

PERCUTANEOUS CORONARY INTERVENTION OF OSTEAL LESIONS IN NEWLY ESTABLISHED TERTIARY CARE CARDIAC HOSPITAL

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ABSTRACT

Objective: To assess the outcome of percutaneous Angioplasty in patients with osteal lesions in Coronary artery Diseases.

Methodology: This is a retrospective analysis of all coronary angiograms performed at the catheterization laboratory of Karachi Institute of Heart Diseases (KIHD), a tertiary referral center in Karachi, Pakistan, between the periods August 2006 to August 2008.

Results: Fifty patients were enrolled which included thirty-five men and fifteen women and all were >40 years of age. Each patient had a single target osteal lesion: twenty nine patients underwent PCI for ostial LAD lesion (among them twenty males & nine females), six had ostial LCX (five males and one female), and fifteen patients had osteal RCA stenosis (10-male patients and 5- female patients). After high-pressure balloon dilatation residual stenosis was reduced. Twelve patients were treated with bare metal stents (BMS) while thirty eight with drug eluting stents (DES). In all cases the procedure was successful without any pre and post procedural complications.

Conclusion: A key issue in the treatment of an osteal lesion is to assure that the stent is inserted proximal enough to fully cover the osteal junction. Improvements in technique, equipment, adjunctive drug therapy and better understanding of the procedure have remarkably changed the practice of interventional cardiology

KEY WORDS: Osteal stenosis, percutaneous transluminal coronary Angioplasty, coronary artery disease.

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INTRODUCTION

The term "ostium" is derived from the Latin os, referring to the mouth. A lesion located at the ostium of a vessel is referred to as an "ostial lesion."¹ Generally, lesions located within 3 mm of the origin of a vessel are considered osteal, but some authors consider lesions up to 5 mm from the origin as osteal.²

Besides atherosclerosis, coronary osteal lesions may occur uncommonly in various other disease states such as syphilis, Takayasu's

arteritis, fibromuscular dysplasia and postradiation fibrosis.^{3,4} Female sex and family history of coronary artery disease and hypercholesterolemia are independent predictors of ostial right coronary artery lesion.⁵ Ostial left main coronary artery and right coronary artery lesions are strongly correlated.^{6,7} Bilateral tight coronary ostial lesion is potentially lethal and can present as sudden cardiac death, myocardial infarction, angina or ischemic cardiomyopathy. There is also imminent risk of sudden death due to global ischemia or infarction.⁸

Coronary angioplasty of ostial lesions has a high initial angiographic and clinical success rate but restenosis remains a limitation. The beneficial action of stents has been attributed to maximizing the initial luminal gain and overcoming the elastic recoil. PTCA using a cutting balloon followed by stenting is technically feasible. Stenosis of the right coronary ostium is a high-risk lesion for stent restenosis.⁹ With improvement in hardware and increasing experience, isolated ostial disease of the LMCA (>3.5mm) can be managed by stenting alone or debulking and stenting.^{10,11} A technique for correct positioning of a stent in an ostial stenosis by using a second wire passed through the last cell of a stent. The anchor wire technique, first described by Szabo will facilitate precise ostial stent placement and eliminate errors of positioning inside or outside the ostial narrowing.¹² DES implantation in de novo ostial LAD lesions appears safe and effective and is associated with a significant decrease in restenotic rates compared with historical experience with BMS.¹³ Stent insertion into saphenous vein graft ostial lesions is associated with a high procedural success rate,¹⁴ but Angioplasty of these lesions is further limited by the high restenosis rate. This may be as a result of the simultaneous presence of atherosclerosis, fibrosis, and calcification in the surrounding aortic wall, and the higher concentration of elastic and muscle fibers around the ostium, which causes recoil after balloon inflations.¹⁵ Conventional balloon dilatation can successfully dilate the majority of coronary artery stenoses. In this study we have analyzed the data of PCI of ostial lesions.

METHODOLOGY

This is a retrospective analysis of all coronary angioplasties performed at the catheterization laboratory of Karachi Institute of Heart Diseases (KIHD), a tertiary referral center in Karachi, Pakistan, between the periods August 2006 to August 2008.

An inclusion criterion was Single de novo lesions and ostial lesions. While patients with LVEF <30%, Left main >50%, coexisting chronic heart failure, valvular heart diseases, chronic renal failure, contraindication to anticoagulation, difficult anatomy, poorly visible lesions, overlap, multiple stenoses within one artery, diffuse disease, left main disease, multivessel disease were excluded from study.

All ostial lesions were crossed by guide wire through respective guiding catheter then predilated with appropriate balloon at approximately 6 to 8 ATM pressure followed by deployment of suitable stent according to lesion size and length. Both types of stents, Bare Metal (BMS) and Drug eluting (DES) were selected but majority were DES (Endeavour). All DES were sirolimus type, because restenosis rate is less with DES stent as compared to BMS but due to financial constraints we also deployed bare metal stents. We used Endeavour because government provided us as well as it is one of good quality DES. Patients were evaluated for the occurrence of major adverse cardiac events comprising death, acute myocardial infarction, and need for repeat revascularization with either coronary artery bypass surgery or PCI. Patients were discharged after 48 hours of procedure.

RESULTS

Table-I gives details of total study population. Fifty patients were recruited: Thirty-five men and fifteen women and all were >40 years of age. Each patient had a single target ostial lesion: twenty nine patients underwent PCI for ostial LAD lesion (among them twenty males & nine females), six had ostial LCX (five males and one female), and fifteen patients had ostial RCA stenosis (10-male patients and 5- female

Table-I: Demographics of patients undergoing PCI for osteal stenosis (n=50)

		Male	Female
Distribution of patients	50	35	15
LAD	29	20	9
LCX	6	5	1
RCA	15	10	5

Table-II: Use of BMS Vs DES stents for osteal stenosis (n=50)

	BMS	DES
Total patients	12	38
LAD	6	23
LCX	2	4
RCA	4	11

patients). After high pressure balloon dilatation residual stenosis was reduced in all the cases. This facilitated stent deployment while post dilatation was done in few cases if required. Table-II shows twelve patients were treated with bare metal stents (BMS) while thirty eight

with drug eluting stents (DES). In all cases the procedure was successful without any pre and post procedural complications. During 6 to 12 months follow up period we observed marked improvement of symptoms in majority of patients. Nine had myocardial infarction, 14 presented with arrhythmias, 11 with unstable

Table-III: Follow up of patients (n=50)

time	□		Gender		Total
			M	F	
6 Month	Myocardial infraction	Count	1	2	3
		% within gender	2.9%	13.3%	6.0%
	Arrhythmias	Count	2	3	5
		% within gender	5.7%	20.0%	10.0%
	Unstable angina	Count	2	3	5
		% within gender	5.7%	20.0%	10.0%
	Re-stenosis	Count	1	0	1
		% within gender	2.9%	0.0%	2.0%
	No disease identified	Count	29	7	36
		% within gender	82.9%	46.7%	72.0%
12 Month	Myocardial infraction	Count	2	4	6
		% within gender	5.7%	26.7%	12.0%
	Arrhythmias	Count	4	5	9
		% within gender	11.4%	33.3%	18.0%
	Unstable angina	Count	3	4	7
		% within gender	8.6%	26.7%	14.0%
	Re-stenosis	Count	3	2	5
		% within gender	8.6%	13.3%	10.0%
	No disease identified	Count	23	0	23
		% within gender	65.7%	0.0%	46.0%
		Count	35	15	50
		% within gender	100.0%	100.0%	100.0%

Marginal statistical association exist among male and females at 6 months (p-value: 0.07) while at 12 months significant difference exist among males and females (p-value<0.001) using Chi square test of significance

angina, six had re-stenosis .while no death has been reported so far.

DISCUSSION

Coronary osteal stenosis involving both the coronary ostia is a rare occurrence. The plaque formation can occur at unusual sites, including the ascending aorta and around the coronary ostia. This atheroma can interfere with aortic valve function.¹⁶ Bilateral coronary osteal stenosis in young subjects without known conventional coronary risk factors may be due to fibromuscular dysplasia, syphilitic aortitis, post radiation and Takayasu's arteritis or without any recognizable etiological background.¹⁷ Debulking before stenting located at the ostium of the LAD is safe and is associated with a high rate of technical success.¹⁸ Mechanism of balloon angioplasty action involves three events: plaque fracture, compression, and stretch.¹⁹ Balloon dilatation within a stenosis results in application of force in a random manner to the components of the stenosis.²⁰ Cutting balloon has been used for osteal lesions with or without stenting²¹ but we can't use cutting balloon because of financial constraints.

In our cases dilatation with the conventional balloon enabled sufficient luminal gain to be obtained, facilitating stent insertion. This combination resulted in a good post procedural result with no immediate or late complications. The guiding catheter is optimally positioned outside the ostium but in sufficient proximity to opacify the adjacent aorta and thereby localizes the target ostium. However, the guiding catheter must be maintained at a sufficient distance from the ostium to avoid dislodging or damaging the stent. The choice of therapy in osteal stenosis of the coronary artery has conventionally been CABG or surgical patch aorto-coronary osteo-plasty. Eversince various balloons, stents, and other devices are introduced the success rate of PCI is over 95%, and the risks of serious complications has also decreased considerably.²² As table 2 shows Drug -eluting stent implantation in osteal lesions achieved excellent results regarding re-stenosis and clinical outcomes compared with BMS

implantation.²³ Osteal lesions, either in native vessels or grafts, can be particularly difficult to dilate with conventional balloon angioplasty. Stent insertion into sapheneous vein graft aorto-osteal lesions is associated with a high procedural success rate,²⁴ but restenosis remains a limitation for plaque excision as the preferred treatment because both balloon angioplasty and stenting presented a very high risk of jeopardizing the ostium of the left anterior descending artery, the mid left circumflex, or both.^{25, 26} The beneficial action of stents has been attributed to maximizing the initial luminal gain and overcoming the elastic recoil. These mechanical advantages may be ideally suited for aorto-osteal lesions.²⁷

CONCLUSIONS

A key issue in the treatment of an osteal lesion is to assure that the stent is inserted proximal enough to fully cover the osteal junction. Although Cutting balloon angioplasty followed by stent insertion is a feasible technique for the treatment of osteal lesions, the cutting balloon proved useful in dilating stenoses resistant to conventional balloon dilatation alone and so facilitating insertion of a Stent and perhaps improving long term outcome. But due to limited resources we did conventional balloon dilatation. Improvements in technique, equipment, adjunctive drug therapy and better understanding of the procedure have remarkably changed the practice of interventional cardiology.

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REFERENCES

1. Pritchard CL, Mudd JG, Barner HB. Coronary osteal stenosis. *Circulation* 1975; 52: 46-48
2. Arima M, Kanoh T, Okazaki S, Iwama Y, Matsuda S, Nakazato Y. Long-term clinical and angiographic follow-up in patients with isolated ostial stenosis of the left coronary artery. *JPN Circ J.* 2009;73(7):1271-7. Epub 2009 May 8.
3. Goldstein J, Hobbs H, Brown M. Familial hypercholesterolemia. In: *The Metabolic and Molecular Basis of Inherited Disease.* New York: 2001, p 2863

4. Ajani AE, Kim HS, Castagna M, Satler LF, Kent KM, Pichard AD, et al. Clinical utility of the cutting balloon. *J Invasive Cardiol.* 2001 ;13(7):554-7.
5. Haridas KK, Kumar V, Rajesh T, Kumar MV, Pannekal B. Percutaneous transluminal angioplasty with cutting balloon and stenting for isolated bilateral aorto-coronary ostial stenosis in a young female. *Indian Heart J.* 2001;53(4):490-2.
6. Swissa M, Ayzenberg O, Caspi A. Symptomatic calcified ostial lesions in both left main and right coronary arteries. *Isr Med Assoc J* 2007; 9(11):829.
7. Darabian S, Amirzadegan A, Sadeghian H, Sadeghian S, Abbasi A, Raeesi M. *Angiology. Osteal Lesions of Left Main and Right Coronary Arteries: Demographic and Angiographic Features.* *Angiology* 2008 Apr 2.
8. Machado MN, Trindade PF, Miranda RC, Maia LN. Bilateral ostial coronary lesion in cardiovascular syphilis: case report. *Rev Bras Cir Cardiovasc.* 2008;23(1):129-31.
9. Chung CM, Nakamura S, Tanaka K, Tanigawa J, Kitano K, Akiyama T, et al. Stenting alone versus debulking and debulking plus stent in branch ostial lesions of native coronary arteries. *Heart Vessels.* 2004;19(5):213-20.
10. Tsunoda Taro, Nakamura Masato, Wada Masamichi, Ito Naoki, Kitagawa et al; Chronic stent recoil plays an important role in restenosis of the right coronary ostium. *Coronary Artery Disease* 2004; 15(1):39-44.
11. Kurbaan AS, Kelly PA, Sigwart U. Cutting balloon angioplasty and stenting for aorto-ostial lesions. *Heart* 1997; 77:350-352.
12. Maja S, Josko B. PCI of an osteal LCX stenosis, with protection of LAD, using two guiding catheters. *Acute Card Care.* 2006; 8(3):161-5
13. Seung KB, Kim YH, Park DW, Lee BK, Lee CW, Hong MK, et al. Effectiveness of Sirolimus-Eluting Stent Implantation for the Treatment of Ostial Left Anterior Descending Artery Stenosis With Intravascular Ultrasound Guidance. *J Am Coll Cardiol* 2005;46(5):787-792.
14. Chen J, Li JJ, Chen JL, Qiao SB, Xu B, Yang YJ. Drug-eluting stents for the treatment of osteal coronary lesions: comparison of sirolimus-eluting stent with paclitaxel-eluting stent. *Coron Artery Dis* 2008; 19(7):507-11.
15. Nassar H, Gotsman I, Gerganski P, Moseri M, Lotan C, Gotsman M. Cutting balloon angioplasty and stent implantation for aorto-ostial lesions: clinical outcome and 1-year follow-up. *Clin Cardiol* 2009;32(4):183-6.
16. Soulis J, Giannoglou G, Dimitrakopoulou M, Papaioannou V, Logothetides S, Mikhailidis D. Influence of Oscillating Flow on LDL Transport and Wall Shear Stress in the Normal Aortic Arch. *Open Cardiovasc Med J* 2009; 17;3:128-42.
17. Kawakami H, Matsuoka H, Koyama Y, Saeki H, Inoue K, Nishimura K, et al. Isolated left coronary osteal stenosis as a result of fibromuscular dysplasia in a young man. *JPN Circ J* 2000; 64: 988-999.
18. Croti UA, Gregori F Jr, Marcial MB, Dallan LA, Gregori TE, Oliveira DS. Coronary bilateral ostial enlargement using the saphenous vein in a patient with syphilitic aortitis. *Arq Bras Cardiol.* 2000;74(2):153-8.
19. Ito H, Piel S, Das P, Chhokar V, Khadim G, Nierzwicki R, et al. Long-term outcomes of plaque debulking with rotational atherectomy in side-branch ostial lesions to treat bifurcation coronary disease. *J Invasive Cardiol.* 2009; 21(11):598-601.
20. Stillabower ME. Longitudinal force focused coronary angioplasty: a technique for resistant lesions. *Cath Cardiovasc Diagn* 1994; 32:196-8.352
21. Chung CM, Nakamura S, Tanaka K, Tanigawa J, Kitano K, Akiyama T, et al. Comparison of cutting balloon vs stenting alone in small branch ostial lesions of native coronary arteries. *JPN Circ J* 2003;67(1):21-5.
22. Botsios S, Maatz W, Sprengel U, Heuer H, Walterbusch G. Patch angioplasty for isolated ostial stenosis of the left main coronary artery. *J Card Surg* 2008;23(6):743-6.
23. Liu P, Zhang S, Schiele F, Meneveau N, Bassand JP. Clinical outcome of patients with left anterior descending artery osteal lesions treated with percutaneous coronary intervention: case-matched comparison with bypass surgery. *Chin Med J (Engl).* 2003;116(6):844-8.
24. Barlis P, Kaplan S, Dimopoulos K, Ferrante G, Di Mario C. Comparison of bare-metal and sirolimus- or paclitaxel-eluting stents for aorto-ostial coronary disease. *Cardiology.* 2008;111(4):270-6. Epub 2008 May 2.
25. Croti UA, Gregori F Jr, Marcial MB, Dallan LA, Gregori TE, Oliveira DS. Coronary bilateral ostial enlargement using the saphenous vein in a patient with syphilitic aortitis. *Arq Bras Cardiol* 2000;74(2):153-8.
26. Gao LJ, Chen JL, Chen J, Yang YJ, Gao RL, Li JJ, et al. Long-term clinical efficacy of cutting balloon angioplasty followed by bare metal stent implantation for treating ostial left anterior descending artery lesions. *Clin Cardiol.* 2009;32(8):E31-5.
27. Tsagalou E, Stankovic G, Iakovou I, Melzi G, Cosgrave J, Ge L, et al. Early outcome of treatment of ostial de novo left anterior descending coronary artery lesions with drug-eluting stents. *Am J Cardiol.* 2006;15;97(2):187-91. Epub 2005 Nov 21.
28. Park DW, Hong MK, Suh IW, Hwang ES, Lee SW, Jeong YH, Kim YH, et al. Results and predictors of angiographic restenosis and long-term adverse cardiac events after drug-eluting stent implantation for aorto-ostial coronary artery disease. *Am J Cardiol* 2007; 15;99(6):760-5. Epub 2007 Jan 22.

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