A Case of Aortic Dissection Complicating Right Subclavian Artery Occlusion and Mimicking Inferior Myocardial Infarction

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ABSTRACT
Introduction: Acute aortic dissection is the most common catastrophic event affecting the aorta. Aortic dissection can present with a wide range of manifestations and is easily misdiagnosed.
Case Report: Here we report a case of aortic dissection that was initially misdiagnosed as inferior wall myocardial infarction although the initial symptoms were related to right subclavian artery occlusion caused by a dissection flap. Aortic dissection was diagnosed after the patient underwent coronary angiography. The patient was managed with immediate surgical repair and was discharged in a stable condition.
Conclusion: This case report illustrates the importance of having a high index of suspicion for aortic dissection to avoid a mistaken diagnosis in patients with an electrocardiogram (ECG) suggestive of ST elevation myocardial infarction (STEMI).
Key words: Aortic dissection, myocardial infarction, subclavian artery occlusion
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Introduction
Aortic dissection (AD) is defined as the separation of layers within the aortic wall. Tears in the intimal layer result in the propagation of dissection (proximally or distally) secondary to the entry of blood into the intima–media space. Acute aortic dissection is an extremely severe condition having a high risk of mortality. The classic symptom may mimic other conditions such as myocardial ischemia, leading to misdiagnosis. Coronary malperfusion associated with aortic dissection is relatively rare, but when it occurs, it may be fatal to the patient. Up to 30% of patients suffering from aortic dissection are initially suspected of having other conditions (1). Here we report a case of initially misdiagnosed acute aortic dissection in which a dissection flap occluded the ostium of the right coronary artery and right subclavian artery. Written informed consent was obtained from patient.

Case Report
A 62-year-old female with a medical history of hypertension was referred to emergency service because of chest and right upper extremity pain and syncope. Her initial vital signs showed blood pressure of 100/60 mmHg and heart rate 80 beats/min. ECG at admission showed ST elevation in the inferior leads (Figure 1). Emergent coronary angiography was performed. Coronary angiography revealed an ascending aortic dissection with normal coronary arteries (Figure 2, 3).

A bedside transthoracic echocardiogram was also performed and showed a dilatation of the ascending aorta with an evidence of intimal tear and moderate aortic regurgitation. The vascular surgery team was immediately consulted, and the patient was transferred for emergency surgery. The Bentall procedure was performed.
Acute aortic dissection is the most common life-threatening disorder affecting the aorta, with an estimated annual incidence of approximately 5 to 30 per million (2). The early mortality rate in cases of acute aortic dissection is very high, with an hourly mortality rate of up to 1%–2% being reported in the first several hours after dissection (3). There are two major classification schemes of aortic dissection, based on the location of the dissection: DeBakey and Stanford classifications. Stanford type A dissections involve the ascending aorta and type B dissections occur distal to the subclavian artery (4). On the other hand, in the DeBakey classification, type 1 dissections begin in the ascending aorta and extend to the descending one, while type 2 dissections involve the ascending aorta only and type 3 dissections begin in the descending aorta distal to the left subclavian artery (5). The treatment depends on the site, with emergency surgery being recommended for acute type A dissections and initial medical therapy being recommended for type B dissections.

In our patient, aortic dissection was diagnosed after the patient underwent coronary angiography. Coronary angiography showed normal coronary arteries. Myocardial infarction most likely resulted from functional coronary artery occlusion due to aortic dissection, in which the dissection flap temporarily occluded the ostium of the right coronary artery and affected the blood flow to the vessel. In addition, in our case, aortic dissection extended to the brachiocephalic trunk, closing down and occluding the ostium of the right subclavian artery. Our patient had upper extremity pain, intermittent pulse deficits, and pseudohypotension that may be due to right subclavian artery occlusion.

In the literature, there have been few cases of ascending aortic dissections presenting with STEMI complicating right subclavian artery occlusion (10). However, our case mimics inferior myocardial infarction and complicating right subclavian artery occlusion and exhibits correlated symptoms and findings.

Conclusion
In conclusion, aortic dissection has still been frequently misdiagnosed as acute coronary syndrome. Aortic dissection should always be considered in the differential diagnosis of patients presenting with acute infarction, particularly inferior infarction, especially when their risk factors, symptoms, or examination findings are compatible with this diagnosis.

Informed Consent: Written informed consent was obtained from patient who participated in this study.

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