Abstract
Primary lesions of the tricuspid valve are rare. Tricuspid disease is an uncommon valvular abnormality that most commonly occurs in association with other valvular lesions, particularly in patients with rheumatic heart disease. For many years, surgical commissurotomy and valvuloplasty were the only available methods to correct tricuspid disease. We present a case of post rheumatic disease of the tricuspid valve in a young girl, associated to a left atrial mass.

Keywords: Tricuspid valve; Stenosis; Girl

Case Report
Miss J is an 18 years old young girl coming for the first time to consult in the Cardiac Centre Shisong presenting with shortness of breath on mild exertion, cough, abdominal swelling and mild lower limbs swelling. She was referred by a general practitioner who heard a murmur.

In the history she agreed repetitive tonsillitis with fever and joint pains. Also has been regular to the dentist for teeth problems.

On physical examination she was very ill looking, afebrile, no pallor, mild peripheral cyanosis, good orientation in time and space. S1S2 were regular with a systolic murmur 2/6 and diastolic murmur 3/6 at the tricuspid area, a systolic murmur 2/6 at the mitral area. The lungs were clear with good entry bilaterally, soft abdomen, not tender mild hepatomegaly, lower limbs with mild pitting oedema.

The cardiac echocardiography showed
Dilated inferior vena cava-18mm, dilated right atrium 87 *63 mm with an area of 51,17 cm² and a circumference of 27 mm, the anterior and posterior leaflets were calcified and fused (Figure 1A), giving a severe stenosis with a peak gradient of 22,5 mmHg, the mean gradient was 9,9 mmHg, and pressure half time of 218 ms (Figure 1B). The tricuspid annulus was 57 mm (Figure 1D). The color continuous Doppler through the tricuspid valve gives also a mild regurgitation with a vena contracta-8 mm, we could see also the retraction of the subvalvular apparatus. The right ventricle is dilated but the systolic function was not compromised with a transannular annular plane systolic excursion-21mm. There was right ventricular systolic overload.

The left atrium 69 *48 mm with an area of 51,17 cm² and a circumference of 21 mm is having at the apex near the entrance of the inferior pulmonary veins a mass homogenous well organised looking like a myxoma having a circumference of 11,1 mm, with a long axis of 44 mm and a short axis-18,5 mm (Figure 1C). The mitral leaflets are mildly calcified with good function. The annulus is not dilated, 29 mm. The left ventricle is not dilated with a paradoxal septum and acceptable systolic function.

The laboratory analysis was within the normal limits: Rapid HIV testing was negative. WBC: 4600 cells/mm³, a hemoglobin of 12.3 g/dl, Serum creat: 0.9 mg/dl, K: 3 mmol/l.

In the trans-oesophageal echocardiography, the mass was having the aspect of a myxoma. The patient could not afford the surgery.
The patient was to undergo an open heart surgery with extracorporeal circulation for the plasty/replacement of the tricuspid valve and the excision of the mass and the confirmation of the diagnosis. Tricuspid plasty was done successfully as well as the excision of the mass which had a benign etiology.

**Discussion**

Tricuspid disease, most of which is of rheumatic origin, is rarely seen in developed countries, although it is still found in developing countries. The majority of cases of tricuspid rheumatic disease present with tricuspid regurgitation or a combination of regurgitation and stenosis [1,2]. The case described above was a post rheumatic tricuspid valve disease, with calcified and fused tricuspid leaflets, restriction motion and retracted subvalvular apparatus. Other causes of obstruction to right atrial emptying are unusual and include congenital tricuspid atresia, right atrial tumors, which can produce a clinical picture suggesting rapidly progressive TS and carcinoid syndrome [3]. Tricuspid disease and mainly stenosis has been also described in some cases due to lead endocarditis. It can be a chronic complication of pacemaker implantation that may occur more often than it is clinically suspected and may become even more common due to the growing number of cardiac devices currently implanted. The diagnosis should always be considered in patients with right heart failure and electrodes present in the right heart chambers [4,5]. Surgical correction was the treatment of choice for pulmonary stenosis until three decades ago, when balloon valvuloplasty was implemented; nevertheless, percutaneous tricuspid valvuloplasty may be a therapeutic alternative to surgery in cases of TS alone without any concomitant valvulopathy [6,7]. In our case surgical procedure was the procedure of choice since balloon valvuloplasty is not yet practices, with excision of the left atrial mass. Tricuspid plasty was done with good outcome as well as the excision of the mass which was benign.

Myxoma is the main differential diagnosis suggested for our patient as we know that this kind of tumor is located in 90% on left atrium. It is in general well circumscribed and has an oval form [8]. The magnetic resonance imaging has an important contribution in the differentiation of thrombus from myxoma. After gadolinium injection, myxoma is lifting up on periphery whereas thrombus remains unchanged. However, thrombus could also lift up on periphery when it is organized [9]. The surgical procedure could be done in this case with a very good outcome, in contradistinction of the huge amount of cases with indication of open heart surgery that die because of financial limitations. This is the tip of the iceberg in countries with financial limitations where patients are poor and cannot afford open heart surgery. It urges our government and stakeholders to find sustainable solutions to save lives of citizens.

**Conclusion**

Tricuspid disease associated to left atrial mass is a rare entity. Surgical correction of the disease is the procedure of choice taking into consideration the clarification of the mass diagnosis.

**Conflict of Interest**

None.

**Acknowledgement**

None.

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