Right coronary artery pseudoaneurysm post everolimus eluting stent implantation causing tamponade

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DESCRIPTION

A 49-year-old man with triple vessel coronary artery disease underwent percutaneous coronary intervention. The right coronary artery (RCA) was treated with two platinum chromium everolimus-eluting stents (Promus Stent; Boston Scientific, Natick, Massachusetts, USA); 3.5×16 mm in proximal RCA and 3.5×38 mm in mid-to-distal RCA, respectively, with stenting of the left anterior descending (LAD) artery and first obtuse marginal artery (OM1) as well.

The patient was readmitted a month later with chest pain and dyspnoea on exertion. On examination, the patient had tachycardia, elevated jugular venous pressures and mild hypotension. A catheter angiogram revealed a small pseudoaneurysm arising from the RCA with occluded mid RCA stent. In view of disproportionately severe symptoms compared with the size of the pseudoaneurysm and to delineate the anatomy of the pseudoaneurysm for planning management, a coronary CT angiography (CTA) was advised.

CTA revealed patent proximal RCA stent, with near occlusion of the RCA just proximal to the mid RCA stent (figure 1A). A small pseudoaneurysm

was seen arising at the level of proximal 1/3rd of the second stent, with a large surrounding haematoma in the atrioventricular groove (figure 1A–D). The stent showed in-stent total occlusion, with reformation of distal RCA and a good-sized posterior descending artery. Mass effect was noted on the right atrium (RA) and right ventricle with dilated RA and inferior vena cava with flattening of the interventricular septum, indicating localised tamponade (figure 1B,C). The LAD and OM1 stents were patent. The patient underwent surgery with repair of the RCA pseudoaneurysm using direct pledgeted sutures and had an uneventful post-operative course.

Coronary artery pseudoaneurysms can develop from 1 week to 4 years after drug-eluting stent (DES) implantation.¹ The postulated underlying mechanisms include antiproliferative and antimetabolite action of the drug and inflammatory response to the drug, the polymer or the stent platform which results in incomplete endothelialisation and poor wound healing.¹ Thus, a combination of initial injury induced by stent placement and the biological response to DES might be the cause behind pseudoaneurysm formation. These pseudoaneurysms may rapidly enlarge leading

Figure 1 Curved multiplanar reformatted image of coronary CT angiography (A) depicts the patent proximal right coronary artery (RCA) stent with near occlusion of the RCA just proximal to the mid-RCA stent and total in-stent occlusion of the mid-RCA stent (block arrow). A small pseudoaneurysm (white arrowhead) is seen arising at the level of proximal one-third of the second stent, with a large surrounding haematoma (*). Reformation of distal RCA is also noted. Four chamber (B) and short axis view (C) reconstructed images show the pseudoaneurysm (white arrowhead) adjacent to the RCA stent (block arrow) with adjacent haematoma (*) seen indenting the right atrium and right ventricle with flattening of interventricular septum (black thin arrows) to the left side. Non-contrast axial CT image (D) shows the hyperdense haematoma (*) (mean attenuation ~64 HU) in the right atrioventricular groove surrounding the stent (block arrow). Abbreviations: LA, left atrium; LV, left ventricle; RA, right atrium; RV, right ventricle.

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Images in...

to rupture and can rarely also cause cardiac tamponade, as was observed in this case.²

Learning points

- ▶ Coronary artery pseudoaneurysms can develop from 1 week to 4 years after drug-eluting stent implantation.
- ▶ A combination of initial injury induced by stent placement and the biological response to drug-eluting stent, including antiproliferative and antimetabolite action of the drug and inflammatory response to the drug, the polymer, or the stent platform, might result in pseudoaneurysm formation.
- ▶ Coronary artery pseudoaneurysms may rapidly enlarge leading to rupture and can rarely cause cardiac tamponade.

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