# **Chapter 8**

## OVERVIEW OF ENDOVASCULAR TREATMENT FOR LOWER EXTREMITY PERIPHERAL ARTERY DISEASE

## Mustafa ALDAĞ<sup>1</sup>

#### Introduction

Atherosclerosis is a systemic disease of the vessels causing luminal focal or diffuse narrowing as a result of the accumulation of lipid and fibrous material between the intimal and medial layers of the artery. Atherosclerosis of the noncardiac vessels is defined as peripheral artery disease (PAD). PAD remains a significant health problem across the globe, also it is the third most common manifestation of cardiovascular disease, following occlusive coronary artery disease and stroke (1,2). PAD is extremely common, particularly in the elderly patients (3), with the elderly population expected to increase 22% by the year 2040. The overall prevalence of lower extremity PAD varies widely depending upon the population studied, but is estimated to be approximately 13 percent of adults older than 55 years (4). Although gold standard therapy is conventional surgery when revascularization is needed, the endovascular treatment has become the mainstay of treatment in recent years.

#### **Clinical Presentations**

PAD represents a spectrum from asymptomatic stenosis to limb-threatening ischemia. Claudication, which is defined as reproducible ischemic muscle pain, is one of the most common manifestations of PAD caused by atherosclerosis. Claudication occurs during physical activity and is relieved after a short rest. Pain develops because of inadequate blood flow. Although symptoms are most frequently localized to the calf, the thigh or buttocks may also be affected. The location of the pain in patients with PAD is determined by the anatomic location of the arterial lesions. PAD is most common in the distal superficial femoral artery, a location that corresponds to claudication in the calf muscle area. When atherosclerosis is distributed throughout the aortoiliac area, thigh and buttock muscle claudication predominates. When claudication is used as an indicator, it is estimated that 2% of the population aged 40-60 years and 6% of the population older than 70 years are affected. Predicted allcause mortality for PAD patients with claudication is approximately 30% at 5 years of follow-up, 50% at 10 years, and 70% at 15 years (5).

<sup>&</sup>lt;sup>1</sup>MD, Bahçeşehir University School of Medicine, Department of Cardivascular Surgery, VM Medicalpark Pendik Hospital, Istanbul, Turkey

### **Endovascular Treatment in Femoro-Popliteal Occlusions**

The most common location (> 70%) for PAD is the femoropopliteal segment (16). In contrast to the aorto-iliac lesions the femoropopliteal arteries represent a unique challenge for endovascular revascularization. Usually, these lesions found to have diffuse, calcific often occlusive atherosclerosis which is likely related to both its length and location (17). In addition, endovascular treatment of the femoropopliteal region is more difficult because this region is mobile and exposed to many external forces. There is no surgical or endovascular therapy indication in asymptomatic patients, however numerous treatment modalities for symptomatic femoropopliteal lesions are currently available (PTA, drug-coated PTA, stenting, atherectomy), with various degrees of technical success and limited long-term data. Percutaneous trans-luminal angioplasty though effective for shorter (<5 cm) non-calcified lesions, fails when lesions are longer, calcified and more complex. Moreover, drug-coated balloon angioplasty with or without atherectomy has become first choice in femoropopliteal lesions nowadays. A multicentric data about primary stenting for TASC C & D femoropopliteal lesions was published by Brouillet in 2018. According to their study, early in-stent thrombosis was observed as 3.4 percent. The 6 and 12 months primary patency rates were 85.2±2. and 67±3.%, respectively. The 12-month instent thrombosis and restenosis rates were 19.6% and 13.9% and also the stent fracture rate was presented as 10.2% (18). However, Katsogridakis and co-workers published a systematic meta-analysis regarding drug-eluting stents (DES) and they concluded that the short-term outcomes of DES for TASC C & D femoro-popliteal lesions are encouraging (19).

### References

- 1. Jelani QU, Petrov M, Martinez SC, Holmvang L, Al-Shaibi K, Alasnag M. Peripheral arterial disease in women: an overview of risk factor profile, clinical features, and outcomes. *Curr Atheroscler Rep.* 2018 Jun 2;20(8):40. Doi: 10.1007/s11883-018-0742-x.
- 2. Klein AJ, Ross CB. Endovascular treatment of lower extremity peripheral arterial disease. *Trends Cardiovasc Med.* 2016 Aug;26(6):495-512. Doi: 10.1016/j.tcm.2016.02.007.
- 3. Selvin E, Erlinger TP. Prevalence of and risk factors for peripheral arterial disease in the United States: results from the National Health and Nutrition Examination Survey, 1999-2000. *Circulation*. 2004;110(6):738–743.
- 4. Fowkes FG, Rudan D, Rudan I et al. Comparison of global estimates of prevalence and risk factors for peripheral artery disease in 2000 and 2010: a systematic review and analysis. *Lancet* 2013; 382:1329-40.
- Norgren L, Hiatt WR, Dormandy JA, Nehler MR, Harris KA, Fowkes FG; TASC II Working Group. Inter-Society Consensus for the Management of Peripheral Arterial Disease (TASC II). J Vasc Surg. 2007 Jan;45 Suppl S: S5-67.
- Crawford F, Welch K, Andras A, Chappell FM. Ankle brachial index for the diagnosis of lower limb peripheral arterial disease. *Cochrane Database Syst Rev.* 2016 Sep 14;9:CD010680. Doi: 10.1002/14651858.CD010680.pub2. Review.
- Rutherford RB, Baker JD, Ernst C, Johnston KW, Porter JM, Ahn S, Jones DN. Recommended standards for reports dealing with lower extremity ischemia: revised version. *J Vasc Surg.* 1997 Sep;26(3):517-38. Erratum in: J Vasc Surg 2001 Apr;33(4):805.
- 8. Fontaine R, Kim M, Kieny R. Surgical treatment of peripheral circulation disorders. *Helv Chir Acta*. 1954 Dec;21(5-6):499-533. German.

#### Cardiovascular Surgery

- Ricco JB, Parvin S, Veller M, Brunkwall J, Wolfe J, et al. Statement from the European Society of Vascular Surgery and the World Federation of Vascular Surgery Societies: Transatlantic Inter-Society Consensus Document (TASC) III and International Standards for Vascular Care (ISVaC). *Eur J Vasc Endovasc Surg.* 2014 Feb;47(2):118. Doi: 10.1016/j.ejvs.2013.11.009.
- 10. Murphy TP, Cutlip DE, Regensteiner JG, Mohler ER, Cohen DJ, et al. Supervised exercise, stent revascularization, or medical therapy for claudication due to aortoiliac peripheral artery disease: the CLEV-ER study. J Am Coll Cardiol. 2015 Mar17;65(10):999-1009. Doi: 10.1016/j.jacc.2014.12.043.
- 11. Fakhry F, Fokkenrood HJ, Spronk S, Teijink JA, Rouwet EV, Hunink MGM. Endovascular revascularisation versus conservative management for intermittent claudication. *Cochrane Database Syst Rev.* 2018 Mar 8;3:CD010512. Doi: 10.1002/14651858.CD010512.pub2.
- 12. Polat A. (2016). *Endovasküler cerrahiye giriş, temel tel ve kateter teknikleri*. Istanbul. Türk Kalp ve Damar Cerrahisi Derneği Bayçınar Tıbbi Yayıncılık
- 13. Indes JE, Mandawat A, Tuggle CT, Muhs B, Sosa JA. Endovascular procedures for aorto-iliac occlusive disease are associated with superior short-term clinical and economic outcomes compared with open sur- gery in the inpatient population. *J Vasc Surg.* 2010; 52:1173–1179,1179. e1.
- 14. Aggarwal V, Waldo SW, Armstrong EJ. Endovascular revascularization for aortoiliac atherosclerotic disease. *Vasc Health Risk Manag.* 2016 Mar 29; 12:117-27. Doi: 10.2147/VHRM.S98721.
- 15. De Donato G, Bosiers M, Setacci F, Deloose K, Galzerano G, et al. 24-Month Data from the BRAVISSIMO: A Large-Scale Prospective Registry on Iliac Stenting for TASC A & B and TASC C & D Lesions. *Ann Vasc Surg.* 2015;29(4):738-50. Doi: 10.1016/j.avsg.2014.12.027.
- 16. Zeller T. Current state of endovascular treatment of femoro-popliteal artery disease. *Vasc Med.* 2007; 12:223-34.
- 17. Klein AJ, Chen SJ, Messenger JC et al. Quantitative assessment of the conformational change in the femoropopliteal artery with leg movement. *Catheter Cardiovasc Interv.* 2009; 74:787-98.
- Brouillet J, Deloose K, Goueffic Y, Poirier M, Midy D, et al. Primary stenting for TASC C and D femoropopliteal lesions: one-year results from a multicentric trial on 203 patients. *J Cardiovasc Surg.* 2018 Jun;59(3):392-404. Doi: 10.23736/S0021-9509.16.09282-X.
- 19. Katsogridakis E, Ballance L, Cawley O, Antoniou GA. Drug-eluting stents for the treatment of complex femoro-popliteal disease: a systematic review and meta-analysis. *J Cardiovasc Surg (Torino).* 2018 Aug 29. Doi: 10.23736/S0021-9509.18.10614-8.