Refactory renovascular hypertension secondary to Takayasu arteritis treated with aorto-mesenteric bypass

DEAR EDITOR, Takayasu arteritis is traditionally seen in young Asian females but is also seen in a worldwide distribution. A recent study found a female-to-male ratio of 12:1, with peak age of onset between 20 and 30 years of age [1]. Here, we present a case of Takayasu arteritis in an unusual patient population with atypical vessel distribution [2].

A 52-year-old Caucasian non-smoker male with a history of renovascular hypertension (RVH), atrio-mesenteric right kidney and hypopituitarism presented with chest pain radiating to his back. Laboratory work revealed a normal troponin level, elevated D-dimer (0.52 μg FEU/ml), elevated ESR (99 mm/h) and elevated CRP (5.1 mg/dl, normal < 0.8 mg/dl). A CT angiography of the chest/abdomen demonstrated thickening and inflammation of his descending thoracic aorta, abdominal aorta, mesenteric vessels, bilateral iliac and bilateral renal arteries. Further history revealed that he was diagnosed with RVH 10 years before, secondary to right renal artery stenosis. Based on the current CT angiography findings, he was diagnosed with possible Takayasu arteritis.

After rheumatological evaluation, our patient was started on high-dose prednison and AZA, with improvement in inflammatory markers (CRP 1.2 mg/dl; ESR 19 mm/h). His AZA was then discontinued owing to transaminitis. Unfortunately, he was then lost to follow-up.

The patient re-presented 1 year later with another acute exacerbation (CRP 14.8 mg/dl; ESR 31 mm/h), whereupon he was treated again with high-dose prednisone. Tocilizumab was added, with symptom improvement and inflammatory marker normalization (CRP 0.1 mg/dl; ESR 8 mm/h). However, our patient had challenges with tocilizumab infusions and compliance, resulting in subsequent flare-ups. Follow-up CT angiography obtained while on high-dose prednisone continued to show worsening vasculitis. He underwent magnetic resonance angiography, which confirmed progression of large vessel vasculitis and near-critical stenosis of the coeliac, superior mesenteric, inferior mesenteric and bilateral renal arteries.

Given his young age and continued progression, multidisciplinary discussions determined that surgical bypass would be the next best step. Pre-operatively, he was not in an acute inflammatory state (CRP 0.1 mg/dl; ESR 4 mm/h) and had received tocilizumab 3 weeks before. This bypass was performed with a bifurcated synthetic graft (Fig. 1A). Intra-operatively, he was noted to have significantly friable vessels. The proximal graft was anastomosed to the supraceliac aorta. One graft limb was anastomosed to the coeliac trunk base (Fig. 1B). The second limb was tunneled to the superior mesenteric artery (Fig. 1C). The left renal artery was ligated from the aorta and anastomosed to the graft. The final anastomosis was performed between the superior mesenteric artery and the graft (Fig. 1D). The patient tolerated the procedure well. Post-operatively, he continued to improve, with down-trending creatinine levels, and was weaned off his three anti-hypertensive medications.

In retrospect, the initial presentation of Takayasu arteritis in this patient was probably the RVH diagnosed 10 years prior. At that time, the patient did not exhibit any additional signs or symptoms of Takayasu arteritis. In Takayasu arteritis patients, it is not uncommon to have renal artery involvement, with upwards of 60% involvement in Asia, which can result in RVH in 33–80% of cases [3]. Often, this RVH becomes refractory to multimodal medical therapy.

Historically, arterial angiography has been the gold standard for diagnosis [4]. However, CT angiography and magnetic resonance angiography have become used increasingly. The main imaging findings are mural thickening and luminal changes. Mural thickening commonly appears as a double ring, with the inner ring representing intimal swelling and the outer ring representing active inflammation in the medial and adventitial layers [5]. Mural thickening is commonly associated with stenosis, seen in 90% of Takayasu arteritis patients. The most commonly stenosed vessels are the thoracic and abdominal aorta, subclavian, common carotid and renal arteries. Additionally, there have been reports of aortic vessel dissections, but not of coeliac or common iliac artery dissection like our patient [6].

Given this patient’s young age and worsening disease on escalating medical therapy, he was evaluated for surgical intervention. Open bypass surgery was chosen over stenting owing to increased risk of restenosis. The higher rate of post-operative restenosis in endovascular approaches is secondary to progression of the native disease process and placement of a foreign body directly at the site of inflammation [7]. In a comparison of open vs endovascular interventions for Takayasu arteritis, the 10-year primary patency rate was 48.8% for open surgical and 31.8% for endovascular [7]. Inflammatory markers were normalized for our patient before surgical intervention, because performing reconstructive procedures during active vasculitis has been
shown to increase complications such as anastomotic dehiscence, restenosis rates and disease progression in other arterial areas [8].

In conclusion, we present a case of refractory Takayasu arteritis in an atypical patient with unusual vessel involvement, managed with surgical bypass. The progression of this patient’s disease is multifactorial, and the lack of compliance with treatment is likely to have played a substantial role in disease progression. This unique case illustrates the successful use of a triple bypass (aorto-coeliac, aorto-superior mesenteric artery and aorto-renal) for the treatment of refractory RVH secondary to Takayasu arteritis.

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**Data availability statement**

Data are available upon reasonable request by any qualified researchers who engage in rigorous, independent scientific research, and will be provided following review and approval of a research proposal and Statistical Analysis Plan (SAP) and execution of a Data Sharing Agreement (DSA). All data relevant to the study are included in the article.

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References

1 Onen F, Akkoc N. Epidemiology of Takayasu arteritis. Presse Med 2017;46:e197–e203.

2 Gudbrandsson B, Molberg Ø, Garen T, Palm Ø. Prevalence, incidence, and disease characteristics of Takayasu arteritis by ethnic background: data from a large, population-based cohort resident in Southern Norway. Arthritis Care Res 2017;69:278–85.

3 Chaudhry MA, Latif F. Takayasu's arteritis and its role in causing renal artery stenosis. Am J Med Sci 2013;346:314–8.

4 Chung JW, Kim HC, Choi YH et al. Patterns of aortic involvement in Takayasu arteritis and its clinical implications: evaluation with spiral computed tomography angiography. J Vasc Surg 2007;45:906–14.

5 Zhu FP, Luo S, Wang ZJ et al. Takayasu arteritis: imaging spectrum at multidetector CT angiography. Br J Radiol 2012;85:e1282–92.

6 Yang KQ, Yang YK, Meng X et al. Aortic dissection in Takayasu arteritis. Am J Med Sci 2017;353:342–52.

7 Diao Y, Yan S, Premaratne S et al. Surgery and endovascular management in patients with Takayasu's arteritis: a ten-year retrospective study. Ann Vasc Surg 2020;63:34–44.

8 Kim Y-W, Kim D-I, Park YJ et al. Surgical bypass vs endovascular treatment for patients with supra-aortic arterial occlusive disease due to Takayasu arteritis. J Vasc Surg 2012;55:693–700.