Uncorrected Tetralogy of Fallot in a 51-Year-Old Patient

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ABSTRACT
Tetralogy of Fallot (TOF) is the most common cyanotic congenital heart disease. This report describes the case of a fifty-one year old man who presents with an exertional dyspnea and polycythemia. Echocardiography showed perimembranous ventricular septal defect, hypertrophy of the right ventricle, over-riding of the aorta and stenosis of the right ventricular outflow tract.

Key words: Tetralogy of Fallot, uncorrected, polycythemia

INTRODUCTION
Tetralogy of Fallot (TOF) is the most common cyanotic congenital heart disease, with 10% of all congenital heart malformations (1). Tetralogy of Fallot includes four major component which are ventricular septal defect (VSD), overriding aorta, right ventricle outflow tract obstruction and right ventricle hypertrophy (RVH) (2). At the present time, the most of these patients are diagnosed in first year of their life’s and moreover the majority of these patients are diagnosed in antenatal period, however this disorder may be occasionally recognized at adulthood (1,3). Echocardiography is an indispensable method for organizing treatment.
and diagnosis of the disease (3). Despite of the fact that TOF is a fatal disease particularly when not being treated, approximately 85% of the patients are able to survive up to adulthood after surgical correction (4).

We have reported a TOF patient who was not diagnosed and was not treated up to 51 years old.

CASE

A fifty-one year old man was referred to the hematology department of our university due to his polycythemia. On his medical history, he had mild to moderate exertional dyspnea for approximately ten years and phlebotomy had been performed several times.

Initial clinical assessment designated a well nourished individual with central cyanosis. His blood pressure was 110/80 mmHg, pulse 92 beats per minute and respiratory rate of 18 breaths per minute. His oxygen saturation was 68% under room air. Auscultation exposed a loud first heart sound with systolic ejection murmur at the left sternal border. The lungs were clear. Electrocardiogram (ECG) showed sinus rhythm with RVH and right bundle branch block (RBBB). Polycythemia was detected by hemoglobin of 21.4 g/dl and hematocrit of 61.1%.

The transthoracic echocardiography was applied. Perimembranous ventricular septal defect (Figure 1), hypertrophy of the right ventricle, over-riding of the aorta and valvular stenosis of the right ventricular outflow tract with a pressure gradient of 73 mmHg were obtained (Figure 2). Left ventricular dimension and functions was normal. Operation couldn’t have been performed because of the patient’s refuse.

DISCUSSION

Tetralogy of Fallot is generally presented with cyanosis at childhood whereas dyspnea, exercise intolerance and cyanosis are mainly occurred at adulthood (5). The cause of cyanosis seen in earlier time is the shunt being from right to left. The shunt becomes from left to right in some patient when resistance of right ventricular outflow tract is lower than left system and thus cyanosis will not be seen and the patient will be diagnosed in a later period consequently (1). It was shown in the previous studies that surgical total correction in an earlier period is the best treatment option. The most patients not being treated are to die probably at childhood (1). Pulmonary hemorrhagia, brain abscess and thromboembolic complications are the most prominent causes of death (4). Only 1% of the patients not being treated can survive up to 50 years old. However the survival rate for those being operated is about 74% (2,5). The oldest
uncorrected TOF in the literature aged 86 was reported from USA (6). In our country, the oldest uncorrected TOF reported by Yokusoglu et al. was 68 years old (2).

In conclusion, our patient was asymptomatic for a long time and the diagnosis were made while polycythemia was being investigated. Even though there are some TFO cases diagnosed at older age in litterateur, this case among the others reported in our country is one of the oldest patients not being diagnosed (1,5,7).

REFERENCES


