Subacute ventricular perforation is a rare complication of pacemaker or implantable-cardioverter defibrillator implantation. However, it can be life threatening. The development of small-diameter active fixation leads may be associated with increased risk for delayed right ventricular perforation. Additionally, the management of this complication has been poorly described. We report an unusual case of subacute right ventricular perforation caused by a passive fixation lead.

Introduction

Myocardial perforation is a rare complication of pacemaker or implantable-cardioverter defibrillator (ICD) implantation and usually happens at the time of lead insertion. Previous use of temporary pacemaker lead seems to increase the risk of myocardial perforation. The development of small diameter leads and active fixation mechanisms have resulted in increased stiffness at the tips of leads, potentially increasing the rate of this uncommon event. We describe an unusual case of subacute myocardial perforation caused by a passive fixation lead. In a hemodynamically stable patient, signs and symptoms suggestive of perforation are presented, as well as the therapeutic management.

Case report

A 77-year-old woman underwent a dual-chamber pacemaker implantation for episodes of sinus pauses and syncope (Zephyr XL, St Jude Medical Inc, St Paul, MN, USA). A 7-French passive fixation lead (St Jude IsoFlex S 1646-T) was inserted via the left subclavian vein approach and positioned in the right ventricular (RV) apex without any immediate complications (Figure 1A and 1B). The atrial and ventricular sensing were measured at 1.0 and 10.4 millivolts, respectively. The pacing thresholds (volts/milliseconds) were 1.0/0.5 and 0.5/0.5 in the atrial and ventricular leads, respectively. Impedances were in the normal range (atrial lead 429 ohms and ventricular lead 727 ohms). Thirty days post-implant she presented to the emergency department due to sudden onset chest pain in the lower right quadrant. There was no evidence of acute coronary syndrome. Pacemaker interrogation revealed ventricular undersensing and loss of capture with high pacing output (7.5 volts at 1.5 ms), as well as intermittent diaphragmatic stimulation. The chest X-ray obtained in the postero-anterior view revealed the RV lead outside the heart silhouette (Figure 1C). Additionally, the lateral view showed the lead in an unusual posterior aspect (Figure 1D). A chest computed tomography with tridimensional reconstruction confirmed RV lead perforation through the RV apex, with 7 cm of lead positioned outside of the heart. There was no pericardial effusion (Figure 2A and 2B). Patient underwent lead removal and repositioning in a slightly different place in the RV apex. A pericardial drain was also inserted to monitor bleeding. The lead sensed R waves at 21.1 millivolts with an impedance of 707 ohms and pacing threshold of 0.5 volts at 0.6 milliamperes. The post-operative course was uneventful. The pericardial drain was removed after minimal drainage and patient was discharged home.

Discussion

Lead perforation is a relatively rare (0.3-1%) complication of pacemakers and ICD. It usually occurs within the first 24 hours after implantation, more commonly with active fixation leads and in the atrial aspect. Late perforation is believed to be very rare. The clinical course is extremely variable with some patients presenting completely asymptomatic, while others can develop cardiac tamponade and hemodynamic instability. The clinical predictors associated with lead perforation are use of temporary pacemaker, helical screw leads and steroids. Potential predictors for late perforation, particularly associated with passive fixation leads, include smaller lead diameters, septal or apical positioning, as well as high degree of slack on the ventricular lead. In the present case the chest X-Ray after the initial implant (Figure 1A) showed a lead positioned in the RV apex and apparently with ideal slack. However, the lateral view shows a distal angle in the RV lead, which could increase the tension in the lead, causing perforation (Figure 1B).

In most patients, the leads can safely be removed under fluoroscopic guidance and close monitoring. Although controversial, the insertion of a prophylactic pericardial drain is based on professional judgment. The emergent risk of pericardial effusion and tamponade in these situations, as well the presence of surgical backup can favors the decision of prophylactic drain insertion. An interesting aspect
observed is that progressive technology has resulted in the development of small diameter leads with modified designs and increased stiffness at the tip, which are related to the increased number of perforation in patients who receive pacemaker or ICD.

The present case describes a patient with recent passive-fixation lead perforation and no hemodynamic instability. It is important for the general cardiologist to pay attention for minor signs that can suggest lead perforation, as major signs such as pericardial effusion are not necessarily present. Chest or upper abdominal pain and a pacemaker interrogation that shows lower impedance, as well as undersensing or failure to capture the involved chamber are suspicious findings.

Finally, the decision to implant leads based only on their size is not justified, as most recent complications described in the literature occurred in the newer models of pacemaker or ICD leads.

Potential Conflict of Interest
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RV perforation by a passive-fixation lead

Case Report

Figure 2 - Chest computed tomography (volume-rendering technique) demonstrating reconstruction of the right ventricular lead perforation (A and B). The tip of the pacemaker lead is marked with asterisk (*).

References


