

A Rare Congenital Anomaly: Interrupted Aortic Arch and Giant Ascending Aortic Aneurysm in a Young Male

Muhammed Hakan Tas, MD, Ziya Simsek, MD, Husnu Degirmenci, MD,
Hayri Ogul, MD, Ugur Aksu, MD, and Abdulmecit Kantarci, MD

Department of Cardiology, Ataturk University, Turkey

Interrupted aortic arch (IAA) is a rare congenital cardiac defect defined as complete loss of luminal and anatomical continuity between the ascending and descending segments of the aorta. Very few patients can reach adult age without surgical correction. These cases showed diagnosis of the nearly asymptomatic Type A IAA with aortography and multislice computed tomography (MSCT). A 27-year-old male patient was applied to the internal medicine department with complaint of fatigue. They heard a diastolic murmur on the aortic focus and referred the patient to our cardiology polyclinic. Physical examination showed regular heart beat with blood pressures of 150/90 mm Hg in the right arm, 140/90 mm Hg in the left arm, 100/60 mm Hg in the right limb and 100/60 mm Hg in the left limb, and there was a diastolic murmur prominently on the aortic focus. Electrocardiogram was in sinus rhythm and normal. His chest roentgenogram (anteroposterior) showed a dilated aortic root. When we made echocardiography, we saw bicuspid aortic valve with

moderate aortic regurgitation and dilatation of the ascending aorta (70 mm). Then an attempt to perform aortography via femoral artery failed because the aortic arch was not able to go through. When we gave contrast, we saw that the aortic arch was interrupted. Aortography via the right brachial artery showed a giant ascending aortic aneurysm (70 mm), severe aortic regurgitation, interruption of the aortic arch distal to the left subclavian artery (Fig. 1, arrow) and prominent collateral circulation (Fig. 2, arrows). After aortography, the patient underwent a MSCT. MSCT confirmed the diagnosis of the IAA distal to the origin of the left subclavian artery and measured the ascending aorta as 75 mm (Fig. 3, arrow). The patient was referred to an experienced surgeon on adult congenital heart diseases, but was denied surgery due to lack of symptoms. In this case, a rare congenital anomaly was detected with transcatheter aortography and MSCT diagnostic techniques, which complement each other.

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Correspondence: Muhammed Hakan Tas, MD, Department of Cardiology, Ataturk University, Turkey
Tel: 904422318481, Fax: 904422318481, E-mail: mhakantas@gmail.com

• The authors have no financial conflicts of interest.

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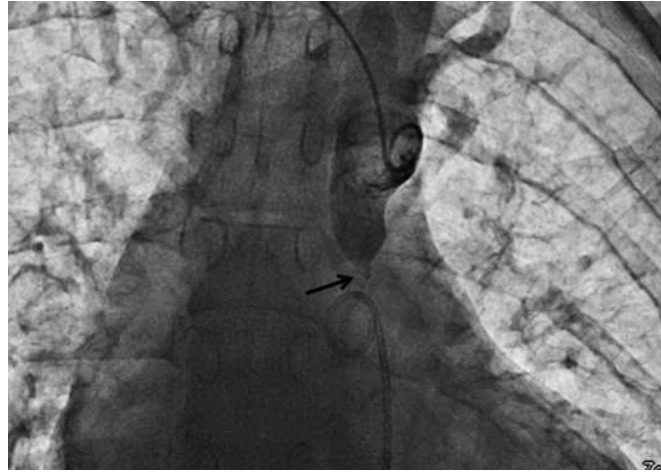


Fig. 1.

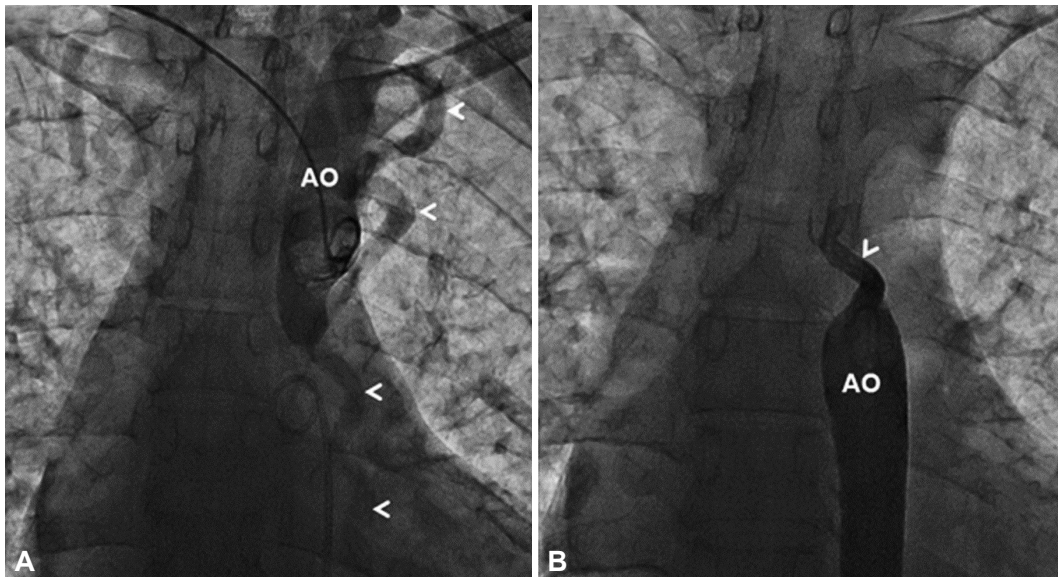


Fig. 2.

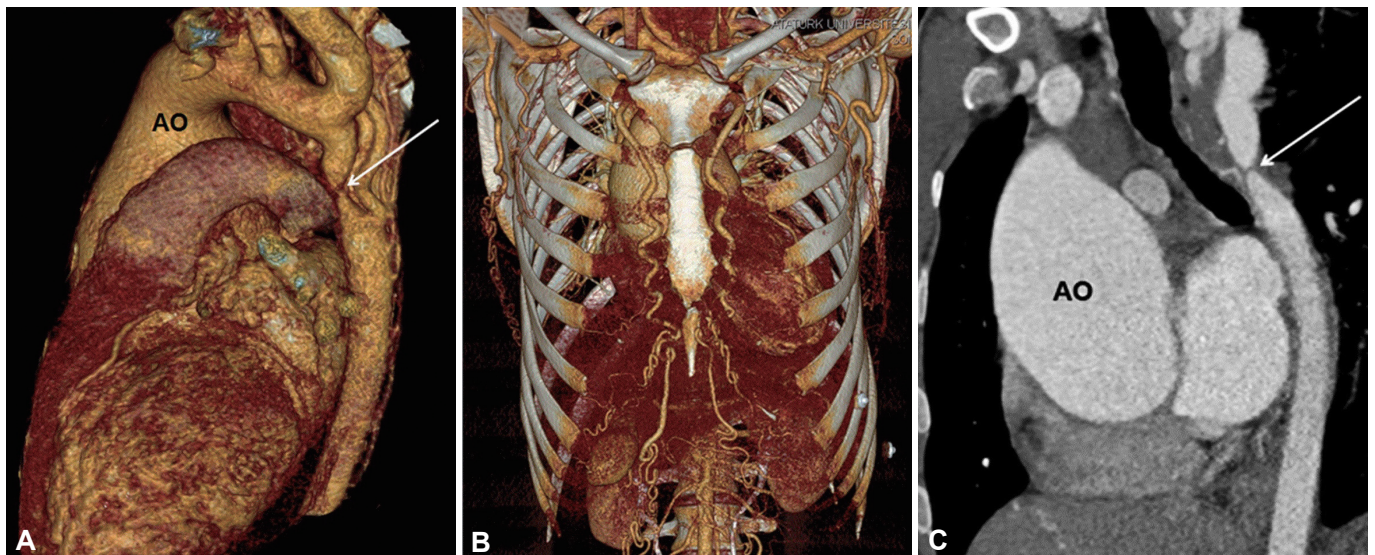


Fig. 3.