Rheumatic heart disease with triple valve involvement

**ABSTRACT**
Acute rheumatic fever (ARF) is a postinfectious, nonsuppurative sequela of pharyngeal infection caused by Streptococcus pyogenes, or Group A β hemolytic Streptococcus (GABHS). Of the associated symptoms, only damage to the heart’s valvular tissue, or rheumatic heart disease (RHD), can become a chronic condition leading to congestive heart failure, stroke, endocarditis, and death. ARF is the most common cause of cardiac disease in children in developing countries. A joint meeting of the World Health Organization and the International Society estimated that 12 million people in developing countries were affected by acute rheumatic fever and rheumatic heart disease, with the majority of these being children. This level of morbidity is comparable to that in developed countries’ in the last century, before the introduction of penicillin. (1) RHD is the most common cause of mitral stenosis (MS) in adults (2) but it is also the most common cause of multivalvular disease. About 60% of patients with rheumatic MS have concurrent mild-to-moderate aortic insufficiency (AI), and 10% of patients with rheumatic MS have concurrent severe AI. (3) The presence of concomitant AI may augment the volume of mitral regurgitation (MR) flow leading to left ventricle (LV) enlargement. The presence of both severe MR and severe AI is poorly tolerated. Significant trivalvular disease involving mitral, aortic and tricuspid valves is uncommon. Although rare, trivalvular disease has been described in the literature. Clinical and hemodynamic manifestations depend on the severity of each lesion. We reported this case because of the rare presentation of an uncommon disorder and to highlight the fact that the presence of trivalvular disease can be difficult to diagnose, even for a trained physician.

**Key words**: rheumatic heart disease, triple valve disease, rheumatic valve disease

**Introduction**
Acute rheumatic fever (ARF) is a postinfectious, nonsuppurative sequela of pharyngeal infection caused by Streptococcus pyogenes, or Group A β hemolytic Streptococcus (GABHS). Of the associated symptoms, only damage to the heart’s valvular tissue, or rheumatic heart disease (RHD), can become a chronic condition. A joint meeting of the World Health Organization estimated that 12 million people in developing countries were affected by acute rheumatic fever and rheumatic heart disease, with the majority of these being children. This level of morbidity is comparable to that in developed countries’ in the last century, before the introduction of penicillin. (1) RHD is the most common cause of mitral stenosis (MS) in adults in developing countries (2) but it is also the most common cause of multivalvular disease. About 60% of patients with rheumatic MS have concurrent mild-to-moderate aortic insufficiency (AI), and 10% of patients with rheumatic MS have concurrent severe AI. (3) The presence of concomitant AI may augment the volume of mitral regurgitation (MR) flow leading to left ventricle (LV) enlargement. The presence of both severe MR and severe AI is poorly tolerated. Significant trivalvular disease involving mitral, aortic and tricuspid valves is uncommon. Although rare, trivalvular disease has been described in the literature. We reported this case because of the rare presentation of an uncommon disorder and to highlight the fact that the presence of trivalvular disease can be difficult to diagnose, even for a trained physician.

**Case report**
A 19-year-old Senegalese female presented to our Emergency Department in April 2013 complaining of dyspno-
ea and severe hypotension. Her vital signs included an undetectable blood pressure, heart rate of 115 beats/min and oxygen saturation of 100% while receiving 9 l/min oxygen via a simple face mask. On examination, she had severe mental deterioration, heart sounds were dysrhythmic and tachycardic but without murmur, bruit or rubs, and her lungs demonstrated a diminished vesicular murmur bilaterally. Hemoglobin was 11.9 g/dl and hematocrit was 34.8%. Her 12 lead electrocardiogram showed atrial fibrillation with a prominent R and S wave in all leads. After surgical reparation of both the aortic and mitral valve and substitution of the tricuspid valve, the patient experienced three episodes of paroxysmal supraventricular tachycardia. Each episode ended spontaneously. The patient was immediately treated with high doses of diuretics (furosemide 160 mg per day and potassium canrenoate 50 mg per day), oral anticoagulant (acenocoumarol 4 mg per day), nitrates (nitroglycerine 60 mg per day) and carbonic anhydrase inhibitor (acetazolamide 500 mg per day). Two months after surgery, echocardiography demonstrated the absence of tricuspid insufficiency, mild AI, mild mitral insufficiency, normalization of left ventricular function with an ejection fraction of 55%. Final oral therapy included diuretics (furosemide 75 mg per day and potassium canrenoate 25 mg per day), ACE inhibitor (ramipril 5 mg per day), beta blocker (bisoprolol 1.25 mg per day), oral anticoagulant (acenocoumarol 1 mg per day) and proton pump inhibitor (omeprazole 20 mg per day).

Discussion
Pathophysiology
The role of GABHS infection as an initial event is clear, and has been supported by outbreaks of ARF following pharyngitis, as well as rising antistreptococcal antibodies in patients with ARF. (4) ARF is believed to be an autoimmune reaction to GABHS infection. In 1945 antihypercardiac autoantibodies were identified. These antibodies could damage host tissue without the continued presence of streptococci. In 1962, cross-reactivity between human heart tissue and GABHS was demonstrated. This cross-reactivity is believed to occur through molecular mimicry: in ARF and RHD, the foreign antigen is the M-protein and it cross reacts with cardiac myosin. Although all four heart valves can be involved in rheumatic carditis, there is marked predominance of mitral valve involvement. The reason for this prevalence is yet unknown. Once heart valves have been damaged, blood is allowed to regurgitate. This can cause acute congestive heart failure, which can be the presenting symptom of ARF. Patients that recover from initial rheumatic carditis are likely to have permanent valve damage in the form of stenosis or regurgitation. RHD is the most common cause of MS. MS typically induces a pressure overload in the left atrium, which leads to progressive atrial dilatation and, finally, to the creation of a profibrillatory environment in the left atrium that manifests clinically as atrial fibrillation. The other main features of MS are the development of pulmonary hypertension (5) and tricuspid regurgitation (6). More than one third of patients with MS have been reported to have at least moderate TR. (7) Atrial fibrillation is a common dysrhythmia in

Figure 1. Chest radiography demonstrating cardiomegaly.
patients with MS (8) and predisposes a patient to the development of TR.

Diagnosis

The diagnostic criteria for ARF were first developed by Jones in 1944 (9) and have been modified several times. The criteria are divided into Major and Minor criteria. Diagnosis is made by the presence of either 2 Major or 1 Major and 2 Minor criteria, plus evidence of recent streptococcal infection, either by a pharyngeal swab culture positive for GABHS or positive rapid GABHS antigen test.

Therapy

Patients with known chronic valvular RHD are managed individually according to lesion severity, location, and number of valves involved. Nonsurgical recommendations include restriction of physical exertion in symptomatic patients to reduce cardiovascular load and prophylactic antibiotics to prevent endocarditis. Medical therapy for patients with MS aims to reduce heart rate and decrease blood volume with beta-blockers, cautious use of oral diuretics, and even digoxin if atrial fibrillation is present. Significant trivalvular disease involving mitral, aortic, and tricuspid valves is uncommon. Yet, for those presenting with advanced heart failure it is recommended that all three valvular lesions should be surgically corrected.

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Conclusion

RHD is an uncommon pathology in developed countries, for this reason we chose to report what we saw. This manuscript aims to remind all clinicians that sooner or later they may come across a forgotten disease.

REFERENCES