Case Report

Late bacterial endocarditis of an Amplatzer atrial septal device

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Abstract

A 59-year-old male with an secundum atrial septal defect status post repair with an Amplatzer occluder in 2001 was admitted with sepsis and MRSA bacteremia. Transesophageal Echocardiography (TEE) showed presence of an overlying mobile echogenic structure on the left atrial surface of the device suggestive of a vegetation/infected thrombus. This is only the 3rd case description of late endocarditis involving the Amplatzer ASD closure device in an adult.

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1. Case description

A 59-year-old male with a history of hypertension, type 2 diabetes, Congestive heart failure s/p Implantable Cardioverter Defibrillator and a Secundum ASD s/p percutaneous repair with an Amplatzer Atrial Septal Occluder Device (ASO; AGA Medical Corporation, Golden Valley, Minnesota) was admitted with sepsis and positive blood cultures for MRSA. Bacteremia persisted despite IV Vancomycin. With a high suspicion for Infective endocarditis (IE) despite normal Transthoracic echocardiography, a TEE was done. It showed a highly mobile echogenic mass suggestive of vegetation/infected thrombus in the left atrial surface of the device Figs. 1 and 2, Videos 1 and 2. After initiation of IV daptomycin, there was significant clinical improvement leading to discharge with 6 weeks of IV antibiotic.

2. Comments

The Amplatzer septal occluder is one of the most frequently used percutaneous devices for the closure of Secundum ASD. Complications of this procedure include device embolization/malposition, arrhythmias, cardiac perforation, thrombus formation and device erosion with infection being the least common.

Only 2 previous cases of late infection of occluder device have been reported in literature. A case of IE involving a CardioSEAL device for patent foramen ovale has also been
Device infection can occur in two ways; either through introduction of microbes during the procedure or secondary to seeding of microorganisms at a later time. After device implantation, it is thought that it takes 6 months for complete neoendothelialization. The current guidelines recommend antibiotic prophylaxis for 6 months following device placement. The organism isolated in this case is classical for nosocomial infection, likely from infection from an abdominal wound. In late IE involving ASD closure device medical management with prolonged antibiotic therapy with constant blood culture monitoring might be adequate in the absence of device dehiscence, septal perforation or fistula formation.

Conflicts of interest

All authors have none to declare.

REFERENCES