeks to 10
18	years. Different complications after ATBD in combination with EVAR were described, such as
19	endograft thrombosis in 8/10 cases, endograft collapse in 8/10 cases, or rupture in 2/10 cases,
20	with often disastrous outcomes. Treatment with aortic rupture is described as difficult or not
21	possible because of risk of multiorgan failure postoperatively.
22	In the case described above, life-threatening type B aortic dissection in normotensive patient
23	
	occurred five years after uneventful treatment for infrarenal aneurysm with EVAR. Normal blood

1 rupture of the infrarenal aorta, but most likely the acute type B dissection was caused by high 2 blood pressure. In the literature, all patients with the same conditions were normotensive<sup>8-9</sup>. 3 The recommended surveillance for endovascular graft is usually 3- and 6-months post repair, and then annually 10. In this case, the patient lost follow up in the last three years prior to the ATBD 4 5 with infrarenal rupture of false lumen. Spontaneous type B aortic dissection is the most probable 6 cause, given the period of five years between ATBD and the repair of AAA. The device-related, 7 procedure-related dissection of aorta after stent graft implantation will usually occur in a shorter 8 time after surgery. The post-procedure contrast-enhanced CT scan of the whole agrta of the 9 patient on the third postoperative day showed no evidence of aortic dissection. 10 Presumably the stress to the calcified, aortic wall from high blood pressure weakened the aorta 11 wall, which caused an acute type B dissection with main entry distal to the left subclavian artery. 12 Apparently, the increased pressure in diastole in the false lumen was sufficient to dissect the 13 aortic wall beyond the proximal origin of the prior stent graft. The radial force exerted by the 14 endograft over the length of the proximal neck was obviously exceeded by the dissection lumen. 15 The false lumen of the dissection terminated in the excluded aneurysm sac, resulting in a lack of 16 outflow, and caused a rupture of infrarenal aorta. 17 In our opinion, the design of the aortic endograft doesn't have an impact on the risk of rupture 18 during TBAD. The aortic dissection is an arterial wall disease, in which the inner layer of the 19 aortic wall tears and separates from the middle layer of the aortic wall. The dissection is not 20 between the aortic wall and the endograft. If the endograft would increase the rupture risk during 21 dissection, it would lead to a local rupture or penetration of the stent graft in the aortic neck. In 22 our case we don't see that. 23 Endovascular treatment of complicated type B dissection using TEVAR for entry occlusion is 24 common. The standard treatment with TEVAR in ATBD with combination of infrarenal aortic

1	rupture of false lumen after EVAR would be ineffective because of multiple reentries, and
2	therefore not a treatment option. Only open abdominal repair in spite of the high risk for
3	complications and operative mortality is possible.
4	In the present case, after suprarenal clamping of the aorta within 30 minutes, there was no celiac
5	artery (CA), superior mesenteric artery (SMA), or lower leg malperfusion. Postoperatively, the
6	patient had a few days of acute kidney insufficiency, while in the intensive care unit which
7	improved gradually despite single kidney perfusion.
8	Postoperative computed tomography scan showed unaltered type B dissection of thoracic aorta,
9	which now classified as uncomplicated, and without conspicuous finding after the surgical
10	treatment.
11	Conclusions
12	We described a very rare case of ATBD in combination with infrarenal false lumen rupture after
13	EVAR. Open surgery of complicated infrarenal aorta rupture and non-functional left kidney after
14	acute type B dissection is the treatment of choice in hemodynamically stable patients despite the
15	high risk of complications, especially dialysis. Treatment of postoperative uncomplicated type B
16	dissection of thoracic aorta is conservative, with follow-up examination with CTA scan of the
17	chest in 6 months and annually. Nevertheless, patient condition, morbidity, and age of patient
18	play a role in surgical outcome, and therefore has to be analyzed individually.
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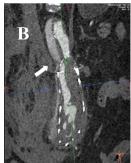
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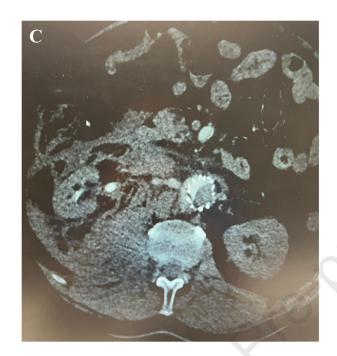
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Journal Pre-problem



John Richard Control









John Marie Control

**Figure 1.** Preoperative CT images of aorta demonstrating acute type B dissection at the level of the descending aorta with (A), hematoma after infrarenal rupture of the false lumen ATBD after EVAR with AFX stentgraft (B), no perfusion of left renal artery after ATBD (C). Arrow shows an entry tear (A) and re-entry tear (B) on the CT scan.

**Figure 2.** Postoperative CT image of a orta after removal of the stent graft AFX and implantation of a bifurcated dacron graft (A) with reperfusion of the right kidney artery (B)