

Secondary prevention is as important as surgical technique

Sigrid Sandner MD¹  | Marc Ruel MD, MPH²

¹Department of Cardiac Surgery, Medical University of Vienna, Vienna, Austria

²Division of Cardiac Surgery, University of Ottawa Heart Institute, Ottawa, Canada

Correspondence: Sigrid Sandner, MD, Department of Cardiac Surgery, Medical University of Vienna, Spitalgasse 23, A1090 Vienna, Austria.
Email: sigrid.sandner@meduniwien.ac.at

KEYWORDS

secondary prevention, vein graft disease, vein graft failure

Coronary artery bypass grafting (CABG) is the treatment of choice for patients with complex coronary artery disease. In the U.S. alone, approximately 300,000 patients undergo CABG annually. Saphenous vein grafts (SVG) are used in more than 90% of CABG procedures.¹ SVGs have a higher rate of failure than arterial grafts, with contemporary rates of SVG occlusion of 6% at 1 month, 10% at 3–6 months, and 13% at 6–12 months.² Thrombosis and technical failure is the predominant pathophysiologic mechanism of SVG failure within the first week and during the first month after CABG, followed by intimal hyperplasia from 1 month to 1 year, and atherosclerosis beyond 1 year.³ A multitude of studies have focused on determining factors associated with SVG occlusion during the first year after CABG.

In this issue of the *Journal of Cardiac Surgery*, Yang et al.⁴ assessed vein graft disease (VGD) progression between 1 week and 1 year after CABG. The current report draws on a previously described cohort of 233 patients who underwent computed tomographic angiography (CTA) 1 week after off-pump CABG,⁵ and includes 218 of these patients (654 SVG) who subsequently underwent CTA at 1 year. The authors found that, overall, VGD progression between 1 week and 1 year occurred in 11.3% of SVG, and 22.1% of patients. VGD progression was particularly evident in those SVG that already demonstrated VGD 1 week after CABG and occurred in 38.6% of SVG with Fitzgibbon B patency, whereas only 11.3% of perfectly patent SVG (Fitzgibbon A) demonstrated signs of VGD at 1 year, a difference that was statistically significant.

In a novel approach to examine the association of the success (or lack thereof) of their implementation of secondary prevention measures with VGD progression, the authors generated a risk

score derived from their assessment of patients achieving defined targets of lipid levels, glucose control, smoking cessation, and hypertension control 1 year after CABG. The failure to meet any one secondary prevention target contributed one point to the risk score, which was included as a variable in the logistic regression model. The authors demonstrated that the risk score was significantly associated with VGD progression both at the graft (odds ratio [OR] = 1.38; *p* for trend = .01) and the patient level (OR = 1.52, *p* for trend = .01).

The findings of Yang et al. underscore the importance of secondary prevention after CABG. To date, only the use of postoperative antiplatelet therapy and statins has been shown in randomized clinical trials (RCT) to reduce SVG failure. A 1993 meta-analysis of seven RCTs including 1443 patients showed that the use of aspirin after CABG was associated with a lower risk of SVG occlusion.⁶ A recent meta-analysis involving 1316 patients and 1668 SVG showed that adding ticagrelor to aspirin was associated with a significantly lower incidence of SVG failure (11.2%) per graft than was aspirin (20%; difference, -8.7% [95% confidence interval, CI, -13.5% to -3.9%]; OR, 0.51 [95% CI, 0.35–0.74]; *p* < .001).⁷ Aggressive lipid-lowering reduced progression of atherosclerosis in grafts in the post-CABG trial.⁸ In addition, higher low-density lipoprotein (LDL) levels during the first postoperative year were associated with significantly greater VGD 3 years after CABG in the aggressive cholesterol therapy to inhibit vein graft events trial.⁹ Diabetes mellitus and active smoking have been postulated among the predictors for early SVG occlusion.² In this regard, the current work confirms that lack of glycemic control and failure of smoking cessation correlates with progression of VGD between 1 week and 1 year after CABG.

This is an open access article under the terms of the Creative Commons Attribution-NonCommercial-NoDerivs License, which permits use and distribution in any medium, provided the original work is properly cited, the use is non-commercial and no modifications or adaptations are made.

© 2022 The Authors. *Journal of Cardiac Surgery* published by Wiley Periodicals LLC.

In an analysis from the SWEDEHEART registry, treatment with statins (hazard ratio [HR]: 0.56, 95% CI, 0.52–0.60), RAAS inhibitors (HR: 0.78, 95% CI, 0.73–0.84), and platelet inhibitors (HR: 0.74, 95% CI, 0.69–0.81) were individually associated with lower mortality risk after CABG.¹⁰ In a subanalysis of the Synergy between PCI With Taxus and cardiac surgery Extended Survival study,¹¹ patients on optimal medical therapy at 5 years had significantly lower mortality at 10 years compared with those on ≤ 2 types of medications (13.1% vs. 19.9%; adjusted HR: 0.470; 95% CI, 0.292–0.757; $p = .002$). An astonishing finding in the current work was that despite 96.3% of patients receiving statins 1 year after CABG, 56.4% of patients did not meet the study-specified target LDL level of < 1.8 mmol/L, which in addition remains well above the European Society of Cardiology guideline-recommended target of < 1.4 mmol/L for secondary prevention of cardiovascular disease.¹²

On the basis of the data of Yang and colleagues, we are reminded that, very likely, a large proportion of patients remain undertreated in terms of secondary prevention after CABG, and that optimal medical therapy is as important as the surgical procedure itself.

CONFLICT OF INTEREST

The authors declare no conflict of interest.

ORCID

Sigrid Sandner  <http://orcid.org/0000-0003-4669-9841>

REFERENCES

- Bowdish ME, D'agostino RS, Thourani VH, et al. STS adult cardiac surgery database: 2021 update on outcomes, quality, and research. *Ann Thorac Surg.* 2021;111:1770-1780.
- Antonopoulos AS, Odutayo A, Oikonomou EK, et al. Development of a risk score for early saphenous vein graft failure: an individual patient data meta-analysis. *J Thorac Cardiovasc Surg.* 2020;160:116-127.
- Xenogiannis I, Zenati M, Bhatt DL, et al. Saphenous vein graft failure: from pathophysiology to prevention and treatment strategies. *Circulation.* 2021;144:728-745.
- Yang Y. Predictors of vein graft disease progression between one week and one year after surgical coronary revascularization: impact of secondary prevention medications. *J Card Surg.* 2022.
- Yang Y, Zhu Y, Tang C, et al. Predictors of early vein graft failure after off-pump coronary artery bypass grafting: angiocomputed tomographic results of 233 patients. *Eur J Cardiothorac Surg.* 2020;57:277-284.
- Fremes SE, Levinton C, Naylor CD, Chen E, Christakis GT, Goldman BS. Optimal antithrombotic therapy following aortocoronary bypass: a meta-analysis. *Eur J Cardiothorac Surg.* 1993;7:169-180.
- Sandner S, Redfors B, Angiolillo DJ, et al. Association of dual antiplatelet therapy with ticagrelor with vein graft failure after coronary artery bypass graft surgery: a systematic review and meta-analysis. *JAMA.* 2022;328:554-562.
- Post Coronary Artery Bypass Graft Trial Investigators. The effect of aggressive lowering of low-density lipoprotein cholesterol levels and low-dose anticoagulation on obstructive changes in saphenous-vein coronary-artery bypass grafts. *N Engl J Med.* 1997;336:153-162.
- Kulik A, Abreu AM, Boronat V, Ruel M. Intensive versus moderate statin therapy and early graft occlusion after coronary bypass surgery: the aggressive cholesterol therapy to inhibit vein graft events randomized clinical trial. *J Thorac Cardiovasc Surg.* 2019;157:151-161.
- Björklund E, Nielsen SJ, Hansson EC, et al. Secondary prevention medications after coronary artery bypass grafting and long-term survival: a population-based longitudinal study from the SWEDEHEART registry. *Eur Heart J.* 2020;41:1653-1661.
- Kawashima H, Serruys PW, Ono M, et al. Impact of optimal medical therapy on 10-year mortality after coronary revascularization. *J Am Coll Cardiol.* 2021;78:27-38.
- Mach F, Baigent C, Catapano AL, et al. 2019 ESC/EAS guidelines for the management of dyslipidaemias: lipid modification to reduce cardiovascular risk. *Eur Heart J.* 2020;41(41):111-188.

How to cite this article: Sandner S, Ruel M. Secondary prevention is as important as surgical technique. *J Card Surg.* 2022;37:3673-3674. doi:10.1111/jocs.16896