



Infective Endocarditis on Ventricular Septal Defect Complicated by Septic Pulmonary Embolism: Case Report

**Bettar Mohamed Ghouleme ^{a*}, Charfo B Mahamdo ^a,
Mullendelle Mayanga Patrick ^a, Nji Maleck ^a,
Haboub Meryeme ^a, Arous Salim ^a, Ghali Bennouna ^a
and Rachida Habbal ^a**

^a *Department of Cardiology, Faculty of Medicine and Pharmacy of Casablanca, Ibn Rochd University Hospital, Morocco.*

Authors' contributions

This work was carried out in collaboration among all authors. All authors read and approved the final manuscript.

Article Information

DOI: 10.9734/CA/2024/v13i1401

Open Peer Review History:

This journal follows the Advanced Open Peer Review policy. Identity of the Reviewers, Editor(s) and additional Reviewers, peer review comments, different versions of the manuscript, comments of the editors, etc are available here: <https://www.sdiarticle5.com/review-history/115045>

Case Report

Received: 25/01/2024

Accepted: 30/03/2024

Published: 02/04/2024

ABSTRACT

Ventricular septal defect (VSD) is a congenital heart disease most likely to cause infective endocarditis. Pulmonary embolism constitutes one of its main complications. Treatment is mostly based on effective antibiotic therapy but can sometimes require additional surgical intervention. We report a case of infectious endocarditis in a 20-year-old patient followed for congenital heart disease in the form of a ventricular septal defect. This was complicated by septic pulmonary embolism due to a right heart vegetation, with a good clinical-biological resolution under medical treatment only.

*Corresponding author;

Keywords: *Infectious endocarditis; ventricular septal defect; pulmonary embolism.*

1. INTRODUCTION

Infectious endocarditis (IE) is a serious condition causing significant morbidity and mortality. Among its risk factors are congenital heart diseases, with VSD being the most frequent [1]. Patients with VSD have an incidence of 1.67/1000 patient per year, or 11 to 15 times greater risk than the general population [2]. Septic pulmonary embolism represents a rare complication, thereby increasing the mortality risk and necessitating appropriate care.

2. CASE PRESENTATION

This is a 20-year-old patient with a history of VSD-type congenital heart disease who consulted for dyspnea at rest associated with subacute onset fever that has been ongoing for 2 months. He was hemodynamically stable blood pressure 111/61 mmHg, and a temperature of 39.6°C, ecchymotic purpura in the lower limbs,. A holosystolic murmur was noted at the left sternal border during heart auscultation; the rest of the clinical examination was unremarkable.

Electrocardiogram (ECG) showed a regular sinus rhythm at 107 beats per minute (bpm),

fixed PR interval at 120ms, normal heart axis, narrow QRS wave, without repolarization disorders. Transthoracic echocardiography showed good bi-ventricular function, the atria were not dilated, no mitral or aortic valvulopathy, presence of a peri-membranous VSD of 11mm long axis with restrictive left-right shunt (Fig. 1), with a maximum gradient across the VSD at 88mmHg. At the VSD site there was a hyperechoic filiform structure measuring 23x12 mm on the lower border of the VSD floating in the right ventricle (Fig. 2). The inferior vena cava (IVC) was not dilated (11 mm) and compliant and there was no pericardial effusion. Faced with this clinical and echocardiographic picture, the diagnosis of infective endocarditis was made.

Biologically: leukocytosis at 18030 G/L with predominantly PNN (14170 G/L), C-reactive protein (CRP): 340g/L, renal and hepatic function as well as the hemostasis assessment were normal whereas the 2 blood cultures were positive for *Staphylococcus aureus*. As part of the assessment, a chest angio-CT scan was carried out and showed the presence of a bilateral distal pulmonary embolism strongly suggestive of septic emboli (Fig. 3); the fundus examination was normal.

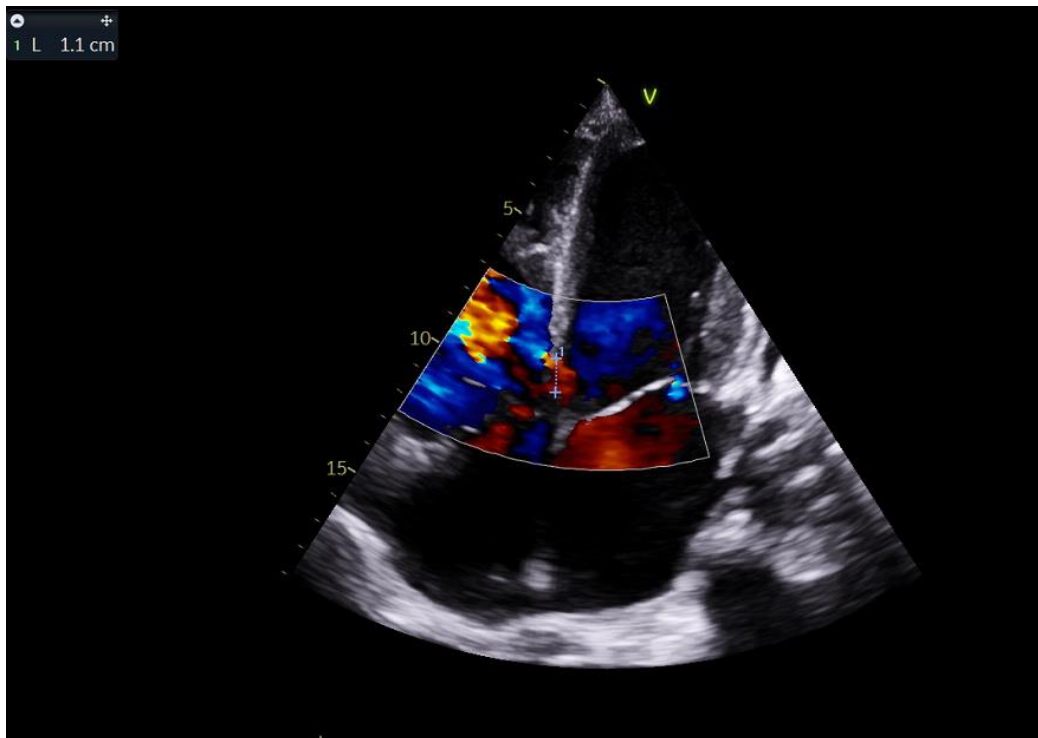


Fig. 1. Apical four-cavity cut showing the septal defect (VSD)

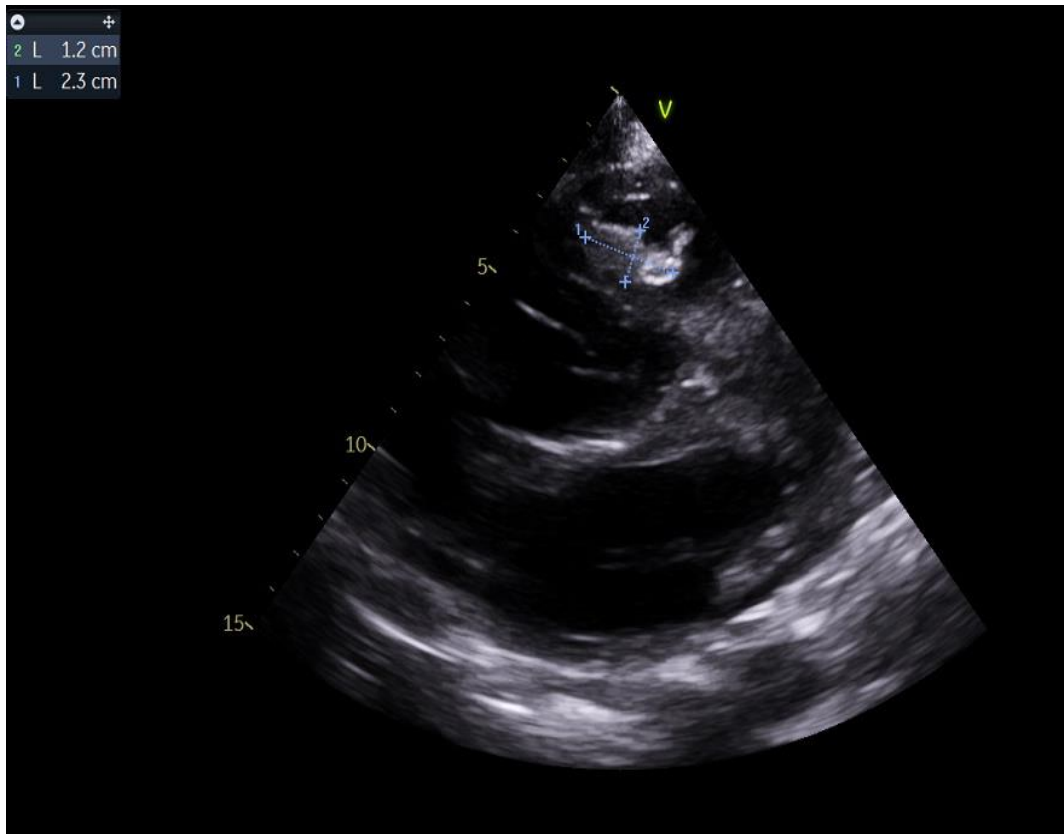


Fig. 2. Modified parasternal long axis view showing a hyperechoic filiform image arising on the IVS and floating in the RV (vegetation image)



Fig. 3. CT image showing distal pulmonary embolization

The port of entry was probably dental given the patient's poor oral condition. He was put on dual antibiotic therapy using ceftriaxone 2g/day and gentamycin 160 mg/day with good clinical-biological. On follow-up clinical exam the patient was afebrile (T°: 36.8°C), saturation of 98% in open air, with negative CRP value and normal white blood count.

A surgical closure of the VSD was considered taking into account the size of the vegetation and the pulmonary vascular resistance on echocardiography findings which was equals to 1 m/s (Vmax of the TR/pulmonary ITV). The patient was transferred to the cardiovascular surgery department for surgical intervention.

3. DISCUSSION

Patients with left-right shunts such as VSD present a risk of IE. Indeed, several studies have documented this risk of developing IE in particular on the right side [3,4] due to the high pressure gradient between the two cardiac chambers, source of endocardial erosive lesions on the right side, are generally the site of vegetations.

Active search for pulmonary embolism is always recommended in patients with endocarditis, particularly in patients with large (> 10 mm) and mobile vegetations. This link between IE and septic pulmonary emboli is well documented in the literature [5,6]. Our patient had septic bilateral distal vegetation embolization which persisted despite appropriate antibiotic therapy [7,8].

Intravenous antibiotic therapy is the mainstay of treatment for IE, and they typically respond to a 4- to 6-week course of parenteral antibiotics [9,10].

Surgical intervention in right-sided IE is indicated only when caused by difficult-to-eradicate microorganisms, e.g. fungus, bacteremia persisting for more than 7 days despite adequate antibiotic therapy, embolisms recurrent pulmonary heart failure with or without concomitant right heart failure, perivalvular abscess, persistent large tricuspid valve vegetation (>20 mm), or right heart failure secondary to severe tricuspid regurgitation [11,12,13].

4. CONCLUSION

Septic pulmonary embolism represents an important complication of endocarditis secondary

to VSD. A systematic workout should be carried out in all patients with right heart endocarditis, especially if the vegetation is large and mobile and treated early. Infective endocarditis is a serious condition and must be looked for in the face of any unexplained fever, particularly in patients with congenital heart disease, especially VSD. Echocardiography plays an essential role in the positive diagnosis, monitoring as well as the search for complications and the study of congenital heart disease.

CONSENT

As per international standards or university standards, patient(s) written consent has been collected and preserved by the author(s).

ETHICAL APPROVAL

It is not applicable.

COMPETING INTERESTS

Authors have declared that no competing interests exist.

REFERENCES

1. Sattwika PD, Hartopo AB, Anggrahini DW, Mumpuni H, Dinarti LK. Right-sided infective endocarditis in patients with uncorrected ventricular septal defect and patent ductus arteriosus: Two case reports. *Clin Case Rep.* 2018;6(11):29.
2. Lee PT, Uy FM, Foo JS, Tan JL. Increased incidence of infective endocarditis in patients with ventricular septal defect. *Congenit Heart Dis.* 2018;13(6).
3. Kim YJ, Sohn DW. Pulmonary valve endocarditis with septic pulmonary thromboembolism in a patient with ventricular septal defect. Sohn DW. Pulmonary valve endocarditis with septic pulmonary thromboembolism in a patient with ventricular septal defect. Sohn DW. Pulmonary valve endocarditis with septic pulmonary thromboembolism in a patient with ventricular septal defect. *J Cardiovascular Ultrasound.* 2009; 17(4): 138-40.
4. Zijlstra F, Fioretti P, Roelandt JR. Echocardiographic evidence of free-wall vegetative endocarditis complicated by pulmonary embolism in a patient with a ventricular septal defect. *Br Heart J.* 1986;55(5):497-9.

5. Teran CG, Antezana AO, Salvani J, Abaitey D. Group B streptococcal endocarditis associated with multiple septic pulmonary emboli. Group B streptococcal endocarditis associated with multiple septic pulmonary emboli. Group B streptococcal endocarditis associated with multiple septic pulmonary emboli. *Practical Clin.* 2011;1(1).
6. Nakauchi Y, Taniguchi M, Miyamura Y, Ishise T, Miyazaki S. [Pulmonary septic embolism with right side infectious endocarditis and ventricular septal defect: A case report]. *J Cardiol.* 2007;50(6): 383-7.
7. Roodpeyma S. Infective endocarditis complicated by septic pulmonary embolism in a case of ventricular septal defect. *J Compr Ped.* 2015;6:e29610.
8. Aydin MS, Hazar A, Demirkol AH. Massive right main pulmonary embolism caused by infective endocarditis of the tricuspid valve. *Heart Asia.* 2013;5: 128–129.
9. Sutcliffe EC, Terasaki GS, Thompson RE. Tricuspid endocarditis with pulmonary embolism. *RespirCare.* 2006;51:1471–1474.
10. Ishak Ahmed Abdi, Abdirahim Ali Adan Nur, Abdirahman Duale. A case of infective endocarditis and pulmonary; 2022.
11. Shmueli H, Thomas F, Flint N, Setia G, Janjic A, Siegel RJ. Right-sided infective endocarditis 2020: challenges and updates in diagnosis and treatment. *J Am Heart Assoc.* 2020;9:e017293.
12. Park HE, Cho GY, Kim HK, Kim YJ, Sohn DW. Pulmonary valve endocarditis with septic pulmonary thromboembolism in a patient with ventricular septal defect. *Cardiovascular ultrasound J.* 2009; 17: 138–140.
13. Saleem M, Ahmed F, Patel K, Munir MB, Ghaffar YA, Mujahid H, Balla S. Isolated pulmonary valve endocarditis: Case report and review of existing literature on diagnosis and therapy. *CAS (Phila).* 2019; 3:227–230.

© Copyright (2024): Author(s). The licensee is the journal publisher. This is an Open Access article distributed under the terms of the Creative Commons Attribution License (<http://creativecommons.org/licenses/by/4.0>), which permits unrestricted use, distribution, and reproduction in any medium, provided the original work is properly cited.

Peer-review history:

The peer review history for this paper can be accessed here:
<https://www.sdiarticle5.com/review-history/115045>