steno-sis, mild mitral regurgitation, MVA was calculated as 1.2 cm² with planimetry and 1.35 cm² with Doppler. The maximum and mean gradients across the mitral valve were 19 and 11 mmHg respectively. TEE revealed a thrombus at LAA base, not protruding into LA and measured as 1.2x1.5 cm. The Wilkins valve score was calculated as 9 (Fig. 3). The patient refused to undergo open-heart surgery for mitral valve replacement. Based on the success of the previous case, and the same type of thrombus, which was restricted to the base of the LAA, PMBV was offered and the risk of the procedure was explained in detail. PMBV was performed with the help of TTE. After completion of the procedure without any complication, echocardiographic parameters were as follows, MVA was 1.7 cm², systolic PAP was 35 mmHg, maximum and mean gradients across the valve were 10 mmHg and 5 mmHg respectively.

**Discussion**

PMBV is the treatment of choice for patients with rheumatic mitral stenosis and suitable valve anatomy. Stroke was one of the catastrophic complications of PMBV. While ACC/AHA guidelines emphasizes LA thrombus as a contraindication for PMBV (level of evidence C), there is no solid evidence for cases with left atrial appendage thrombus (3).

Some operators do not consider LA thrombus as an absolute contraindication (4).

Koca et al. (5) performed PMBV to nine patients with symptomatic mitral stenosis and thrombus restricted to the LAA. These procedures were performed under TEE guidance and there were no thromboembolic events. Consequently, researchers concluded that in selected cases, PMBV under TEE guidance was safe and thrombus restricted to the LAA is not an absolute contraindication to the procedure.

The most comprehensive research about this comes from small scale trials comprising 28 and 30 patients. Manjunath et al. (4) performed PMBV to 30 patients with LAA thrombus and observed no systemic thromboembolic events. In this study, Manjunath et al. (4) grouped LA thrombus into five subtypes. In patients with type 1a (LAA thrombus confined to appendage), type 1b (LA appendage thrombus protruding into LA cavity) and type 2a (LA roof thrombus limited to a plane above the plane of fossa ovalis) thrombus, PMBV was considered as safe and effective with modified techniques. Shaw et al. (6) performed PMBV to 28 patients with LAA thrombus and none of the patients experienced embolic event. Our patients had type 1a thrombus according to the classification denoted by Manjunath et al. (4). The procedures were performed with lower septal puncture and with less manipulation. Contrary to previous researchers, we used TTE.

**Figure 3. Transesophageal echocardiography shows LAA thrombus at mid-esophageal aortic valve short-axis view (Case 2). Arrow-thrombus in LAA**

LA - left atrium, LAA - left atrial appendage, RA - right atrium

**Conclusion**

PMBV is a safe option for patients with suitable valve anatomy and thrombus localized to LAA. Systemic thromboembolism is rare if performed by an experienced operator. TTE seems as a safe and effective alternative to TEE.

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The role of two-dimensional speckle-tracking echocardiography in a patient with Behçet's disease

**Behçet hastalığı olan hastada iki boyutlu benecek takip yöntemi ekokardiyografinin önemi**

**Introduction**

Behçet’s disease (BD) is a systemic inflammatory disorder of unknown origin characterized by variable clinical manifestations. Most
common forms of clinical presentation are recurrent oral and genital aphthous ulceration, uveitis and erythema nodosum (1). Cardiac involvement is relatively uncommon (2) and may present in the form of pericarditis, conduction system disturbances, coronary arteritis, intracavitary thrombosis, endomyocardial fibrosis or valvular insufficiency (3). We reported a case of 24-year-old man with right ventricular thrombus and endomyocardial fibrosis diagnosed with BD during routine echocardiography examination.

Case Report

A 24-year-old man, with a family history of sudden cardiac death of a brother at age of 26, was referred to our institution for cardiovascular investigation. He had no complained on admission. He was not a smoker. Past medical history was unenlightening except recurrent aphthous stomatitis and tonsillitis. Clinical examination detected 2/6 mid-systolic murmur over the left second space. He was a febrile and his blood pressure was 110/76 mmHg with a heart rate of 95. A pedunculated homogenous mass (16x20 mm) mobile with tricuspid valve suspected to be a thrombus in the right ventricle (RV) and increased thickness and echogenity (Fig. 1A, Video 1. See corresponding video/movie images at www.anakarder.com) of right ventricular free wall and slightly dilated main pulmonary artery (3.2 cm) with normal estimated pulmonary artery pressure (Fig. 1B) were found on echocardiographic examination. Speckle tracking echocardiographic evaluation of right ventricular free wall revealed decreased regional longitudinal strain value of -10 (Fig. 2) (4). CT pulmonary angiography pointed out pulmonary arterial aneurysms. Coagulation studies and immune indexes including lupus anticoagulant were normal. Serologic investigation did not yield any evidence of bacterial or fungal infections. On further investigation, genital aphthous ulcerations, positive pathergy test and positive human leukocyte antigen B51 (HLA-B51) serologic typing were detected. According to international study group criteria for BD (5), he was diagnosed with BD. Bright echoes, increased thickness and decreased longitudinal strain and strain rate in the right ventricular free wall were thought to be result of endomyofibrosis secondary to BD. A regimen of pulse methylprednisolone and monthly intravenous cyclophosphamide infusion combined with oral colchicines was initiated. Concomitantly, we started unfractionated heparin infusion and activated partial thromboplastin time was maintained 1.5 to 2.5 times the control value. After two weeks of therapy, there was complete resolution of right ventricular thrombus on echocardiography. At 3 months’ follow-up, he was doing well on the same therapy.

Discussion

Behçet’s disease, currently classified as a vasculitis, is a systemic inflammatory disorder first described in 1937 as a triple symptom complex of aphthae, genital ulcers, and hypopyon uveitis (6). Although mucocutaneous features such as recurrent oral ulceration are the most common presenting symptoms, BD lacks pathognomonic clinical or laboratorial findings and the diagnosis is made on the basis of a group of clinical features.

Cardiac involvement of BD is uncommon and found in 7-46% of the patients in clinical series (3). BD usually involves the right side of the heart. Endomyocardial fibrosis is a sequel of vasculitis that involves the endocardium, myocardium, or both, and has a tendency toward bacteerial endocarditis or intraventricular thrombosis. It is seen as diffuse, bright, thickened endocardium on echocardiography and can cause segmental wall motion abnormalities. Two-dimensional speckle tracking echocardiography (STE) allows the study of regional myocardial deformation expressed by strain and strain rate. Subtle cardiac involvement of various diseases can be detected by STE reliably. Moreover, it has been proven that; peak regional longitudinal strain and strain rate values are significantly lower in fibrotic areas than normal myocardium (7). Decreased strain and strain rate values are clues to myocardial or endomyocardial involvement of the disease process. Yaşmur et al. (8) have proven that left ventricular longitudinal systolic strain is significantly impaired and NT-proBNP is increased in correlation with mean left ventricular longitudinal strain in patients with BD compared with the healthy controls.

Conclusion

Speckle tracking echocardiography, a novel non-invasive method, can be used for the assessment of subclinical right ventricular dysfunction in BD. Further clinical studies are needed to define the eventual role of STE in the determination of RV functions in BD.

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Video 1. Views of a pedunculated homogenous mass (16x20 mm) mobile with tricuspid valve suspected to be a thrombus in the right ventricle (RV) with increased thickness and echogenity
Aneurysm of ascending and descending aorta in a 10-year-old boy with Wiskott-Aldrich syndrome

Wiskott–Aldrich syndrome (WAS) is a recessive genetic disorder linked to the X-chromosome characterized by immune deficiency, eczema and thrombocytopenia. To the best of our knowledge, a few cases of vasculitis or aneurysmal formation have been reported in this syndrome, but the association has not been well established (1-6). We report a patient with WAS and extensive aortitis causing severe aneurysmal dilatation in the everywhere of the aorta who underwent suc- cessful first stage operation involving replacement of ascending aorta.

Case Report

A 10-year-old boy had been followed up with the diagnosis of chronic idiopathic thrombocytopenic purpura (ITP) for 2 years. There was no history of eczema and repeating infections suggesting immunodeficiency. Because of persistent caught in the last two months, chest X-ray was performed and it suggested an ascending aortic aneurysm (Fig. 1). Echocardiography showed mild aortic regurgitation without aortic stenosis and aneurysmatic dilatation of ascending and descending aorta. 3D computed tomography revealed dilatation of the ascending, descending and abdominal aorta with extreme calcification and plaque (Fig. 1). He had two maternal cousins with the diagnosis of WAS. One of them had died with septicemia after splenectomy. There was no history of aneurysm in his cousins. Flow cytometry showed absence of WAS protein. We planned two-stage operation beginning with replacement of the ascending aorta due to high risk of rupture. Pulse methyl-prednisolone therapy (30 mg/kg/day for 3 days) was tried to correct the thrombocytopenia prior to cardiac surgery. The platelet count increased from 45,000 to 103,000/mm². 0.5 gr/kg IVIG was administered monthly. Apheresis thrombocytes were also given before the operation. He underwent valve sparing surgery. Vascutek® graft (24 mm) was replaced to aortic root and ascending aorta (Fig. 2). The luminal surface of the aorta was found to be covered by ulcerated and calcified necrotic plaques. Postoperative recovery was uneventful and no excessive