CASE REPORT

Spontaneous Isolated Common Iliac Artery Dissection Treated with Self-Expandable Stent in a 38-year-old Patient: A Case Report

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Introduction: Isolated iliac artery dissection (ISIAD) without the involvement of the aorta is a rare medical condition.

Report: A case of a 38-year-old man with sudden onset of rest pain and paraesthesia on the right lower limb (RLL) is presented. Upon admission, the RLL was pulseless, with mild paraesthesia in the foot. The patient underwent computed tomography angiography, which revealed isolated common iliac artery (CIA) dissection followed by endovascular treatment (stenting) of the CIA dissection, with an instant therapeutic effect. Hospital stay was uneventful. The patient was discharged on the third post-procedural day.

Discussion: Endovascular treatment of ISIAD is a viable treatment modality, with low periprocedural complications, mortality, and morbidity. Owing to its mini-invasiveness, it is a viable treatment modality.

INTRODUCTION

Isolated iliac artery dissection (ISIAD) without the involvement of the aorta is a rare medical condition. Only a few cases regarding this condition have been published. The causes for this condition can be classified as either traumatic or non-traumatic. The most common “traumatic” causes of an ISIAD are blunt trauma, extreme physical activity, pregnancy, and iliac catheterisation.1–4 The “non-traumatic” group of medical conditions that cause ISIAD include various connective tissue disorders (CTD) such as Marfan syndrome, Ehlers-Danlos syndrome, fibromuscular dysplaia, cystic medial degeneration, Erdheim-Gsell, and atherosclerosis.5–9 The most frequent and potentially fatal complication of ISIAD is rupture.

Most of the reports presented in the literature are associated with a CTD. An extremely rare case of a 38-year-old man who presented with an ISIAD without a CTD is discussed.

Case presentation

A 38-year-old man was referred for sudden onset of rest pain and paraesthesia on the right lower limb (RLL). Upon admission, the RLL was pulseless with mild paraesthesia in the foot. Acute limb ischaemia was classified as Rutherford grade IIb. Left lower limb showed no signs of ischaemia with palpable peripheral pulsation on the anterior tibial artery and left dorsal artery of the foot. The patient’s medical history revealed only compensated arterial hypertension treated with amlodipine. The patient underwent computed tomography angiography (CTA) revealing isolated common iliac artery (CIA) dissection. False lumen compressed the true lumen of the CIA just above the iliac bifurcation. The external iliac artery (EIA), internal iliac artery, and proximal part of the common femoral artery had no blood flow on CTA (Fig. 1). The procedure was performed through the contralateral groin under local anaesthesia. Through a 6-Fr sheath, a hydrophilic guidewire was used to cross through the true lumen of the dissected CIA into the common femoral artery. The entry of the dissection was localised in the right distal CIA above the iliac bifurcation. The proximal part of the EIA was spastic without atherosclerotic infiltration. The entry of the dissected CIA and proximal part of the EIA were treated with self-expanding nitinol OptiMed Sinus stent 7 × 60 mm (OptiMed Medizin-ische Instrumente, Ettingen, Germany). Post-dilatation was completed with Admiral Xtreme PTA balloon 6 × 60 mm (Admiral Xtreme; Medtronic, Minneapolis, MN, USA). The procedure was technically successful, with blood flow fully restored into the RLL (Fig. 2). The left femoral artery was closed with an AngioSeal vascular closure device (AS, St. Jude Medical, Minneapolis, MN,
USA). Post-procedural hospital stay was uneventful. No reperfusion compartment syndrome occurred after the procedure. The patient was discharged on the third post-procedural day with palpable peripheral pulsation. The patient was given a daily dose of 100 mg acetylsalicylic acid (aspirin) and will undergo an annual follow-up with a Doppler ultrasound check-up. The patient’s collagen connective tissue work-up was negative.

Written informed consent was obtained from the patient for publication of this case report and accompanying images. A copy of the written consent is available for review by the Editor-in-Chief of this journal.

**DISCUSSION**

To date, only a few cases of ISIAD have been reported in the literature. The most frequent causes of ISIAD are connective tissue disorders. They are characterised as “non-traumatic causes”. Under extreme circumstances, ISAID can be caused by so-called “traumatic causes”. However, idiopathic cases are very sporadic. Based on the literature search, only two previous papers have reported an ISIAD.

Criteria for the treatment of asymptomatic patients with ISIAD are not yet defined, owing to the rarity of the condition. The study by Liang et al. reported that asymptomatic patients could be safely treated conservatively. However, the question regarding further follow-up is still not answered. Owing to the nature of the disease, these patients are prone to aneurysmatic degeneration of arteries. Patients should undergo an annual check-up for screening an arterial aneurysms formation and their eventual progression. The urgency of the intervention is determined by the presentation mode, and those presenting with acute limb ischaemia or rupture require emergency endovascular or open repair.

Endovascular treatment is associated with high technical and clinical success rates with low periprocedural mortality and morbidity when compared with open repair.
implantation of these devices can, in some cases, be limited by tortuosity of the iliac vessels. The risk of continued false lumen perfusion from distal tears has to be also taken into the account when performing an endovascular treatment. Embolisation of the false lumen can be combined with stent/stent-graft implantation to achieve an optimal result.13

CONCLUSIONS
Endovascular treatment of ISIAD is a viable treatment modality with low periprocedural complications, mortality, and morbidity.

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CONFLICTS OF INTEREST
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