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Reply

We greatly appreciate the interest in our paper (1) and the astute comments by Drs. Tsigkas, Xanthopoulos, and Alexopoulos who highlight the use of embolic protection devices (EPDs) in saphenous vein graft (SVG) percutaneous coronary interventions (PCI) and the importance of the amount of myocardium in jeopardy. While we agree that the amount of jeopardized myocardium is an important predictor of outcome, the information necessary to calculate the amount of myocardium in jeopardy during PCI is not available in the CathPCI Registry. EPDs were used in 19.64% of SVG PCIs during the study period. EPD use was not associated with in-hospital mortality in univariable analysis (odds ratio: 0.988, 95% confidence interval: 0.870 to 1.122) or multivariable analysis (odds ratio: 0.935, 95% confidence interval: 0.813 to 1.075). Longer-term follow-up is likely needed to detect an impact from EPD use on clinical outcomes.

EPDs have been proved to reduce the incidence of post-SVG PCI myocardial infarction, and have a Class I indication in the American College of Cardiology/American Heart Association PCI guidelines. Yet, EPDs remain underutilized both in the United States and in Europe (2), due to device complexity, difficulties assessing the embolization risk of each SVG lesion, unavailability of a universally applicable EPD, and lack of reimbursement (3,4).

The similarity of the findings from the population of Xanthopoulos et al. (5) and from NCDR (1) strengthen the conclusion

that native coronary artery PCI is preferable to SVG PCI in prior coronary artery bypass graft patients, if technically feasible. Given the rapid advances in complex PCI techniques, especially chronic total occlusion PCI (6,7), native coronary artery interventions are likely to be increasingly utilized in the future in patients presenting with SVG failure.

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