

Figure 1. Embolization of Amplatzer atrial septum defect occluder device into the main pulmonary artery

ostium secundum (OS) ASD measuring 22 mm on the transthoracic echocardiography (TEE) was underwent transcatheter closure with a 30 mm Amplatzer device. The procedure was failed due to dislodgment of the device into MPA (Figure 1). The retrieval of the dislodged device was attempted, but faced with failure since the guidewire got entangled into the chords of the tricuspid valve. Due to hemodynamically instability, emergency surgical intervention was advised to retrieve the device.

On surgery, a large secundum ASD (30×40 mm) was found with no tissue “shelf” at the inferior margin. The device was covered with fresh thrombus and embolized entirely into the main pulmonary artery. It was retrieved without injury to main pulmonary artery, and entangled part of the guidewire was removed through femoral route. Subsequently, the ASD closed with an autologous pericardial patch. The patient had an uneventful recovery and discharged a week later. He remained well during the four-month follow-up.

Case 2

A 48-year-old male with OS-ASD measuring 24 mm on TEE with adequate rims was taken for device closure. The transcatheter closure was attempted; however, the device closure failed and dislodged into the right ventricular outflow tract. The patient became hemodynamically unstable, shifted to the operation theatre. On operation, the right atrium opened longitudinally, and the Amplatzer device was retrieved from the right ventricular outflow tract (Figure 2). Subsequently, the ostium secundum ASD (25×15 mm) was closed with fresh autologous pericardial patch. The patient had an uneventful recovery and was discharged a week later. He remained well during the one-year follow-up.

Case 3

An 18-month-old female infant with perimem-

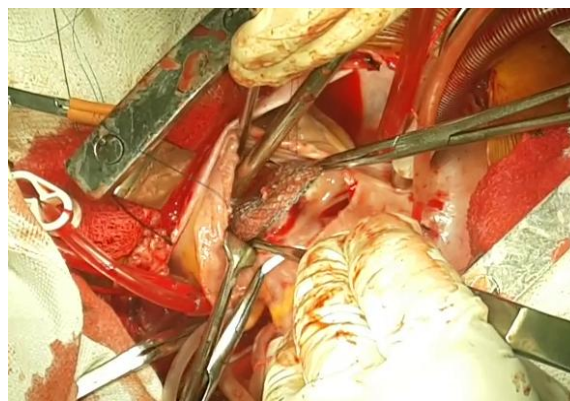


Figure 2. Removal of Amplatzer device from the right ventricular outflow tract

branous and subaortic restrictive VSD (4 mm) with left to right shunt on the TTE was underwent for transcatheter closure. The transcatheter closure was performed under fluoroscopic control. As the device deployed, it embolized into the right pulmonary artery (Figure 3). The patient was shifted to the operation room, and the median sternotomy was performed. The right pulmonary artery was dissected till lobar branches, no device felt externally by the finger. When the right pulmonary artery was opened, the device was visualized intraluminally at the lobar branch of the middle lobe. It was retrieved gently from the right pulmonary artery. Afterwards, the VSD was closed using PTFE felt patch. The patient recovered without any postoperative complications and was discharged a week later. She was well after the six-month follow-up.

Case 4

A five-year-old asymptomatic female patient with perimembranous VSD (7 mm) with left to right shunt, showing no pulmonary arterial hypertension on the TTE was taken for device closure in the cathlab. The attempted device closure was failed and the device embolized into the descending thoracic aorta below the level of the crossing left renal vein. The patient was

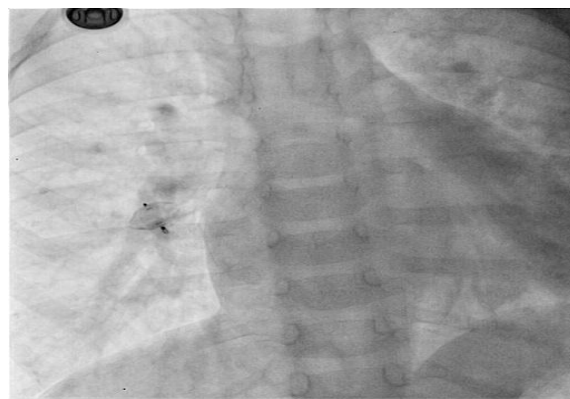


Figure 3. Embolized Amplatzer ventricular septal defect occluder device into the right pulmonary artery

thermodynamically stable and shifted to the operation room. Laprotomy was performed, and after mobilizing complete bowel, vertical aortotomy was carried out just below the level of crossing left renal vein. Therefore, the device was retrieved through aortotomy. The patient had an uneventful recovery and was discharged a week later. She was followed up, and the VSD closure was preformed surgically after six months without any postoperative complications. She was discharged a week later.

Discussion

Although the surgical repair of ASD and VSD yields extremely low morbidity and mortality, the interventional transcatheter closure for all types of cardiac defects with left to right shunting gained wide popularity. The recent studies strongly suggest that transcatheter device closure is safe method for both adult and child patients, with shorter hospital stay less mortality and morbidity rates. Moreover, this therapeutic approach obviates surgical risk factors and inevitable surgery scar. None the less, this procedure is not complications free. The reported complications of the transcatheter closure include, device malposition, residual shunt, caval thrombosis, systemic or pulmonary embolization, thromboembolism, erosion and perforation of the heart, and atrial arrhythmia (3, 4). The device embolization may cause serious damage due to embolization at unexpected sites of circulatory system. The reported embolization rates are 4% and 20% in 1991 (5) and 1996 (6), respectively, which dramatically decreased to 0.55% in 2005 (7). Out of the 81 transcatheter closures (i.e., 50, 12, and 19 ASD, VSD, and patent ductus arteriosus, respectively) performed in our institute in the last year (December 2015-December 2016), four cases (i.e., 2 ASD and 2 VSD) needed emergency surgical management due to septal occluder device embolization. All the four devices were embolized within 10 min of deploying the device.

The most common cause of device failure is improper patient or device selection. One of the the major cause of device dislodgement is size of the defect greatly exceeding the waist diameter of the device. On the other hand, erosion and perforation can occur with oversizing of device, especially when there is a defect with a deficient anterosuperior rim. The patients in whom we can not achieve satisfactory device position or significant residual defect is remaining inspite of device closure, device retrieval is usually best option at the time of implantation and subsequent elective surgical closure should be done. However, if embolization (in which transcatheter snaring and retrieval is unsuccessful) occurs, urgent surgical therapy is necessary.

Device removal should be performed mainly through the right atrium depending on the position of the embolized device. Performing right atriotomy provides a good surgical exposure to all intracardiac structures even to the left ventricle via ASD. The intracardiac structure damages at valvular and subvalvular apparatus caused by percutaneous device retrieving attempts should be repaired. Those devices, which embolize to the ascending or descending aorta should be removed via aortotomy after confirming their positions with TEE. Perioperative TEE guidance is essential particularly in these patients since the device migration might occur after aortic cannulation and initiation of cardiopulmonary bypass.

Conclusion

The device closure for congenital septal defects is less invasive than the surgery; furthermore, this technique obviates the surgical risk factors and the inevitable surgery scar. The device closure failures may be prevented by appropriate patient selection, and also proper selection of the size of the devices. However, embolization is still a major complication in interventional closure of the cardiac defects. In case of the failure of the percutaneous catheter retrieval attempts, surgical management would be the only method to remove the embolized devices. Therefore, the surgeon should be present in place and also be prepared for the removal of the embolized device.

Informed consent

Written informed consent was obtained from the patient for publication of this Case Report and any accompanying images.

Disclaimers

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Conflict of Interest

The authors declare no conflict of interest.

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