UNSUCCESSFUL PRIMARY PCI FOR POSTPARTAL DISSECTION OF ALL THREE MAJOR CORONARY ARTERIES

Svetlana Apostolović¹, Miodrag Damjanović¹, Dragan Mihailović², Aleksandra Antović³, Miloje Tomašević^{4,5}, Slobodan Obradović⁶, Nemanja Stepanović⁷

Coronary artery dissection is rare but very dangerous condition which can result in myocardial infarction. It mostly occurs in young postpartum women. The left anterior descending artery is affected in 75% of cases and in some extremely rare cases dissection may include all three coronary arteries. Treatment guides are still not consistent. While some authors recommend medical treatment other recommend surgical one. Case report. We present a 36-year-old female patient who was admitted to our hospital with intermittent chest pain and dyspnea that occurred two hours before admission. After premedication with Aspirin 300 mg per os, Clopidogrel 300 mg per os and Enoxaparin 30 mg iv, the pain disappeared. Chest pain was repeated after five days. Coronary angiography revealed spiral dissection in medial segment of LAD. After PCI attempt, dissection progressed and ended fatal. Pathological finding was intimal dissection of the coronary arteries with loss of some parts of internal elastic lamina. It is interesting that in the literature there is no consistent opinion about therapeutic approach to SDCA and we hope that this case report will contribute to elucidating the problem. *Acta Medica Medianae 2014;53(1):47-52.*

Key words: acute spontaneous coronary artery dissection, fibromuscular dysplasia, PCI treatment of SCAD

University of Niš, Faculty of Medicine, Clinic of Cardiology, Niš, Serbia¹ University of Niš, Faculty of Medicine, Institute of Pathology, Niš, Serbia² University of Niš, Faculty of Medicine, Institute of Forensic Medicine, Niš, Serbia³ Clinical Center Belgrade, Belgrade, Serbia⁴ University of Kragujevac, Faculty of Medicine, Kragujevac, Serbia⁵ Military Medical Academy, Belgrade, Serbia⁶ University of Niš, Faculty of Medicine, PHD study group, Niš, Serbia⁷

Contact: Nemanja Stepanović Neretvljanska 20 Serbia, 18000 Niš E-mail: nemstep@gmail.com

Introduction

Spontaneous coronary artery dissection (SCAD) is generally considered a rare event which, if not recognized and treated on time, can be fatal. SCAD is an unusual cause of acute coronary syndrome which is not associated with conventional risk factors. It was first described in 1931 in a 42-year-old healthy woman who died shortly after the first symptoms appeared (1). It is difficult to estimate real epidemiology trends due to the limited number of reported cases. However, approximately 500 cases of SCAD have been described in literature. The majority of patients (71.9%) were women and 28.1% were men. SCAD has been reported in 10% of women

younger than 50 years presenting with acute coronary syndrome. Same studies have shown that the prevalence of SCAD increases with ageing of the population, so its highest prevalence (76%) is among patients under the age of 40 (2). However, some latest reports showed that SCAD is not such a rare manifestation. In the population of women who underwent coronary angiography, 29.3% had SCAD (3). One third of them were young, healthy women in the last trimester of pregnancy or shortly after delivery, as Koul et al. reported. Most of them presented in the postpartum period (78%) and the rest of them during pregnancy. SCAD usually occurs within two weeks of delivery, but in some patients it could present very late (10-12 weeks postpartum). The left anterior descending artery (LAD) is affected in 75%, the right coronary artery (RCA) in 20%, the left circumflex artery (LCx) in about 4%, and the left main coronary artery (LMCA) in less than 1% of the cases (4). In extremely rare cases, SCAD may include all tree coronary arteries simultaneously (5).

Case report

We present a case of a 36-year-old woman who was admitted to our hospital with intermittent chest pain and dyspnea that occurred two hours before admission. Blood pressure was 120/ 80 mm Hg with heart rate 100/min. It was tenth postpartum day after an uncomplicated spontaneous vaginal delivery. She had two successful pregnancies (7 and 3 years before) and was on hormone fertility stimulation therapy in the last three years. Two months before delivery she was taking Hexoprenaline (0.5 mg per day) and Verapamil 80 mg (per day). She had been an active smoker for 10 years, (approximately 20 cigarettes per day) without other risk factors for coronary artery disease.

Baseline electrocardiogram (ECG) showed negative T waves in D2, D3 and aVF, V2-V6 leads which suggested anterior-lateral and inferior ischemia (Figure 1A). Troponin I level was 1.10 ng/ml. Echocardiography revealed akinetic apex of the left ventricle and estimated ejection fraction was 63%. After premedication with aspirin 300 mg per os, clopidogrel 300 mg per os and enoxaparin 30mg iv, the pain disappeared. Chest pain was repeated after five days with ECG changes - ST elevation in D2, D3, aVF, V2-V6 leads (Figure 1B). Troponin I level was 3.14ng/ml. The patient was immediately transferred to invasive coronary unit for further diagnosis and treatment.

Coronary angiography revealed spiral dissection of LAD and LCx (Figure 2A) with 80-90% diameter stenosis in the medial segment of LAD (Figure 2B). The lesion in the medial segment of LAD had the appearance of a sharp narrowing of the artery lumen with irregularities.



Figure 1. ECG showing: negative T waves in D2, D3 and aVF, V2-V6 leads (A); ST elevation in D2, D3, aVF, V2-V6 leads (B)

Right coronary artery catheterization revealed dissection at the ostium (Figure 1C). Dissection continued to spread proximally from both LAD and LCx towards the LMCA. Because the patient was hemodinamically unstable with signs of cardiogenic shock and bradycardia, we placed contrapulsatory intra-aortic balloon pump and temporary pace maker. Stents implantation was done from the medial segment of LAD to the LMCA (Figures 2D, 3A, 3B). After stent placement in the LMCA, patient was disoriented and lost consciousness, with Thrombolysis in Myocardial Infarction (TIMI) 2 coronary flow in both LAD and LCx. After the implantation of another stent in the medial segment of LAD due to the uncovered dissection (Figure 3C), no-reflow phenomenon and consequently cardiac arrest developed (Figure 3D). Cardiopulmonary resuscitation was unsuccessful.

An autopsy was done.

Pathological finding was intimal dissection of the coronary arteries with the loss of some parts of internal elastic lamina which is labeled as fibromuscular dysplasia (Figure 4).



Figure 2. Coronary angiography series showing: LAO CRAN, (LJ 4.0; 6 F) LCA where we can see sharp lumen narrowing in medial to distal segment of LAD, there are no any other changes in coronary arterial system (A); LAO CAUD, (RJ, 4.0; 6F) RCA, dissection of ostial part of RCA (B); RAO CAUD: First projection of LCA with guding catheter (GEBU 3.75) where spiral dissection is present in the main stem of LAD and ACx (C); BMW wire placed in LAD and ACx. First stent was implanted in the medial part of LAD (D)



Figure 3. Coronary angiography series show: Second stent placed in the proximal part of LAD and overlapped with the first one. TIMI flow in LAD was 3 (A); Stent placed in the main stem (B); Dissection progressed to aorta and flow in distal part of LAD was lost. We placed one more stent in the proximal part (C); Last projection before exitus letalis (D)



Figure 4. Pathological findings show: Coronary artery dissected with blood. HE, Ob.x4 (A); Thickened intima and disrupted internal elastic lamina. Media destroyed with blood. Autofluorescence, obj.x10 (B)

Discussion

All SCAD patients have been traditionally divided into tree groups: patients with coronary artery disease (CAD), women in the last trimester of pregnancy or shortly after delivery, and the idiopathic group. SCAD related to pregnancy is usually labeled as p-SCAD (6). The first peripartum case of SCAD was reported in 1952 (1). Etiology of SCAD remains unclear. De Maio showed that only 30% of patients had risk factors for atherosclerosis but they found that both multiparity and advanced age are related to SCAD diagnosis. However, Hering et al. showed that atherosclerosis is more often the cause of SCAD. They reported that rupture of atherosclerotic plaque occurred in 35 of 42 patients with SCAD (3). Intimal thickening caused by hormone level increase during pregnancy is also reported to be the possible cause of pregnancy associated SCAD as Gammal showed (7). He showed significant intimal thickening in female rats exposed to synthetic estrogen and progesterone. Literature review of histological finding showed that the main findings are: eosinophiles infiltrate, cystic medial necrosis, intimal tear, but, in some cases, there were no pathological findings. Role of eosinophiles in SCAD has been well described in literature. Basso et al. described hematoma between tunica media and adventitia. Study from 1996 showed that SCAD was the cause of acute myocardial infarction or sudden cardiac death. In all cases women were involved. Histology findings included eosinophilic infiltrate in four cases, cystic medial necrosis in one case and in the rest of them there was no histological finding (8). Significantly different eosinophile presence in SCAD group was shown when compared to control group. The explanation of possible relation between eosinophiles and dissection is well documented (9). Connective tissue disorders have been lately reported as possible causes of SCAD. In our case, segmental disruption of internal elastic lamina was present. Internal lamina loss can be part of arterial fibromuscular dysplasia (FMD), which is lately described as one of relatively important causes of SCAD. Relations between fibromuscular dysplasia and SCAD have been shown in literature (3,10). According to classification described by Harrison, our case can be diagnosed as intimal fibromuscular dysplasia (11).

The treatment of SCAD can include medical therapy and revascularization procedures such as percutaneous coronary intervention or coronary artery bypass graft surgery. The decision which treatment to use depends on the clinical presentation and the extent of coronary dissection. Conservative treatment with antiplatelet medications, antithrombin agents, nitrates, and beta blockers may be a reasonable approach in asymptomatic, stable patients with limited dissections (12,13). The literature today abounds with data which have described PCI as successful main treatment of SCAD (14). However, in all these cases dissection occurred in one vessel, mostly LMCA, unlike our case where dissection occurred in all tree arteries simultaneously (15). For dissection involving several coronary vessels, beating heart surgery is a recommended first line treatment since it can restore blood flow distal to the dissection. This intervention also provides reinforcement for false and true channel (16). Tweet et al. showed that patients with a single lesion in the left or right coronary artery had significantly better outcome when treated with an early aggressive strategy. Early treatment with CABG or percutaneous coronary intervention following the diagnosis of SCAD leads to less need for further intervention (17).

If we consider this case, we are sure that the operator (the author of this paper) was not up to the task. It is correct that we immediately set up a balloon pump and temporary pacemaker to provide cardiac output and hemodynamic support. Should the patient be immediately referred to an emergency surgery unit? (In our center there is no such possibility). Was it a basic mistake to place stents in the LAD from the distal segment to the main stem? Considering this case, knowing that most SCAD have fundamentally less valuable media layer and that it usually progresses proximally and distally through the vessel wall, we believe that it is correct to put a stent in the LMCA toward the LAD and than refer the patient to immediate surgery in other center. We hope that the comments of readers of this text will help interventionalists in better understanding of this potentially fatal event and point out the best therapeutic approach in these and similar cases.

Conclusion

This case report shows one of the types of fibromuscular dysplasia as a cause of spontaneous dissection of the coronary arteries and acute coronary syndromes (ACS) in the postpartum period. It also draws attention to this population of patients experiencing ACS and high probability of SDCA as a main cause of ACS. It is interesting that in the literature there is no consistent opinion about therapeutic approach to SDCA and we hope that this case report will contribute to elucidate the problem.

References

- Pretty HC. Dissecting aneurysm of coronary artery in a woman aged 42. Br Med J 1931; 1:667.
 Hering D, Piper C, Hohmann C, Schultheiss HP,
- Hering D, Piper C, Hohmann C, Schultheiss HP, Horstkotte D. Incidence, etiology and therapy of spontaneous coronary artery dissection. A prospective monocenter study of 3800 consecutive patients. Z Kardiol 1998; 87:961–70. [CrossRef] [PubMed]
- 3. Saw J, Poulter R, Fung A, Wood D, Hamburger J, Buller CE. Spontaneous coronary artery dissection in patients with fibromuscular dysplasia. Circ Cardiovasc Interv 2012; 5:134-7. [CrossRef] [PubMed]
- Koul AK, Hollander G, Moskovits N, Frankel R, Herrera L, Shani J. Coronary artery dissection during pregnancy and the postpartum period: Two case reports and review of literature. Catheter Cardiovasc Interv 2001; 52:88–94. [CrossRef] [PubMed]
- Maresta A, Varani E, Balducelli M, Vecchi G. Spontaneous coronary dissection of all three coronary arteries: A case description with mediumterm angiographic follow-up. Ital Heart J 2002; 3:747-51. [PubMed]
- 6. Lovitt WV, Corzine WJ. Dissecting intramural hemorrhage of anterior descendingbranch of left coronary artery. Arch Pathol 1952; 54: 458-62.
- Gammal EB. Intimal thickening in arteries of rats treated with synthetic sex hormones. Br J Exp Pathol 1976; 57(2):248-54. [PubMed]
- Basso C, Luigi MG, Thiene G. Spontaneous coronary artery dissection: a neglected cause of acute myocardial ischaemia and sudden death. Heart 1996; 75:451-4. [CrossRef] [PubMed]
- 9. Azam MN, Roberts DH, Logan WF. Spontaneous coronary artery dissection associated with oral

contraceptive use. Int J Cardiol 1995; 48:195-8. [CrossRef] [PubMed]

- Stanley JC, Gewertz BL, Bove EL, Sottiurai V, Fry WJ. Arterial fibroplasia, histopathologic character and current etiologic concepts. Arch Surg 1975; 110:561–6. [CrossRef] [PubMed]
- 11. Harrison EG, McCormack LJ. Pathologic classification of renal arterial disease in renovascular hyper tension. Mayo Clin Proc 1971; 46:161–7. [PubMed]
- 12. Zampieri P, Aggio S, Roncon L, Rinuncini M, Canova C, Zanazzi G, et al. Follow up after spontaneous coronary artery dissection: A report of five cases. Heart 1996; 75:206–9. [CrossRef] [PubMed]
 13. Almeda FQ, Barkatullah S, Kavinsky CJ. Sponta
- Almeda FQ, Barkatullah S, Kavinsky CJ. Sponta neous coronary artery dissection. Clin Cardiol 2004; 27:377–80. [CrossRef] [PubMed]
- 14. Nogueira de Macedo R, de Paula Miranda S, Vieira da Costa RL. Spontaneous coronary artery dissection — a diagnosis to be considered in young patients presenting with acute myocardial infarction. J Invasive Cardiol 2009; 21(12):E245-7. [PubMed]
- 15. Tu CM, Taso TP, Chu KM, Cheng SM, Lin WS. Long segmental right coronary artery dissection success fully treated by percutaneous coronary intervention. J Med Sci 2010; 30(5):221-4. [CrossRef]
- 16. Shamloo BK, Chintala RS, Nasur A, Ghazvini M, Shariat P, Diggs JA, et al. Spontaneous coronary artery dissection: aggressive vs. conservative therapy. J Invasive Cardiol 2010; 22(5):222-8. [PubMed]
- 17. Tweet M, Hayes S, Pitta S, et al. Clinical features, management, and prognosis of spontaneous coronary artery dissection. Circulation 2012; 126 (5):579-88. [CrossRef] [PubMed]

SPONTANA DISEKCIJA SVE TRI KORONARNE ARTERIJE KAO UZROK AKUTNOG INFARKTA MIOKARDA

Svetlana Apostolović, Miodrag Damjanović, Vanja Miloradović, Valentina Čupić, Miloje Tomašević, Goran Davidović, Nemanja Stepanović

Spontana disekcija koronarnih arterija je redak, ali vrlo opasan događaj, koji može da se završi infarktom miokarda. Najčešće se javlja kod mladih žena pred kraj ili neposredno nakon porođaja. Leva koronarna arterija zahvaćena je u 75% slučajeva, a u nekim, vrlo retkim slučajevima, disekcija može da zahvati sve tri koronarne arterije. Terapijske opcije još uvek nisu jedinstvene. Prikazaćemo ženu staru 36 godina, koja je primljena u bolnicu sa naizmeničnim bolom u grudima i dispnejom koji su se javili dva sata pre prijema. Nakon medikacije Aspirinom 300mg per os, Klopidogrelom 300mg per os i Enoxaparinom 30mg iv, bol je prestao, da bi se nakon pet dana ponovo javio. Koronarna angiografija je otkrila spiralnu disekciju u srednjem segmentu LAD. Pokušan je PCI, međutim, bezuspešno. Patološki nalaz pokazao je intimalnu disekciju koronarnih arterija sa gubitkom unutrašnje elastične membrane. Interesantno je da u literaturi ne postoji uniformno mišljenje o terapijskom pristupu disekciji. Nadamo se da će ovaj slučaj pomoći u rešavanju problema. *Acta Medica Medianae 2014;53(1):47-52.*

Ključne reči:: spontana disekcija koronarnih arterija, fibromuskularna displazija, PCI tretman SDKA