



Brief Review of Right Aortic Arch with Aberrant Left Subclavian Artery

Didem Melis Oztas, MD¹ Muzaffer Umutlu, MD² Melike Ertan, MD³ Metin Onur Beyaz, MD⁴
Serdar Badem, MD⁵ Ibrahim Erdinc, MD⁶ Mustafa Ozer Ulukan, MD⁴ Orcun Unal, MD⁷
Cenk Conkbayir, MD⁸ Ufuk Alpagut, MD³ Murat Ugurlucan, MD⁴

¹ Department of Cardiovascular Surgery, Bagcilar Training and Research Hospital, Istanbul, Turkey

² Department of Radiology, Istanbul University Istanbul Medical Faculty, Istanbul, Turkey

³ Department of Cardiovascular Surgery, Istanbul University Istanbul Medical Faculty, Istanbul, Turkey

⁴ Department of Cardiovascular Surgery, Istanbul Medipol University Medical Faculty, Istanbul, Turkey

⁵ Department of Cardiovascular Surgery, Bursa City Hospital, Bursa, Turkey

⁶ Department of Cardiovascular Surgery, Izmir Bozyaka Training and Research Hospital, Izmir, Turkey

⁷ Department of Cardiovascular Surgery, Yedikule Chest Diseases and Thoracic Surgery Training and Research Hospital, Istanbul, Turkey

⁸ Department of Cardiology, Near East University, Nicosia, Cyprus

Address for correspondence Murat Ugurlucan, MD, TEM Avrupa Otoyolu Goztepe Cikisi, No: 1, 34214 Bagcilar, Istanbul, Turkey (e-mail: muratugurlucan@yahoo.com).

AORTA 2019;7:179–180.

Abstract

Keywords

- ▶ subclavian artery
- ▶ aberrant
- ▶ aortic arch

Development anomalies of the aortic arch and its major branches are rare congenital cardiovascular disorders. In this article, we present aberrant left subclavian artery associated with right aortic arch.

Right aortic arch (RAA) is a rare malformation and is reported at a range of 0.04 to 0.1% in autopsy series.¹ The anomaly occurs in embryonic life due to the persistence of the right-fourth aortic arch, while regressing the left-fourth arch between the left common carotid artery and the left subclavian artery.² There are three types of the right-sided aortic arch as follows: Type I involves right aortic arch with mirror image branching, Type II involves right aortic arch with aberrant left subclavian artery, and Type III involves right-sided aortic arch with isolated left subclavian artery communicating with the pulmonary artery.³

Right aortic arch is generally an asymptomatic malformation and diagnosed incidentally. In the Type II form, in which the left subclavian artery is aberrant (▶ **Video 1**), patients may present to the clinic with the symptoms occurring secondary to trachea or esophagus compression or aneurysm or dissection of the vessels.¹ Dysphagia and

dyspnea are usually the symptoms at the infant period, whereas atherosclerotic changes, dissection, or aneurysm may be seen in adulthood.^{1,3,4}

Video 1

Computed tomography angiography video shows right aortic arch with aberrant left subclavian artery. Online content including video sequences viewable at: <https://www.thieme-connect.com/products/ejournals/html/10.1055/s-0039-3401999>.

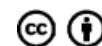
Computed tomography angiography is a valuable tool for the diagnosis because of the high resolution and the

received
July 13, 2018
accepted after revision
November 2, 2019

DOI <https://doi.org/10.1055/s-0039-3401999>.
ISSN 2325-4637.

Copyright © 2019 by Thieme Medical Publishers, Inc., 333 Seventh Avenue, New York, NY 10001, USA.
Tel: +1(212) 760-0888.

License terms



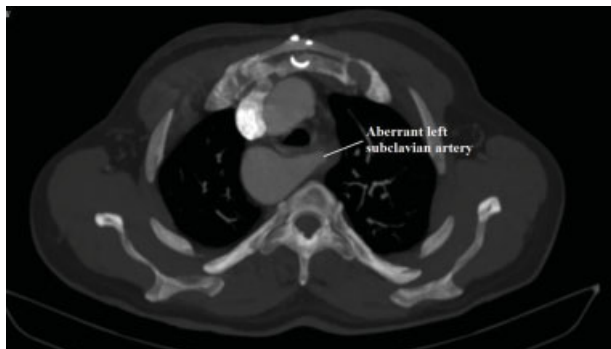


Fig. 1 Right aortic arch with aberrant left subclavian artery.

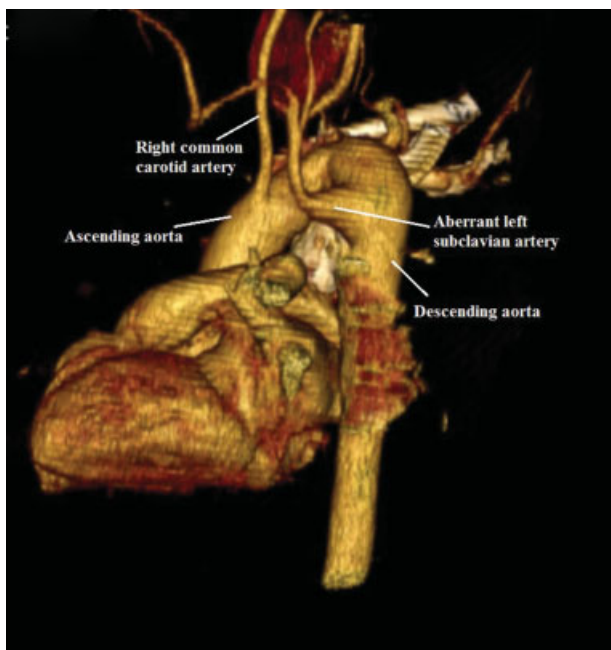


Fig. 2 3D computed tomography angiography image. Right aortic arch with aberrant left subclavian artery. 3D, three-dimensional.

speed of scanning (►Figs. 1 and 2). Also, magnetic resonance imaging is another option which may be used for diagnosis.⁵

The complications of this pathology include aneurysm formation and dissection which may be secondary to atherosclerosis in latter ages, as well as recurrent lower respiratory tract infections, and growth retardation in the early years of childhood³; hence, these patients should be followed-up lifelong.

In conclusion, the symptoms are the most important determinants for the treatment of the patients with RAA. Careful follow-up is necessary for the prevention of fatal complications.

Funding

None.

Conflict of Interest

None.

Acknowledgments

None.

References

- 1 Barr JG, Sepehrpour AH, Jarral OA, et al. A review of the surgical management of right-sided aortic arch aneurysms. *Interact Cardiovasc Thorac Surg* 2016;23(01):156–162
- 2 Salantri J. MR angiography of aberrant left subclavian artery arising from right-sided thoracic aortic arch. *Br J Radiol* 2005;78(934):961–966
- 3 Bhatt TC, Muralidharan CG, Singh G, Jain NK. Kommerell's diverticulum: A rare aortic arch anomaly. *Med J Armed Forces India* 2016;72(01, Suppl 1):S80–S83
- 4 Mubarak MY, Kamarul AT, Noordini MD. Right-sided aortic arch with aberrant left subclavian artery from Kommerell's diverticulum. *Iran J Radiol* 2011;8(02):103–106
- 5 Türkvatan A, Büyükbayraktar FG, Olçer T, Cumhuri T. Multidetector computed tomographic angiography of aberrant subclavian arteries. *Vasc Med* 2009;14(01):5–11