Successful treatment of acutely occluded left main coronary artery in a patient with an aortic valve bioprosthesis

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Introduction
Flow-limiting left main coronary artery (LMCA) thrombus is a life-threatening condition, unless immediate and successful treatment is undertaken.

We present a case of a patient with large thrombus formation within the LMCA, which occurred one day after diagnostic coronary angiography.

Case report
The 63-year-old male patient was hospitalized due to non-ST elevation myocardial infarction. His previous history included aortic valve replacement with bioprosthesis implantation and saphenous vein graft (SVG) bypass to the right coronary artery (RCA) 2 years ago, diabetes mellitus type II, previous thrombophlebitis and arterial hypertension.

The current pharmacological treatment included bisoprolol, angiotensin-converting enzyme inhibitors (ACE-I), aspirin and statin. A loading dose (600 mg) of clopidogrel was administered before admission to the hospital.

Echocardiography examination showed a well-functioning aortic bioprosthesis, preserved left ventricle ejection fraction and mild left ventricle wall hypertrophy.

Troponin level on admission was slightly elevated to 0.11 ng/ml (ULN 0.014 ng/ml).

The angiography demonstrated trifurcation of the LMCA with borderline lesions in the left coronary artery (RCA) 2 years ago, with competitive flow to its distal segment from both the native RCA and patent SVG.

One day after the catheterization and just before planned discharge, the patient developed acute chest pain with signs of cardiogenic shock. ECG showed ST-segment elevation in I, aVL, aVR and V4–V6 leads with concomitant right bundle branch block and left anterior hemiblock (Figure 2 E).

Immediate (within 10 min), repeated angiography revealed a large thrombus within the LMCA, almost totally filling the lumen with TIMI 2 flow downstream in the main epicardial arteries (Figures 1 A, B). Abciximab and a loading dose of ticagrelor were administered. Despite several passages of manual thrombectomy, LMCA patency was not improved. Thus, percutaneous coronary intervention (PCI) with implantation of two everolimus-eluting stents (3.5 × 20 mm and 4.0 × 12 mm) into the LMCA and ostium of the intermediate branch (InB) was performed. A good angiographic result in the LMCA was achieved. The patient’s haemodynamic status improved and ECG normalized (Figure 2 F). Taking into account the given contrast amount and stabilized course, the borderline ostial lesion in the left anterior descending coronary artery (LAD) was left untreated. This lesion was visualized again in control angiography 3 days later and assessed as significant. Thus, same-time PCI with a radial approach was performed. After predilatation of the ostia of the LAD and left circumflex coronary artery (LCX), a 3.5 × 18 mm sirolimus-eluting stent at 14 atm was implanted into the LAD ostium. Finally sequential kissing balloon postdilatation was performed in the LAD/InB (3.5 mm and 3.5 mm at 12 atm), LCX postdilatation (3.0 mm at 12 atm), LAD/InB kissing re-postdilatation and LMCA postdilatation with 3.5 mm at 18 atm (Figures 2 H–I). A good angiographic result was achieved (Figure 2 J). A maximal troponin T increase to 1250 ng/ml was noted.

Transesophageal echocardiography performed subsequently did not show any thrombotic material in the Valsalva sinus.
Figure 1. A, B – Left coronary artery without significant lesions. C, D – Borderline lesion in proximal segment of the right coronary artery with distal concomitant filling from both native artery and patent saphenous vein graft. E, F – Sequential images of non-coaxial intubation of the left main coronary artery.
Figure 2. A – Large thrombus in the left main coronary artery almost totally occluding the lumen. B – Magnification of the image “A”. C – Stent positioning within the left main coronary artery. D – Final angiographic result after implantation of two stents into the left main coronary artery and intermediate branch. E – ECG during chest pain. F – ECG after coronary angioplasty. G – Ostial stenosis in the left anterior descending coronary artery. H – Stent positioning in the ostium. I – Kissing balloon. J – Final angiographic result.
Double antiplatelet treatment (aspirin and ticagrelor) was prescribed for 12 months.

Discussion

The true incidence of LMCA thrombus remains unknown, as many of the patients die before reaching the hospital. The origin of LMCA thrombus in this case remains unclear. However, the close time-course correlation with angiography indicates two most likely event explanations. Firstly, angiographically non-visible catheter-induced LMCA dissection might have led to thrombus formation or subintimal hematoma propagation. Secondly, bioprosthetic-associated thrombus mobilization and migration into the LMCA as a complication of the angiography cannot be ruled out.

Non-coaxial LMCA intubation was seen in our case (Figures 1 E, F), but no traces of any LMCA dissection were visible at the end of LMCA intubation. What is not seen in the angiography may be well visualized by optical coherence tomography (OCT). The OCT-based studies of the fate of stent implantation-induced edge dissections indicate that this is usually a benign phenomenon. Severe catheter-induced LMCA dissection is a rare and dramatic complication, and LMCA lumen compromise can happen immediately thereafter. Our patient did well nearly 24 h following diagnostic catheterization. Bioprosthetic valve-associated thromboembolism is a much rarer cause of acute coronary syndrome (ACS) compared to mechanical valves [1]. The ESC guidelines do not recommend anticoagulation therapy after aortic bioprosthetic valve implantation. This is supported by observations from the ANSWER registry, where the patients taking warfarin had a similar incidence of overall thromboembolism as those not taking warfarin [2].

Whatever the source, the consequences of coronary thromboembolism depend on the location and lumen compromise. Already the worst scenario had presented in our case, and only quick restoration of normal blood flow guaranteed the therapeutic success. In a study by Luca et al., among 24 patients with ACS due to thrombotic LMCA occlusion, the in-hospital mortality was as high as 58%, and 80% ended in cardiogenic shock [3]. The therapeutic options include fibrinolysis, balloon angioplasty, stenting and adjunctive pharmacotherapy such as IIb/IIIa inhibitors. Some authors even suggest a Fogarty maneuver [4], but distal embolization is a risk. Thus, the appropriate treatment choice remains a challenge.

The role of thrombectomy after ambivalent results of four main randomized trials (TAPAS, INFUSE-AMI, TASTE and TOTAL) is questioned, and the number of successfully managed LMCA thrombus reports with this tool is very limited [5].

Next, what made the stenting even more challenging in our case was the presence of LMCA trifurcation, although only the LMCA body was affected by the thrombus and not the trifurcation region. According to the modified Medina classification for trifurcation or quadrifurcration lesions [6], the thrombus in the LMCA trunk represented the “1,0,0,0” scheme. However, plaque shift after stent implantation from the LMCA into the InB during the first PCI changed the anatomical scenario to “0,1,0,0”. This lesion (LAD ostium) seemed angiographically significant during control angiography. Still, one might question its significance and ask for assessment by fractional flow reserve. While embarking on re-PCI, an additional stent for the LAD ostium was required with obligatory kissing balloon postdilatation. When LMCA trifurcating branches are comparable in diameter, simultaneous triple balloon kissing balloon postdilatation may be the option. Although not applied in this case, intravascular ultrasound guidance seems to be justified in such procedures, as in bifurcating LMCA PCI [7].

Conclusions

Finally, taking into account the complex PCI of LMCA trifurcation in the settings of ST elevation myocardial infarction and concomitant diabetes mellitus, a more potent P2Y12 inhibitor (ticagrelor) may be preferred over clopidogrel.

Conflict of interest

The authors declare no conflict of interest.

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