ORIGINAL ARTICLE PERI-PROCEDURAL OUTCOME OF SAPHENOUS VEIN GRAFT INTERVENTION

Tariq Shakoor, Nadeem Hayat Mallick, Ahmad Noeman Punjab Institute of Cardiology, Lahore, Pakistan

Background: Patients who develop recurrent myocardial ischemia after coronary artery bypass graft (CABG) surgery are often referred for percutaneous coronary intervention. The objective of this study was to evaluate the clinical characteristics and peri-procedural outcomes in patients with prior CABG referred for percutaneous coronary intervention (PCI) over a 3 year period. Methods: Data were collected on patients who underwent coronary interventional procedures following CABG surgery. We evaluated angiographic procedural success and immediate outcome among patients who had undergone such procedures from Nov 2006 to Oct 2009 (n=113). Results: Patients in the 2006-2009 cohort had mean age 58.2 years, more patients were male (109 vs 4) and were more likely to have hypertension (57.5%), hyperlipidaemia (72.6%) and family history of ischemic heart disease (IHD) (65.5%), but less likely to have smoking (42.5%). Acute closure of stent leading to procedural failure was seen in 1 (0.9%) patient, sub-acute thrombosis of stent was seen in 1 (0.9%) patient, dissection or perforation of target vessel was seen in 3 (2.7%) and 1 (0.9%) patients respectively. Slow flow phenomenon was seen in 13 (11.5%) and post-procedural cardiac enzymes were raised in 6 (5.3%) patients. Conclusion: Success rates of saphenous vein graft (SVG) intervention and survival rate have improved with time as a result of improvements in technique and greater use of stents, filter devices and adjunctive medications.

Keywords: Percutaneous coronary intervention, saphenous vein graft, Ischemic heart disease, IHD

INTRODUCTION

As surgical coronary revascularization enters its fifth decade, a growing patient population faces the need for repeat revascularization. It is estimated that up to 10% of saphenous vein grafts (SVG) become occluded by the end of 1 year, followed by an additional attrition rate of 3–5% per year. Thus, at 10 years after operation, only 40–60% of bypass grafts are functional.^{1,2} Annual incidence of acute ischemic events in survivors of first bypass operation is 4–8%. These patients mostly have multi-vessel disease and they are usually receiving aggressive medical treatment at the time when they develop symptoms. Thus the crucial therapeutic decision is whether patient needs repeat coronary artery bypass graft surgery (CABG) or catheter based percutaneous coronary intervention (PCI).

Percutaneous coronary intervention (PCI) has been used to treat medically refractory myocardial ischemia in patients with prior CABG since the early 1980s, but even with the addition of stents, PCI of patients with prior CABG has been associated with worse outcomes than PCI of patients without prior CABG.^{3,4}

MATERIAL AND METHODS

This prospective study was conducted in Punjab Institute of Cardiology over a period of three years (from November, 2006 to October, 2009). Patients were included in the study if they fulfilled following clinical, laboratory and angiographic criteria: i) History of having CABG previously; ii) Development of ischemic symptoms after CABG; iii) Graft study showing critical stenosis in sephaneous vein graft; iv) Patient willing for undergoing percutaneous intervention to saphenous vein graft.

Detailed history and physical examination was done at the time of admission. Before procedure every patient was loaded with 300 mg of clopidogrel and graft study was reviewed. During procedure all patients received intravenous un-fractioned heparin according to body weight. Use of filter device, glycoprotein IIa IIIb inhibitor and type of stent (drug eluting stent or bare metal stent) was at the discretion of operator.

Following points were noted after procedure to evaluate procedural outcome: (i) Acute closure; (ii) slow flow phenomenon; (iii) sub-acute thrombosis of stent; (iv) dissection of target vessel; (iv) perforation of target vessel; (v) procedural failure; (vi) post-procedural cardiac enzymes; (vii) contrast reaction; (viii) Major haematoma formation and need of blood transfusion.

RESULTS

Between November 2006 and October 2009, 113 patients were enrolled. Baseline characteristics are shown in Table-1. Although mean age was 58.2 ± 9.4 years, our study contain both young (age less than 40 years) as well as old patients (age above 75 years). Majority of population was male (96.5%). Among risk factors of atherosclerosis, 73.5% had hyperlipidaemia, 66.4% had family history of IHD, 57.5% were hypertensive, 44.2% were diabetic and 42.5% were smokers. Majority of patients presented with either

unstable angina or stable angina while one third of patients presented with STEMI or NSTEMI.

 Table-1: Baseline characteristics of the study population [n (%)]

Variables	Value
Age (Mean±SD)	58.2±9.4
Gender	
Male	109 (96.5%)
Female	4 (3.5%)
Diabetes	50 (44.2%)
Hypertension	65 (57.5%)
Smoking	48 (42.5%)
Hyperlipidemia	83 (73.5%)
Family History of IHD	75 (66.4%)
Mode of Presentation	
Unstable Angina	34 (30.1%)
Stable Angina	33 (29.2%)
ST elevation MI	20 (17.7%)
NSTEMI	26 (23.0%)

Table-2 shows angiographic and procedural characteristics. Majority of stents were deployed in SVG to OM or RCA (36.3% and 30.1% respectively). Filter wire EZ was used in 14 (12.4%) and Export Aspiration catheter was used in 4 (3.5%). Bare metal stents were used in 58 (51.3%), drug eluting stents were used in 53 (46.9%) and in 2 (1.8%) patients, only balloon angioplasty was done. Among bare metal stents (stainless steel), Flex master, Coro flex and Liberte were used while among drug eluting stents (sirolimus, evorolimus or paclitaxel coated), Xience V, Promus, Cypher and Texus were used.

Table-2:- Angiographic and procedural characteristics

D (Value
Parameter	n (%)
Name of Target vessel: (SVG to)	15 (12 20/)
LAD (Left Anterior Descending)	15 (13.3%)
D1 (First Diagonal)	10 (8.8%)
CX (Circumflex)	3(2.7%)
OM1 (First Obtuse Marginal)	41 (36.3%)
OM2 (Second Obtuse Marginal)	4 (3.5%)
RCA (Right Coronary artery)	34 (30.1%)
PDA (Posterior Descending Artery)	3 (2.7%)
Ramus Inter-medius	3 (2.7%)
Filter Wire or Device used	14 (12 404)
Filter Wire EZ	14 (12.4%)
Export Aspiration Catheter	4 (3.5%)
Type of Stent used	
Drug eluting stent	53 (46.9%)
Bare metal stent	58 (51.3%)
Not used (balloon only)	2 (1.8%)
Brand name of stent	
Flex Master	55 (48.7%)
Coro Flex	2 (1.8%)
Liberte	1 (0.9%)
Xience V	18 (15.9%)
Promus	8 (7.1%)
Cypher	22 (19.5%)
Taxus	5 (4.4%)
Glycoprotein IIa/IIIb Inhibitor	
Tirofiban	19 (16.8%)
Eptifibatide	39 (34.5%)
Abciximab	42 (37.2%)
Clopidogrel	113 (100%)
Mean Inflation Pressure	15.18±3.3
Mean Diameter of stent used	3.0686±0.61
Mean Length of stent used	19.82±6.7

Table-3 shows the peri-procedural outcome of percutaneous intervention of saphenous vein grafts. The angiographic primary success rate per lesion was 86% for bypass grafts stenosis while adverse outcomes were seen in 16 out of 113 patients. Twenty seven complications occurred in 16 patients. Death occurred in only 1 (0.9%) patient and cause of death in this patient was sub-acute thrombosis of stent. Slow flow phenomenon was observed in 13 (11.5%) patients. Dissection of target vessel occurred in 3 (2.7%) patients. Acute Closure and Perforation occurred in 1 (0.9%) patients each. The Procedural failure occurred in only 1 (0.9%) patient. In this target lesion was calcified and could not be crossed despite use of various wires. The Procedure was abandon in this patient without any complication. Myocardial injury as evidenced by significant increase in cardiac enzymes within 24 hrs of procedure occurred in 6 (5.3%) patients. Among these 6 patients, 5 patients developed MI after slow flow phenomenon and only one patient had MI due to perforation of target vessel.

Table-3:	Peri-	procedural	Outcome

	No. (%)
Any adverse Outcome	16 (14.2)
Acute Closure	1 (0.9)
Slow flow phenomenon	13 (11.5)
Sub-acute thrombosis of stent	1 (0.9)
Dissection	3 (2.7)
Perforation	1 (0.9)
Procedural failure	1 (0.9)
Post procedural raised CPK	6 (5.3)
Death during hospital stay	1 (0.9)
Emergency CABG	0 (0)
Contrast Reaction	0 (0)
Major Hematoma and need for blood transfusion	0 (0)

DISCUSSION

Coronary artery bypass surgery (CABG) has enabled a large number of patients to obtain 5–10 years of useful life with anginal symptoms relieved or improved. However, symptoms recur or progress in about 5% of patients per year.^{5–8} Percutaneous treatment of SVG obstructions is difficult because it often results in inadequate dilatation, a high likelihood of distal embolization associated with mortality and significant morbidity, and a high restenosis rate.⁹

In patients who had coronary bypass surgery and in which vein grafts were used, majority are reluctant to see a cardiovascular surgeon professionally again after the first CABG operation. Re-operation is technically more difficult and is generally associated with a higher morbidity and mortality than the first surgical procedure. Furthermore, following re-operation 30–40% of patients may not experience improvement in their angina.^{10–15} Thus there is continued need for development of approaches to treat such patients.

Although in our study, mean age is 58.2 ± 9.4 years, some patients are more than 75 years old. There are many unique characteristics of patients treated for

vein graft disease: they are older than when they had their initial procedure, and more importantly, their vein grafts are also older. Beyond 1 year and particularly >3years after surgery, atherosclerosis of vein grafts becomes increasingly prevalent.^{1,16–19} It may coexist with intimal fibro-muscular proliferation and, in patients with acute ischemic syndromes, thrombus of varying ages. Atherosclerotic plaques in vein grafts are often large compared with plaques in native coronary arteries and are usually soft and friable, rich in necrotic debris, cholesterol crystals, blood elements, and foam cells. Thrombus has been documented by angioscopy in up to 70% of vein graft lesions undergoing treatment.²⁰ This substrate is primarily responsible for the problems encountered during interventional procedures, which include distal embolization, inadequate dilatation, increased restenosis rates and reduced ante-grade blood flow leading to slow or no reflow phenomenon. The incidence of peri-operative acute myocardial infarction is reported to range between $2.7-10.6\%^{21}$, which also corresponds to our study, i.e., 5.3%.

Since SVGs lack side branches, distal occlusion results in stasis and thrombus formation throughout the graft, which makes PCI more difficult.²² Thus, as with bypass surgery, outcomes after PCI for SVG stenosis are often worse than in native vessels.²² This was illustrated in a pooled analysis of five randomized glycoprotein IIb/IIIa inhibitor trials in which PCI for SVG stenosis was associated with significant increase in mortality at both 30 days (2.1 versus 1.0 % compared to PCI in native vessels) and six

months (4.7 versus 2.0 %).²³ Mortality rate in our study is 0.9% which is less than other studies. This may be due to better case selection as well as use of stents, filter devices and adjunctive medications.

The mechanism of slow or no-reflow phenomenon is complex and various theories have been proposed. This phenomenon is probably due to a combination of multiple patho-physiological mechanisms which may have a different role in different clinical and procedural settings. The postulates include: spasm of the distal microcirculation and recently the distal embolization of pieces of lipid rich plaque.²⁴ Slow flow phenomenon complicates 10–15% of PCI in SVG.²⁴ In our study, it occurred in 11.5% of cases which compare favourably with other reports.

It is especially important that graft problems be investigated early; since stenosis in these vessels can progress rapidly to total thrombotic occlusion.²⁵ A heavy thrombus burden in an SVG markedly increases the difficulty in obtaining a satisfactory result using a catheter technique. In our study procedural failure occurred in only 1 (0.9%) patient. In this target lesion was calcified and lesion could not be crossed despite using various wires. Although most patients with recurrent angina due to SVG stenosis can be managed medically, cardiac catheterisation should be performed at the earliest signs of recurrent ischemia to detect critical graft lesions that can be treated before irreversible loss of the graft.

Table-4 compares some of our results with other reports.^{21,26–35}

	Patients	SVG attempted lesions	Primary Success	Myocardial Infarction	Emergency CABG	Deaths
First Author	(n)	(n)	(%)	(%)	(%)	(%)
Meester ²¹	84	93	84	8.3	2.4	1.2
Pinkerton ²⁶	236	100	93	NA	3	0.4
Block ²⁷	40	40	78	0	2.5	NA
Dorros ²⁸	61	33	79	4.9	1.6	3.3
Douglas ²⁹	116	62	94	1.7	2.6	0
El Gamal ³⁰	31	44	93	6.5	0	0
Cote ³¹	82	101	85	3.7	1.2	0
Corbelli ³²	94	47	92	2.1	4.3	1.1
Ernst ³³	83	33	97	2.4	0	0
Reeder ³⁴	19	19	84	5.3	0	5.3
Cooper ³⁵	59	24	75	5.1	0	1.7
Present Study	113	113	86	5.3	0	0.9

Table-4: Comparison of results with other studies

CONCLUSION

Percutaneous revascularisation of the saphenous vein grafts can be performed with high procedural success and low rate of in-hospital complications and is a good alternative to re-operation in some patients with previous bypass surgery. Success rates and survival have improved with time as a result of improvements in techniques and greater use of stents, filter devices and adjunctive medications in patients who undergo percutaneous intervention following CABG surgery.

REFERENCES

- Bourassa MG, Enjalbert M, Campeau L, Lesperance J. Progression of atherosclerosis in coronary arteries and bypass grafts. Ten years later. Am J Cardiol 1984;53:102C–107C.
- Johnson WD, Kayser KL, Pednaza PM: Angina pectoris and coronary bypass surgery: Patterns of prevalence and recurrence in 3105 consecutive patients followed upto 11 years. Am Heart J 1984;108:1190–7.
- Savage MP, Douglas JS Jr, Fischman DL, Pepine CJ, King SB 3rd, Werner JA, *et al.* Stent placement compared with balloon angioplasty for obstructed coronary bypass grafts. N Engl J Med 1997;337:740-7.

- Verghese M, Berger PB, Lennon RJ, Gersh BJ, Holmes DR, Jr. Comparison of percutaneous interventions for unstable angina pectoris with and without previous coronary artery bypass grafting. Am J Cardiol 2000;86:931–7.
- Laird-Meeter K, ten Katen HJ, Brower RW, van den Brand MJ, Serruys PW, Haalebos MM, *et al.* Angina pectoris, 1 to 10 years after aortocoronary bypass surgery. Eur Heart J 1983;5:35–42.
- Brower R, Laird-Meeter K, Serruys P, Meester G, Hugenholtz PG. Long-term follow-up after coronary artery bypass graft surgery: progression and regression of disease in native coronary circulation and bypass grafts. Br Heart J 1983;50:42–7.
- Campeau L, Lesperance J, Hermann J, Corbara F, Grondin CM, Bourassa MG. Loss of improvement of angina between 1 and 7 years after aortocoronary bypass surgery: correlations with changes in vein grafts and in coronary arteries. Circulation 1979;60:11–5.
- Dorros G, Johnson WD, Tector AJ, Schmahl TM, Kalush SL, Janke L. Percutaneous transluminal coronary angioplasty in patients with prior coronary artery bypass grafting. J Thorac Cardiovasc Surg 1984;87:17–26.
- de Feyter PJ, van Suylen RJ, de Jaegere PP, Topol EJ, Serruys PW. *et al.* Balloon angioplasty for the treatment of lesions in saphenous vein bypass grafts. J Am Coll Cardiol 1993;21:1539–49.
- Reul GJ, Cooley DA, Ott DA, Coelho A, Chapa L, Eterovic I. Reoperation for recurrent coronary artery disease. Arch Surg 1979;14:1269–75.
- Vouhe P, Grondin CM. Reoperation for coronary graft failure: clinical and angiographic results in 43 patients. Ann Thorac Surg 1979;328–34.
- Loop RD, Cosgrove DM, Kramer JR, Lytle BW, Taylor PC, Golding LA, *et al.* Late and arteriographic results in 500 coronary artery reoperations. J Thorac Cardiovasc Surg 1981;675–85.
- Krause AH, Page US, Bigelow JC, Okies JE, Dunlap SF. Reoperation in symptomatic patients after direct coronary artery revascularization. J Thorac Cardiovasc Surg 1978;75:499–504.
- Lytle BW, Loop FD, Cosgrove DM, Taylor PC, Goormastic M, Peper W, *et al.* Fifteen hundred coronary reoperation: results and determinants of early and late survival. J Thorac Cardiovasc Surg. 1987 Jun;93:847-59.
- Laird-Meeter K, van Domburg R, van den Brand M, Lubsen J, Bos E, Hugenholtz PG. Incidence, risk and outcome of reintervention after aortocoronary bypass surgery. Br Heart J 1987;57:427–35.
- Kalan JM, Roberts WC. Morphologic changes in saphenous veins used as coronary arterial bypass conduits for longer than one year: necropsy analysis of 53 patients, 123 saphenous veins and 1865 five millimeter segments of veins. Am Heart J 1990;119:1164–84.
- Bourassa MG. The national history of saphenous vein bypass graft disease. In: Bates ER, Holmes DR, Eds. Saphenous Vein Bypass Graft Disease. New York, NY: Marcel Dekker; 1998.p. 61–76.
- Nwasokwa ON. SsCoronary artery bypass graft disease. Ann Intern Med 1995;123:528–45.
- 19. Lawrie GM, Lie JT, Movis GC, Beagley HC. Vein graft patency and intimal proliferative after aortocoronary bypass: early and

long-term angiopathologic correlations. Am J Cardiol 1976;38:856-62.

- White CJ, Ramee SR, Collins TJ, Mesa JE, Jain A. Percutaneous angioscopy of saphenous vein coronary bypass grafts. J Am Coll Cardiol 1993;21:1181–5.
- Meester B J, Samson M, Suryapranata *et al.* Long-term followup after attemted angioplasty of saphenous vein grafts: the Thoraxcenter experience 1981–88. Eur Heart J 1991;12:648–53.
- 22. Stone GW; Reifart NJ; Moussa I, Hoye A, Cox DA, Colombo A, *et al* Percutaneous recanalization of chronically occluded coronary arteries: a consensus document: part II. Circulation 2005;112:2530–7.
- 23. Roffi M; Mukherjee D; Chew DP, Bhatt DL, Cho L, Robbins MA, *et al.* Lack of benefit from intravenous platelet glycoprotein IIb/IIIa receptor inhibition as adjunctive treatment for percutaneous interventions of aortocoronary bypass grafts: a pooled analysis of five randomized clinical trials. Circulation 2002;106:3063–7.
- 24. Sharma S. Current Management Of Saphenous Vein Graft Disease. Internet J Cardiol 2004;2:2 (internet edition)
- Chen L; Theroux P; Lesperance J, Shabani F, Thibault B, De Guise P. Angiographic features of vein grafts versus ungrafted coronary arteries in patients with unstable angina and previous bypass surgery. J Am Coll Cardiol 1996;28(6):1493–9.
- Pinkerton CA, Slack JD, Orr CM, Vantassel JW, Smith ML. Percutaneous transluminal angioplasty in patients with prior myocardial revascularization surgery. Am J Cardiol 1988;61:15G–22.
- Block PC, Cowley MJ, Kaltenbac h M, Kent KM, Simpson J. Percutaneous angioplasty of stenoses of bypass grafts or of bypass graft anastomotic sites. Am J Cardiol 1984;53:666–8.
- Dorros G, Johnson WD, Tector AJ, Schmahl TM, Kalush SL, Janke L. Percutaneous transluminal coronary angioplasty in patients with prior coronary artery bypass grafting. J Thorac Cardiovasc Surg 1984;87:17–26.
- Douglas JS, Gruentzig AR, King SB, Hollman J, Ischinger T, Meier B, *et al.* Percutaneous transluminal angioplasty in patients with prior coronary artery bypass surgery. J Am Coll Cardiol 1983;745–54.
- El Gamal M, Bonnier H, Michels R, Heijman J, Stassen E. Percutaneous transluminal angioplasty of stenosed aortocoronary bypass grafts. Br Heart J 1984;52:617–20.
- Cote G, Myler RK, Stertzer SH, Clark DA, Fishman-Rosen J, Murphy M, *et al.* Percutaneous transluminal angioplasty of stenotic coronary artery bypass grafts: 5 year's experience. J Am Coll Cardiol 1987;9:8–17.
- Corbelli J, Franco I, Hollman J, Simpfendorfer C, Galan K. Percutaneous transluminal angioplastyafter previous coronary artery bypass surgery. Am J Cardiol 1985;56:398–403.
- Ernst SM, Vander Feltz TA, Ascoop CA, Bal ET, Vermeulen FE, Knaepen PJ, *et al.* Percutaneous transluminal coronary angioplasty in patients with prior coronary artery bypass grafting: long-term results. J Thorac Cardiovasc Surg 1987;93:268–75.
- Reeder GS, Bresnahan JF, Holmes DR, Mock MB, Orszulak TA, Smith HC, *et al.* Angioplasty for aortocoronary bypass graft stenosis Mayo Clin Proc 1986;61:14–9.
- Cooper I, Ineson N, Demirtas E, Coltart J, Jenkins S, Webb-Peploe M. Role of angioplasty in patients with previous coronary artery bypass surgery. Cathet Cardiovasc Diagn 1989;16:81–6.

Address for Correspondence:

Dr. Tariq Shakoor, 2 Makkah Street, Muslim Colony, Samanabad, Lahore, Pakistan. **Cell:** +92-300-9418546 **Email:** drtariqshakoor@hotmail.com