



# Endovascular stenting of a complicated type B aortic dissection in an 11-year-old patient

Case Report

Arkadiusz Kazimierczak, MD, PhD<sup>a,\*</sup>, Paweł Rynio, MD<sup>a</sup>, Piotr Gutowski, MD, PhD<sup>a</sup>, Tomasz Jedrzejczak, MD, PhD<sup>b</sup>

### **Abstract**

Rationale: Endovascular aortic repair in children in the case of aortic dissection (AD) is currently unavailable. This is the first report of aortic dissection type B in an 11-years old child treated in endovascular way.

Patient concerns: Complicated AD. Since bowel malperfusion and aortic rupture the surgery was necessary in emergency.

Diagnoses: Computed angio tomography.

Intervention: The modified PETTICOAT technique (Provisional Extension To Induce Complete Attachment) was used.

Outcome: Full recovery.

**Lessons:** For the first-time telescope modification were used, to allow aorta to grow with a child. Such a strategy seems to be possible with long overlap and lack of oversizing between implants.

**Abbreviations:** AD = aortic dissection, Angio-CT = angio computed tomography, BMS-XL = bare metal stents extra large, BMT = best medical treatment, CT = celiac trunk, FL = false lumen, LCCA = left common carotid artery, LRA = left renal artery, LSA = left subclavian artery, PETTICOAT = provisional extension to induce complete attachment, RA = renal artery, RRA = right renal artery, SMA = superior mesenteric artery, TEVAR = thoracic endovascular aortic repair, TL = true lumen.

Keywords: aortic dissection, children, PETTICOAT, stent, stent-graft

# 1. Introduction

Reports regarding thoracic endovascular aortic repair TEVAR/PETTICOAT for children in the case of aortic dissection (AD) were not found, thought 1 traumatic aortic rupture TEVAR report was available. The lack of data is mainly due to the fact that AD in children is a highly rare disease (less then 30 cases in literature for < 25 years old). It is normally seen in cases of Marfan syndrome, as well as anomalies of the aortic valve and arch, or in cases of chest trauma (this was not being the case in our patient). In complicated AD, TEVAR or PETTICOAT strategy is usually recommended in adults. Phon This is the first report of aortic dissection type B in an 11-year-old child treated in endovascular way. Since bowel malperfusion and aortic rupture, the surgery was necessary in emergency. The PETTICOAT technique (Provisional Extension to Induce Complete Attachment) was used.

Editor: N/A.

The authors report no conflicts of interest.

Copyright © 2018 the Author(s). Published by Wolters Kluwer Health, Inc. This is an open access article distributed under the Creative Commons Attribution-NoDerivatives License 4.0, which allows for redistribution, commercial and non-commercial, as long as it is passed along unchanged and in whole, with credit to the author.

Medicine (2018) 97:14(e0279)

Received: 29 December 2017 / Received in final form: 28 February 2018 / Accepted: 7 March 2018

http://dx.doi.org/10.1097/MD.000000000010279

# 2. Report

An 11-year-old girl presented with a sudden onset of chest and mesenteric ischemia. No injury was reported, with no marfanoid features and a normal aortic valve. A final diagnosis of type B aortic dissection was established with the aid of angio-CT. Entry was located below left subclavian artery (LSA) with complete intima tear, which was causing dynamic stenosis of the aorta. True lumen (TL) was compromised along the dissection. Celiac trunk (supply from TL) was patent. Superior mesenteric artery (SMA) was occluded at a short distance. However, efficient collateral circulation to the bowels was present. Kidney arteries with dynamic stenosis were supplied via a simultaneous false lumen (FL) and TL. Best medical treatment (BMT) was first applied. Nevertheless, it was found that effective hypertension control was impossible (250 mmHg systolic). After 2 days, mesenteric ischemia deteriorated to peritonitis and sepsis (without perforation). Second angio-CT was applied showing static/critical aortic stenosis below the LSA due to partially thrombosed false lumen. SMA static occlusion at a distance of 6 cm with thrombosis of the FL due to complete TL collapse in the visceral and infra-renal aorta. Renal arteries (RA) were dissected completely and supply was only from FL. The size of the aorta at the arch was 25 mm, below LSA 31 mm, CT/SMA level 25 mm, RA 21 mm, infra-renal aorta was 13 mm, iliac common arteries were 7.5 mm, iliac external arteries were 6.5 mm.

(Fig. 1). True-lumen collapse

# 3. Methods

Surgical intervention was performed as an emergency. Therefore, Ethics Committee approval was waived at that time and patient consent was obtained from the parents. Extra large bare metal stents (BMS-XL) Medicut (Pforzheim, Germany, 28 mm

<sup>&</sup>lt;sup>a</sup> Department of Vascular Surgery, <sup>b</sup> Department of Cardiosurgery, Pomeranian Medical University in Szczecin, Powstancow, Wielkopolskich Szczecin, Poland.

<sup>\*</sup> Correspondence: Arkadiusz Kazimierczak, Pomeranian Medical University. Department of Vascular Surgery, Szczecin, Poland (e-mail: biker2000@icloud.com).

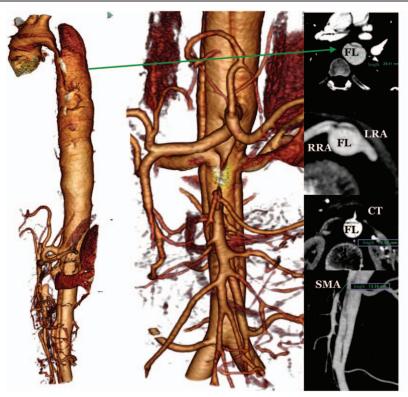


Figure 1. True-lumen collapse.

size, 200 mm length) were deployed to the thoracic and visceral aorta ending 1 cm below RA, followed by Medtronic Vailant II (Medtronic, Santa Rosa, CA, USA; VAMF2828C156TE) below LSA with long overlap (10 cm) with BMS-XL. Force ballooning

of the graft was performed to restore normal TL size and close FL completely (Initially, implants were highly compressed below the LSA). Angiography showed no endo-leak, patent CT, and both RA. SMA remained occluded. Therefore, stenting and ballooning

Initial (Short occlusion SMA, aortic syndrome)

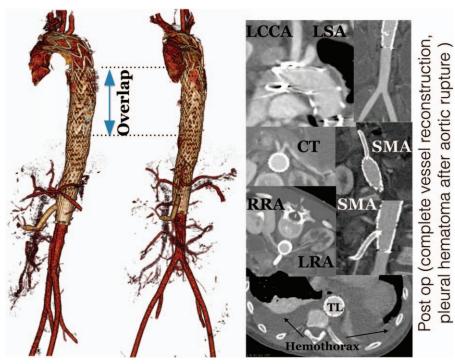


Figure 2. Aorta after endovascular treatment.

was preformed through a BMS-XL mesh with effective restoration of the flow in SMA. Two cardiac arrests occurred on the table (Pulseless Electrical Activity), though effective resuscitation was carried out. The initial arrest was before the retrieval of TL (in angiography blood and contrast extravasation along the aorta), and the second was after the reopening of the SMA, due to the bowel reperfusion syndrome. The patient was subsequently treated in the Intensive Care Unit for another 2 days. Diarrhea continued for another 3 days. Kidney function was found to be completely normal after 4 days. Control angio-CT showed very good early remodelling (Fig. 2).

The patient was discharged 3 weeks later without any symptoms. A next angio-CT is planned in 6/12 months time.

# 4. Discussion

Open surgical repair is possible, but is seen as risky in children.<sup>[11]</sup> Moreover, it is accepted that a prosthesis will not able to grow with the child's body. In this case, an endovascular procedure was considered. However, this was significantly modified due to expected body growth (weighed 40 kg high 159 cm). Therefore, a long (10cm) overlap between the stent-graft and BMS-XL was carried out. This is hoped to act as a telescope for the next few years. The lack of oversizing between implants should facilitate this requirement. Moreover, implant oversizing according to a ortic size varies from 0 mm to 7 mm from the top to the bottom and should keep them attached at the landing zones. It should be noted that PETTICOAT was the only strategy to re-open the orifice of CT, SLA, and RA. Implanting BMS-XL to the TL is usually enough to restore flow, provided that there is no FL thrombosis in the branches. Unfortunately, SMA was thrombosed and occluded, therefore, this required immediate reopening and subsequent stenting. Follow up will continue with this patient into adulthood.

# 5. Conclusion

To our knowledge, this is the youngest patient to be successfully treated with PETTICOAT (TEVAR+BMS-XL) in a complicated

type B aortic dissection. Such a strategy only seemed possible with long overlap and a lack of oversizing between implants (telescope phenomenon).

# Author conctributions

Conceptualization: A. Kazimierczak, P. Rynio.

Supervision: P. Gutowski.

Writing - original draft: A. Kazimierczak.

Writing - review & editing: P. Rynio, T. Jedrzejczak.

# References

- Gunabushanam V, Mishra N, Calderin J, et al. Endovascular stenting of blunt thoracic aortic injury in an 11-year-old. J Pediatr Surg 2010;45:2010.
- [2] Zalzstein E, Hamilton R, Zucker N, et al. Aortic dissection in children and young adults: diagnosis, patients at risk, and outcomes. Cardiol Young 2003;13:341–4.
- [3] Goertz K, Diehl AM, Vaseenon T, et al. A catastrophic complication. Acute dissection of an aortic aneurysm in a child with Marfan's syndrome. J Kans Med Soc 1978;79:115–7.
- [4] Burak Onan , Alper Güzeltaş\* , Ender Ödemiş\* İB. Acute aortic dissection in a 10-year- old boy with bicuspid aortic valve. Anadolu Kardiyol Derg 2012;12:693–5.
- [5] Lindinger A, Volkmer I, Reichert H, et al. Dissection of an aneurysmic ascending aorta in a child with atresia of the aortic isthmus. Pediatr Cardiol 1997;18:146–8.
- [6] Panja M, Kumar S, Panja S, et al. Aortic dissection in a non-marfanoid child. J Assoc Physicians India 1990;38:369–71.
- [7] Khanna PC, Rothenbach P, Guzzetta PC, et al. Lap-belt syndrome: Management of aortic intimal dissection in a 7-year-old child with a constellation of injuries. Pediatr Radiol 2007;37:87–90.
- [8] Civilibal M, Sever L, Numan F, et al. Dissection of the abdominal aorta in a child with Takayasu's arteritis. Acta Radiol 2008;49:101–4.
- [9] Hughes GC, Andersen ND, McCann RL. Management of acute type B aortic dissection. J Thorac Cardiovasc Surg 2013;145(3 Suppl):S202-7.
- [10] He H, Yao K, Nie WP, et al. Modified Petticoat technique with preplacement of a distal bare stent improves early aortic remodeling after complicated acute Stanford type B aortic dissection. Eur J Vasc Endovasc Surg 2015;50:450–9.
- [11] Deml K-F, Schoepf UJ, Henzler T. Acute aortic dissection in a 9-year-old boy with chest pain. J Am Coll Cardiol 2010;56:e49.