Saphenous Vein Graft Failure after Coronary Artery Bypass Surgery: Insights from PREVENT IV

Running title: Hess et al.; Predictors of VGF after CABG

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Abstract

**Background**—Coronary artery bypass grafting (CABG) success is limited by vein graft failure (VGF). Understanding factors associated with VGF may improve patient outcomes.

**Methods and Results**—We examined 1828 participants in the PREVENT IV trial undergoing protocol-mandated follow-up angiography 12–18 months post-CABG or earlier clinically-driven angiography. Outcomes included patient- and graft-level angiographic VGF (≥75% stenosis or occlusion). Variables were selected using Fast False Selection Rate methodology. We examined relationships between variables and VGF in patient- and graft-level models using logistic regression without and with generalized estimating equations. At 12–18 months post-CABG, 782 of 1828 (42.8%) patients had VGF, and 1096 of 4343 (25.2%) vein grafts had failed. Demographic and clinical characteristics were similar between patients with and without VGF, though VGF patients had longer surgical times, worse target artery quality, longer graft length, and more frequently underwent endoscopic vein harvesting. After multivariable adjustment, longer surgical duration (odds ratio [OR] per 10-minute increase 1.05, 95% confidence interval [CI] 1.03–1.07), endoscopic vein harvesting (OR 1.41, 95% CI 1.16–1.71), poor target artery quality (OR 1.43, 95% CI 1.11–1.84), and postoperative use of clopidogrel or ticlopidine (OR 1.35, 95% CI 1.07–1.69) were associated with patient-level VGF. The predicted likelihood of VGF in the graft-level model ranged from 12.1–63.6%.

**Conclusions**—VGF is common and associated with a number of patient and surgical factors. These findings may help identify patients with risk factors for VGF and inform the development of interventions to reduce VGF.

**Clinical Trial Registration Information**—ClinicalTrials.gov. Identifier: NCT00042081.

**Key words:** coronary disease, coronary artery bypass graft surgery, revascularization
Coronary artery bypass grafting (CABG) is one of the most frequently performed surgical procedures in the United States, with over 400,000 procedures performed annually. Although CABG improves survival and symptoms in selected patients, surgical success depends on the continued patency of grafts, and graft failure has been associated with worse outcomes. Saphenous vein grafts remain the most widely used conduit during CABG, and rates of vein graft failure (VGF) during the first 12 to 18 months after surgery have been reported to be as high as 25%. Many studies have examined factors associated with VGF and have inconsistently reported associations between multiple clinical and surgical characteristics and VGF. These previous efforts have been limited by the absence of systematic angiographic follow-up. In addition, results from these studies may be outdated, given advances in surgical techniques and adjunctive medical therapies that could impact graft failure. We therefore sought to examine factors associated with VGF assessed by coronary angiography 12–18 months after CABG using data from the PRoject of Ex-vivo Vein graft ENgineering via Transfection IV (PREVENT IV) trial.

Methods

Data source and patient population

We used data from the PREVENT IV trial (ClinicalTrials.gov: NCT00042081), the design and results of which have been previously described. Briefly, PREVENT IV was a phase 3 randomized, double-blind, placebo-controlled trial of ex-vivo vein graft treatment with edifoligide in patients undergoing primary CABG with ≥2 planned vein grafts. A total of 3014 patients were enrolled between August 2002 and October 2003 at 107 centers across the U.S., the
first 2400 of whom were scheduled for follow-up angiography between 12–18 months after
CABG. The PREVENT IV protocol was approved by institutional review boards of all
participating sites and all enrolled patients provided written informed consent.

We included patients in the angiographic cohort who were scheduled to undergo follow-
up angiography 12–18 months after the index CABG (n=2400). Patients in the angiographic
cohort who had VGF documented during earlier angiography for clinical indications in place of
(n=64) or in addition to (n=107) routine protocol angiography were included. We excluded
patients who did not undergo angiographic follow-up (n=477), who received only arterial grafts
(n=4), or who died prior to their 12–18 month repeat angiogram (n=91). Our final analysis
population consisted of 1828 patients enrolled at 100 sites (Figure 1).

Definitions and outcomes

VGF was defined as ≥75% stenosis or occlusion detected at follow-up angiography 12–18
months after CABG or earlier angiography performed for clinical indications. All angiograms
were analyzed at a core laboratory (PERFUSE Angiographic Core Laboratory, Boston, MA). For
grafts with multiple distal anastomoses (m-SVG), failure of any component was considered
VGF.17 Outcomes for our analyses were defined as failure of 1 or more vein grafts (patient-level
angiographic VGF) and graft-level angiographic VGF.

Statistical analysis

Baseline patient and procedure characteristics were examined according to patient-level absence
or presence of VGF at 12–18 months post-CABG. Continuous variables were summarized using
medians and interquartile ranges (IQR), while categorical variables were presented as
frequencies and percentages. Comparisons within continuous and categorical variable groups
were performed using Wilcoxon 2-sample test and Chi-square test, respectively.
We analyzed surgical features at both the patient- and graft-levels. When describing patient-level characteristics, we used the “worst” status to describe procedure characteristics for patients with multiple vein grafts. The following hierarchies (worst status listed first) were used: target artery quality= poor, fair, good; graft quality= poor, fair, good; distal connection technique= non-suture, suture; graft length= longest measurement; graft source= arm vein, lesser saphenous vein, greater saphenous vein; vein harvest technique= endoscopic, open; and m-SVG use= yes, no.

We developed patient- and graft-level models to determine factors associated with VGF. For the main analysis, patient-level variables were created by assessing graft-level data for each patient and, for patients with multiple grafts, determining the worst status for each characteristic among all grafts. We also performed a secondary analysis to examine graft-level variables associated with VGF. For both models, variables associated with VGF were selected using Fast False Selection Rate (Fast FSR).¹⁸ Fast FSR is a conservative variable selection method that accounts for the percentage of variables incorrectly identified as associated with the outcome of interest. Logistic regression models were then fit using the chosen variables to estimate the association of each factor with VGF and odds ratios (OR) with associated 95% confidence intervals (CI) were reported. For graft-level analyses, in order to account for the correlation among multiple grafts within the same patient, generalized estimating equations were used to fit a generalized linear logistic model that allows for an exchangeable correlation matrix between grafts within a single patient.

The following candidate variables were chosen based on clinical judgment and considered for inclusion in both patient- and graft-level models: age, female sex, weight, race, smoking status, chronic lung disease, hypertension, dyslipidemia, prior myocardial infarction,
prior percutaneous coronary intervention, prior cancer, history of liver disease, peripheral artery disease, cerebrovascular disease, prior congestive heart failure, current New York Heart Association class, diabetes (no history, non-insulin therapy, insulin therapy), renal failure, atrial fibrillation/flutter, ejection fraction, type of CABG procedure (emergent/salvage, urgent, elective), use of cardiopulmonary bypass (CPB), CPB time, aortic cross-clamp time, surgical time, graft source (greater saphenous, lesser saphenous), vein harvest technique (endoscopic, open), graft quality, maximum stenosis of target vessel (<75%, ≥75%), target artery quality, proximal anastomosis connection technique (suture, non-suture), graft length, and use of m-SVG. For both patient- and graft-level models, linear splines were used to determine appropriate knot points for the following non-linear variables (see Online supplement for knot points): aortic cross-clamp time, ejection fraction, graft length (patient-level model only), and CPB time (graft-level model only). Significant (p<0.1) levels were then included as candidate variables (see Online supplement). We hypothesized that chronic use of certain medications might be associated with VGF. In PREVENT IV, data regarding medication use were collected at the discrete time points at baseline, discharge, 30 days, and 1 year. We chose to examine 30-day medication use as covariates, as these were thought to best represent chronic postoperative use following the initial surgery. However, since medication use at 30 days is a post-baseline variable, it was included in models as a sensitivity analyses. Rates of missingness for data in our models were ≤1.5%, and no imputation was performed for missing data. Multivariable models were derived from complete cases. For the Fast FSR method, the desired false selection rate was set to 0.05. All analyses were performed at the Duke Clinical Research Institute using SAS version 9.2 (SAS Institute, Cary, NC).
Results

Patient and procedure characteristics

Among a total of 1828 patients included in our study, 782 (42.8%) had VGF at 12–18 months after CABG. At the graft-level, 1096 (25.2%) of the 4343 grafts placed during the index CABG had failed at 12–18 months after CABB. Demographic characteristics and comorbid conditions were similar between patients with and without VGF with the exception of cerebrovascular disease, which was more prevalent among patients with VGF (Table 1).

Patient-level CABG procedure characteristics among patients with and without VGF are shown in Table 2. Compared with patients without VGF, those with VGF had longer surgical and cross-clamp times and worse target artery quality. Patients with VGF also more frequently underwent endoscopic versus open vein graft harvest and had slightly longer graft length than patients without VGF. At 30 days after the index CABG, patients with subsequent VGF were more frequently taking clopidogrel or ticlopidine (26.1% vs. 19.2%, p<0.001) and had similar use of warfarin (9.1% vs. 8.5%, p=0.66) and statins (74.6% vs. 74.9%, p=0.88) than patients who did not have subsequent VGF.

Factors associated with VGF

We first examined patient-level factors associated with VGF at 12–18 months after CABG. Longer duration of surgery (OR per 10-minute increase 1.05; 95% CI 1.03–1.07; p<0.01), endoscopic vein graft harvest technique (OR 1.44; 95% CI 1.19–1.75; p<0.01), and poor target artery quality (OR 1.45; 95% CI 1.13–1.87; p<0.01) were significantly associated with VGF. Adding medications continued at 30 days after CABG to the variable selection model revealed that the use of clopidogrel or ticlopidine was significantly associated with VGF (OR 1.35; 95% CI 1.07–1.69; p=0.01); addition of clopidogrel or ticlopidine to the model did not substantially
change the relationship between the other significant predictors and VGF (Table 3). Goodness of
fit of the model as measured by the Hosmer-Lemeshow statistic indicated that the model fits the
data well (p = 0.85). The c-statistic for the model was 0.61.

Next, we assessed the relationship of graft-level variables with VGF (Table 4). Factors
that were significantly associated with per-graft VGF (Table 4) included fair or poor target
artery quality (OR 1.31; 95% CI 1.11–1.56; p<0.01 and OR 2.34; 95% CI 1.89–2.91; p<0.01,
respectively), longer duration of surgery (OR per 10-minute increase 1.04; 95% CI 1.02–1.05;
p<0.01), endoscopic vein harvest technique (OR 1.37; 95% CI 1.16–1.62; p<0.01), and history of
cerebrovascular disease (OR 1.39; 95% CI 1.06–1.81; p=0.02). After including 30-day
medication use, clopidogrel or ticlopidine use was again associated with VGF (OR 1.30; 95% CI
1.07–1.58; p<0.01).

**Distribution of predicted VGF risk**

We examined the distribution of predicted VGF risk using the full (including 30-day medication
use) graft-level model of VGF. Predicted probability of VGF at 12–18 months post-CABG
ranged from a low of 12.1% to a high of 63.6%. The median predicted risk of VGF among our
patient cohort was 23.4% (interquartile range 19.5% to 29.2%) (Figure 2).

**Discussion**

In this analysis from PREVENT IV which included over 1800 patients, more than 4300
implanted vein grafts, and systematic 12–18 month angiographic follow-up, we found that longer
duration of surgery, endoscopic vein graft harvesting, poor target artery quality, and the use of
clopidogrel or ticlopidine at 30 days post-CABG were factors associated with VGF in both per-
patient- and per-graft-level models. The broad range of predicted VGF using our per-graft-level
model (12.1–63.6%) suggests that VGF is prevalent and hence, these data may be clinically useful to inform efforts to reduce VGF.

Interest in understanding factors associated with VGF after CABG has been longstanding, but prior efforts have been limited.15 Previous studies have consistently reported 1 year VGF rates of 10–20%, with another 5–10% of vein grafts failing between 1–5 years after CABG.10,19-24 These studies have identified patient characteristics, including younger age,11,12 female sex,12,13 prior heart failure or low ejection fraction,12,13 and increased serum cholesterol,11,25 as predictors of VGF. Surgical factors, including temperature of graft solution,25 multiple distal anastamoses,13,26 poor distal vessel,13,26 target artery stenosis,12 and endoscopic harvest technique,26,27 have also been identified as predictive of VGF. Importantly, these analyses were based on data from patients undergoing CABG several decades ago, prior to the widespread use of antiplatelet therapy and the introduction of newer surgical CABG techniques.28-30 Some prior reports were also based on single-center studies, reducing the generalizability of their results, or analyzed data at either the patient- or graft-level, which may account for some of the inconsistency in previous findings. Furthermore, a number of prior studies examined patients undergoing clinically-driven coronary angiography, which may under or overestimate the rate and influence of factors associated with VGF.

Our study extends knowledge in the field in several ways. First, this analysis represents one of the largest analyses of factors associated with VGF to date and includes data from over 100 sites. Second, our study included patients undergoing angiography for clinical reasons as well as relatively complete, protocol-mandated follow-up angiography, allowing for a more unbiased assessment of VGF and the factors associated with it. Third, our analysis was based on data representing more contemporary practice and was strengthened by the detailed clinical and
procedural data that were collected for PREVENT IV. Finally, whereas prior studies have assessed VGF at either the graft- or patient-level, we examined both, as each provides useful and potentially different information. We found that the factors associated with VGF in patient-and graft-level models were almost identical.

We found a number of surgical factors that were associated with VGF. Pathologic studies have demonstrated that atherosclerosis is the main etiology of late (more than 12 months) VGF, whereas early (less than 1 month) and subacute (up to 12 months) graft failure is due to thrombosis, surgical technical errors, and intimal hyperplasia. Intraoperative processes of vein graft harvesting, graft manipulation, and graft implantation can all lead to endothelial dysfunction, inflammation, and ultimately thrombosis and graft occlusion. Accordingly, there is mechanistic feasibility to explain our study results. Longer duration of surgery may reflect technical difficulty, thus contributing to risk of VGF. Endoscopic vein graft harvesting, though less invasive than open vein graft harvesting, can damage vein graft endothelium, causing inflammation and thrombosis with early graft failure or increased intimal hyperplasia and subacute VGF. Observational data regarding the benefits of endoscopic vein harvesting are mixed, with some studies reporting associations of this technique with VGF and worse outcomes, while others have not confirmed these findings. Definitively determining whether endoscopic graft harvesting is associated with VGF will require a prospective randomized clinical study. The Randomized Endo-Vein Graft Prospective (REGROUP) Trial (ClinicalTrials.gov: NCT01850082) which is currently under development will provide important insight into this topic.

We also found that poor target artery quality was associated with VGF. In PREVENT IV, assessments of target artery quality were based on qualitative surgeon judgment and not
systematic classification. However, this qualitative rating likely incorporates the elements of smaller vessel diameter that might reflect challenging surgical anatomy and poor distal run-off, which has been previously associated with VGF.\(^7\)

Two of the factors significantly associated with VGF in our analyses were not related to the surgical procedure. The first was a clinical history of cerebrovascular disease, which was associated with VGF in the graft-level model. Cerebrovascular disease may represent a marker of both more advanced vascular disease and also poor target vessel distal run-off. We also found that use of clopidogrel or ticlopidine at 30 days was associated with an increased risk of VGF. Given the pathologic contribution of thrombosis to early VGF, antiplatelet therapy would be expected to reduce VGF, and randomized data support the use of aspirin to reduce graft failure.\(^{35,36}\) In this study, since use of antiplatelet therapy was not randomized, we hypothesize that the relationship between antiplatelet therapy and VGF is likely due to confounding. Data to support the use of clopidogrel to improve early venous graft patency after CABG are limited,\(^{29,37}\) and clopidogrel is more frequently prescribed to patients with acute coronary syndrome, patients undergoing off-pump CABG, or patients with extensive coronary artery disease.\(^{38,39}\)

In our study, the majority of VGF events were clinically silent. Only 7.1% of the patients with VGF had VGF identified during early repeat angiography for clinical indications. However, studies have demonstrated that VGF identified either during clinically-driven or routine follow-up angiography is associated with significant morbidity.\(^{4,5,10,41}\) Thus, reducing overall VGF after CABG is an important goal that may improve patient outcomes and the durability of CABG surgery.

Research efforts to date have focused on a multifaceted approach to prevent VGF, including modifications in patient behavior, especially smoking cessation, and exploration of
optimal postoperative antiplatelet regimens, as a large proportion of CABG patients are resistant to aspirin.\textsuperscript{15} Given the wide range of predicted VGF risk of our model, these data might help to identify patients at higher risk for VGF who might be considered for CABG with non-vein graft conduits and who should be followed more closely for post-CABG VGF events. However, some of the factors associated with VGF in our study are non-modifiable, suggesting that the greatest use of our data may be to help direct further research into strategies to prevent VGF. The high rate of VGF also emphasizes the importance of investigational surgical techniques to reduce vein graft injury, such as external vein graft support through either stenting or fibrin glue, exploration of novel gene-based molecular therapies to reduce VGF, and the development of synthetic, non-vein graft conduits.\textsuperscript{15}

**Limitations**

This is a retrospective, post-hoc analysis. We assessed VGF at routine angiography 12–18 months after CABG, and the predictors of VGF may change over time. We were not able to assess VGF in patients who died prior to angiography or who did not return for protocol-mandated angiography and have excluded these patients from the analysis. We chose to study VGF and did not include arterial conduits in our analysis. The factors associated with arterial graft failure may differ.\textsuperscript{19,20,42} Some other factors that have previously been associated with vein graft patency were not collected in PREVENT IV.\textsuperscript{11,28,30,35} PREVENT IV only included patients undergoing first-time CABG, and the vein graft handling techniques and pressurized delivery system used in PREVENT IV were unique to the trial. Although our models fit the data well (Hosmer-Lemeshow p=0.85), there was low discriminatory power (C-statistic 0.61). We also included use of clopidogrel and ticlopidine in sensitivity analyses, though these were post-baseline variables that might be associated with non-VGF factors. We were not able to account
for clustering by specific surgeon, as these data were not available. Finally, it should be recognized that both the study timeframe and identification of VGF based on routine angiography impacted the selection of collected data elements, and strategies to reduce VGF have evolved since the time of this study\(^{15}\); all of these factors may limit the generalizability of our results.

**Conclusions**

VGF is common and associated with both patient and surgical factors including, poor target artery quality, longer duration of surgery, use of endoscopic vein harvesting, use of clopidogrel or ticlopidine, and cerebrovascular disease. These data may be useful in identifying patients with risk factors for VGF and to inform the development of strategies to prevent VGF. Further investigation of VGF should be pursued in contemporary datasets.

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Johnson & Johnson, Ortho-McNeil-Janssen Pharmaceuticals, Inc., Pfizer, Sanofi-Aventis, and WebMD (all modest). Dr. Alexander reports consulting for Sohmalution and Moeræ Matrix (all modest). The remaining authors have no conflicts to disclose.

References:


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**Table 1.** Baseline patient characteristics according to presence or absence of VGF.

<table>
<thead>
<tr>
<th>Characteristic</th>
<th>With VGF (n=782)</th>
<th>Without VGF (n=1046)</th>
<th>P Value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Age, median (IQR), yrs</td>
<td>63.0 (55.0–69.0)</td>
<td>63.0 (55.0–70.0)</td>
<td>0.62</td>
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<tr>
<td>Female sex</td>
<td>158 (20.2)</td>
<td>184 (17.6)</td>
<td>0.16</td>
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<td>Weight, median (IQR), kg</td>
<td>88.7 (77.0–100.0)</td>
<td>88.0 (78.0–100.0)</td>
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<td>Race: White</td>
<td>701 (89.6)</td>
<td>954 (91.2)</td>
<td>0.26</td>
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<td>AF/flutter</td>
<td>54 (6.9)</td>
<td>60 (5.7)</td>
<td>0.31</td>
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<tr>
<td>Cancer</td>
<td>72 (9.2)</td>
<td>77 (7.4)</td>
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</tr>
<tr>
<td>Prior CHF</td>
<td>52 (6.6%)</td>
<td>69 (6.6%)</td>
<td>0.96</td>
</tr>
<tr>
<td>Cerebrovascular disease</td>
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<td>88 (8.4%)</td>
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<td>Diabetes mellitus</td>
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<tr>
<td>No diabetes</td>
<td>489 (62.5%)</td>
<td>678 (64.9%)</td>
<td></td>
</tr>
<tr>
<td>Diabetes, no current treatment</td>
<td>14 (1.8%)</td>
<td>23 (2.2%)</td>
<td></td>
</tr>
<tr>
<td>Diabetes, insulin treatment</td>
<td>85 (10.9%)</td>
<td>77 (7.4%)</td>
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</tr>
<tr>
<td>Diabetes, non-insulin treatment</td>
<td>194 (24.8%)</td>
<td>267 (25.6%)</td>
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</tr>
<tr>
<td>EF, median (IQR), %</td>
<td>50.0 (40.0–60.0)</td>
<td>52.5 (43.0–60.0)</td>
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<td>Hypercholesterolemia</td>
<td>169 (21.6)</td>
<td>254 (24.3)</td>
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<tr>
<td>Hypertension</td>
<td>574 (73.4)</td>
<td>760 (72.7)</td>
<td>0.72</td>
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<td>History of liver disease</td>
<td>16 (2.0)</td>
<td>17 (1.6)</td>
<td>0.50</td>
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<td>Chronic lung disease</td>
<td>101 (12.9)</td>
<td>146 (14.0)</td>
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<td>NYHA class</td>
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<tr>
<td>I</td>
<td>312 (40.4)</td>
<td>427 (41.1)</td>
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<td>PAD</td>
<td>87 (11.1)</td>
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<td>History of renal failure</td>
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<td>Smoking status</td>
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<tr>
<td>Never</td>
<td>257 (32.9)</td>
<td>339 (32.4)</td>
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<tr>
<td>Former</td>
<td>345 (44.1)</td>
<td>483 (46.2)</td>
<td></td>
</tr>
<tr>
<td>Current</td>
<td>180 (23.0)</td>
<td>224 (21.4)</td>
<td></td>
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<tr>
<td>Prior MI</td>
<td>343 (43.9)</td>
<td>432 (41.3)</td>
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<tr>
<td>Prior PCI</td>
<td>220 (28.1)</td>
<td>279 (26.7)</td>
<td>0.49</td>
</tr>
</tbody>
</table>

Data presented as no. (%), unless otherwise indicated.

AF indicates atrial fibrillation; CHF, congestive heart failure; EF, ejection fraction; IQR, interquartile range; MI, myocardial infarction; NYHA, New York Heart Association; PAD, peripheral artery disease; PCI, percutaneous coronary intervention; VGF, vein graft failure.
Table 2. Baseline procedural characteristics at the patient-level according to presence or absence of VGF.

<table>
<thead>
<tr>
<th>Characteristic</th>
<th>With VGF (n=782)</th>
<th>Without VGF (n=1046)</th>
<th>P Value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Angiographic classification</td>
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</tr>
<tr>
<td>Per protocol angiography only</td>
<td>655 (83.8)</td>
<td>1002 (95.8)</td>
<td></td>
</tr>
<tr>
<td>Early angiography only</td>
<td>64 (8.2)</td>
<td>0 (0.0)</td>
<td></td>
</tr>
<tr>
<td>Early and per protocol angiographies</td>
<td>63 (8.1)</td>
<td>44 (4.2)</td>
<td></td>
</tr>
<tr>
<td>Maximum stenosis of any target vessel ≥75%</td>
<td>790 (72.3)</td>
<td>2317 (71.5)</td>
<td>0.61</td>
</tr>
<tr>
<td>Endoscopic vein harvest technique</td>
<td>468 (60.1)</td>
<td>531 (50.9)</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td>Any use of composite graft</td>
<td>286 (36.6)</td>
<td>344 (32.9)</td>
<td>0.10</td>
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<tr>
<td>Longest graft length, median (IQR), cm</td>
<td>17.0 (14.3–19.3)</td>
<td>16.0 (14.0–19.0)</td>
<td>0.02</td>
</tr>
<tr>
<td>Any proximal (non-suture)</td>
<td>21 (2.7)</td>
<td>19 (1.8)</td>
<td>0.21</td>
</tr>
<tr>
<td>Any distal (non-suture)</td>
<td>23 (2.9)</td>
<td>27 (2.6)</td>
<td>0.65</td>
</tr>
<tr>
<td>Graft source*</td>
<td></td>
<td></td>
<td>0.32</td>
</tr>
<tr>
<td>Arm vein</td>
<td>0 (0.0)</td>
<td>2 (0.2)</td>
<td></td>
</tr>
<tr>
<td>Lesser saphenous</td>
<td>12 (1.5)</td>
<td>22 (2.1)</td>
<td></td>
</tr>
<tr>
<td>Greater saphenous</td>
<td>770 (98.5)</td>
<td>1022 (97.7)</td>
<td></td>
</tr>
<tr>
<td>Worst target artery quality</td>
<td></td>
<td></td>
<td>&lt;0.01</td>
</tr>
<tr>
<td>Good</td>
<td>308 (39.4)</td>
<td>484 (46.3)</td>
<td></td>
</tr>
<tr>
<td>Fair</td>
<td>281 (36.0)</td>
<td>363 (34.7)</td>
<td></td>
</tr>
<tr>
<td>Poor</td>
<td>192 (24.6)</td>
<td>198 (18.9)</td>
<td></td>
</tr>
<tr>
<td>Worst graft quality</td>
<td></td>
<td></td>
<td>0.12</td>
</tr>
<tr>
<td>Good</td>
<td>537 (68.7)</td>
<td>764 (73.1)</td>
<td></td>
</tr>
<tr>
<td>Fair</td>
<td>206 (26.3)</td>
<td>231 (22.7)</td>
<td></td>
</tr>
<tr>
<td>Poor</td>
<td>39 (5.0)</td>
<td>44 (4.2)</td>
<td></td>
</tr>
<tr>
<td>Use of cardiopulmonary bypass</td>
<td>617 (78.9)</td>
<td>825 (78.9)</td>
<td>0.99</td>
</tr>
<tr>
<td>Pump time, median (IQR), min</td>
<td>95.0 (62.0–123.0)</td>
<td>86.0 (51.0–111.0)</td>
<td>&lt;.0001</td>
</tr>
<tr>
<td>Cross-clamp time, median (IQR), min</td>
<td>60.0 (33.0–78.0)</td>
<td>53.0 (30.0–72.0)</td>
<td>0.01</td>
</tr>
<tr>
<td>Surgical time, median (IQR), min</td>
<td>240.0 (201.0–284.0)</td>
<td>221.0 (186.0–261.0)</td>
<td>&lt;.0001</td>
</tr>
<tr>
<td>Type of procedure</td>
<td></td>
<td></td>
<td>0.66</td>
</tr>
<tr>
<td>Emergent/salvage</td>
<td>20 (2.6)</td>
<td>32 (3.1)</td>
<td></td>
</tr>
<tr>
<td>Urgent</td>
<td>373 (47.7)</td>
<td>480 (45.9)</td>
<td></td>
</tr>
<tr>
<td>Elective</td>
<td>389 (49.7)</td>
<td>533 (51.0)</td>
<td></td>
</tr>
</tbody>
</table>

Data presented as no. (%), unless otherwise indicated.
IQR indicates interquartile range; VGF, vein graft failure.
* For patients with multiple graft sources, the “worst” source according to the following hierarchy was used (worst status listed first): arm vein, lesser saphenous vein, greater saphenous vein.
### Table 3. Factors associated with patient-level VGF.

<table>
<thead>
<tr>
<th>Variable</th>
<th>Chi-Square</th>
<th>OR</th>
<th>95% CI</th>
<th>P Value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Without 30-day medications*</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Duration of surgery (per 10-min increase)</td>
<td>34.66</td>
<td>1.05</td>
<td>1.03–1.07</td>
<td>&lt;0.0001</td>
</tr>
<tr>
<td>Endoscopic harvest technique (vs. open)</td>
<td>14.07</td>
<td>1.44</td>
<td>1.19–1.75</td>
<td>&lt;0.0001</td>
</tr>
<tr>
<td>Worst target artery quality (vs. good)</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Fair</td>
<td>3.72</td>
<td>1.24</td>
<td>1.00–1.53</td>
<td>0.05</td>
</tr>
<tr>
<td>Poor</td>
<td>8.35</td>
<td>1.45</td>
<td>1.13–1.87</td>
<td>&lt;0.01</td>
</tr>
<tr>
<td>Including 30-day medications</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Duration of surgery (per 10-min increase)</td>
<td>32.51</td>
<td>1.05</td>
<td>1.03–1.07</td>
<td>&lt;0.0001</td>
</tr>
<tr>
<td>Endoscopic harvest technique (vs. open)</td>
<td>12.16</td>
<td>1.41</td>
<td>1.16–1.71</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td>Worst target artery quality (vs. good)</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Fair</td>
<td>3.13</td>
<td>1.22</td>
<td>0.98–1.51</td>
<td>0.08</td>
</tr>
<tr>
<td>Poor</td>
<td>7.55</td>
<td>1.43</td>
<td>1.11–1.84</td>
<td>&lt;0.01</td>
</tr>
<tr>
<td>Clopidogrel or ticlopidine use</td>
<td>6.62</td>
<td>1.35</td>
<td>1.07–1.69</td>
<td>0.01</td>
</tr>
</tbody>
</table>

CI indicates confidence interval; OR, odds ratio.

*1817 patients with non-missing covariates were included in the “without 30-day medications” model, and 1812 patients were included in the “30-day medications” model.

### Table 4. Factors associated with graft-level VGF.

<table>
<thead>
<tr>
<th>Variable</th>
<th>Chi-Square</th>
<th>OR</th>
<th>95% CI</th>
<th>P Value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Without 30-day medications*</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Duration of surgery (per 10-min increase)</td>
<td>27.3</td>
<td>1.04</td>
<td>1.02–1.05</td>
<td>&lt;0.0001</td>
</tr>
<tr>
<td>Endoscopic harvest technique (vs. open)</td>
<td>14.03</td>
<td>1.37</td>
<td>1.16–1.62</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td>Target artery quality (vs. good)</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Fair</td>
<td>9.85</td>
<td>1.31</td>
<td>1.11–1.56</td>
<td>&lt;0.01</td>
</tr>
<tr>
<td>Poor</td>
<td>59.19</td>
<td>2.34</td>
<td>1.89–2.91</td>
<td>&lt;0.0001</td>
</tr>
<tr>
<td>History of cerebrovascular disease</td>
<td>5.82</td>
<td>1.39</td>
<td>1.06–1.81</td>
<td>0.02</td>
</tr>
<tr>
<td>Including 30-day medications</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Duration of surgery (per 10-min increase)</td>
<td>25.30</td>
<td>1.03</td>
<td>1.02–1.05</td>
<td>&lt;0.0001</td>
</tr>
<tr>
<td>Endoscopic harvest technique (vs. open)</td>
<td>12.17</td>
<td>1.35</td>
<td>1.14–1.59</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td>Target artery quality (vs. good)</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Fair</td>
<td>9.35</td>
<td>1.31</td>
<td>1.10–1.55</td>
<td>&lt;0.01</td>
</tr>
<tr>
<td>Poor</td>
<td>58.29</td>
<td>2.34</td>
<td>1.88–2.91</td>
<td>&lt;0.0001</td>
</tr>
<tr>
<td>History of cerebrovascular disease</td>
<td>4.92</td>
<td>1.35</td>
<td>1.04–1.77</td>
<td>0.03</td>
</tr>
<tr>
<td>Clopidogrel or ticlopidine use</td>
<td>7.10</td>
<td>1.30</td>
<td>1.07–1.58</td>
<td>&lt;0.01</td>
</tr>
</tbody>
</table>

CI indicates confidence interval; OR, odds ratio.

*4288 grafts over 1813 patients with non-missing covariates were included in the “without 30-day medications” model, and 4279 grafts over 1808 patients were included in the “30-day medications” model.
Figure Legends:

**Figure 1.** Flowchart of patient selection for the final analysis population.

**Figure 2.** Distribution of predicted VGF risk. Shown is the distribution of predicted risk of VGF using the full (including 30-day medication use) graft-level VGF model among the patient cohort. Listed above each bar is the observed probability of VGF. IQR, interquartile range; VGF, vein graft failure.
All patients enrolled in PREVENT IV (n= 3,014 patients; 107 sites)

Exclude non-angiographic cohort (n=614 patients)

Angiographic cohort (n= 2,400)

Exclude patients who were lost to follow-up and did not return for angiographic follow-up (n= 477 patients)

Exclude patients with only arterial grafts (n= 4)

Exclude patients who died (n=91)

Final analysis population (n= 1,828 patients; 100 sites)
Figure 2

Hosmer-Lemeshow p = 0.85

Predicted Probability of VGF (%) vs. Observed Probability of VGF (%) for different age groups.
Saphenous Vein Graft Failure after Coronary Artery Bypass Surgery: Insights from PREVENT IV

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Knot point selection for non-linear variables encountered during model variable selection

Patient-level model

The following variables were found to be non-linear: aorta cross-clamp time (acctime), pre-operative ejection fraction (ef), and longest graft length (p Glen). Each one was split into 3 linear pieces with the cutoff points chosen appropriately. The definition for each spline follows:

\[\text{acctime}_{low} = \begin{cases} \text{acctime} & \text{if } \text{acctime} \leq 45 \\ 45 & \text{if } \text{acctime} > 45 \\ 45 & \text{if } \text{acctime} < 45 \end{cases}\]
\[\text{acctime}_{mid} = \begin{cases} \text{acctime} & \text{if } 45 \leq \text{acctime} \leq 95 \\ 95 & \text{if } \text{acctime} > 95 \\ 95 & \text{if } \text{acctime} < 95 \end{cases}\]
\[\text{acctime}_{high} = \begin{cases} \text{acctime} & \text{if } \text{acctime} \geq 95 \end{cases}\]
\[\text{ef}_{low} = \begin{cases} \text{ef} & \text{if } \text{ef} \leq 50 \\ 50 & \text{if } \text{ef} > 50 \\ 50 & \text{if } \text{ef} < 50 \end{cases}\]
\[\text{ef}_{mid} = \begin{cases} \text{ef} & \text{if } 50 \leq \text{ef} \leq 60 \\ 60 & \text{if } \text{ef} > 60 \\ 60 & \text{if } \text{ef} < 60 \end{cases}\]
\[\text{ef}_{high} = \begin{cases} \text{ef} & \text{if } \text{ef} \geq 60 \end{cases}\]
\[\text{p Glen}_{low} = \begin{cases} \text{p Glen} & \text{if } \text{p Glen} \leq 10 \\ 10 & \text{if } \text{p Glen} > 10 \\ 10 & \text{if } \text{p Glen} < 10 \end{cases}\]
\[\text{p Glen}_{mid} = \begin{cases} \text{p Glen} & \text{if } 10 \leq \text{p Glen} \leq 20 \\ 20 & \text{if } \text{p Glen} > 20 \end{cases}\]
\[\text{p Glen}_{high} = \begin{cases} \text{p Glen} & \text{if } \text{p Glen} \geq 20 \end{cases}\]

We entered each variable if it was significant (<0.1) in a logistic regression model including only the other levels of the variable. The variables included in our model were acctime_low, acctime_mid, ef_mid, ef_high, and p Glen_mid. Variables were allowed to enter the model separately.
Graft-level model

The following variables were found to be non-linear: aorta cross-clamp time (acctime), pre-operative ejection fraction (ef), and time on bypass pump in minutes (pumptime). Acctime and ef were split into 3 linear pieces with 2 cutoffs, while pumptime was split into 2 linear pieces with 1 cutoff. The definition for each spline follows:

\[
acctime_{low} = \begin{cases} 
acctime & \text{if } acctime \leq 50 \\
50 & \text{if } acctime > 50 \\
50 & \text{if } acctime < 50 
\end{cases}
\]

\[
acctime_{mid} = \begin{cases} 
acctime & \text{if } 50 \leq acctime \leq 90 \\
90 & \text{if } acctime > 90 \\
90 & \text{if } acctime < 90 
\end{cases}
\]

\[
acctime_{high} = \begin{cases} 
acctime & \text{if } acctime \geq 90 
\end{cases}
\]

\[
ef_{low} = \begin{cases} 
ef & \text{if } ef \leq 50 \\
50 & \text{if } ef > 50 \\
50 & \text{if } ef < 50 
\end{cases}
\]

\[
ef_{mid} = \begin{cases} 
ef & \text{if } 50 \leq ef \leq 60 \\
60 & \text{if } ef > 60 
\end{cases}
\]

\[
ef_{high} = \begin{cases} 
ef & \text{if } ef \geq 60 
\end{cases}
\]

\[
pumptime_{low} = \begin{cases} 
pumptime & \text{if } pumptime \leq 80 \\
80 & \text{if } pumptime > 80 
\end{cases}
\]

\[
pumptime_{high} = \begin{cases} 
pumptime & \text{if } pumptime \geq 80 
\end{cases}
\]

We entered each variable if it was significant (<0.1) in a logistic regression model including only the other levels of the variable. Variables included in our model were acctime_low, acctime_mid, ef_low, ef_mid, ef_high, and pumptime_high. Variables were allowed to enter the model separately.
관상동맥우회술 후 이식한 복재정맥의 재협착이나 폐쇄는 흔하며, 환자 및 수술적 요인과 관련이 있다 : PREVENT IV 연구로부터의 건해

나승운 교수 고려대학교 구로병원 순환기내과

Summary

배경
관상동맥우회술(coronary artery bypass grafting, CAGB)의 성공은 복재정맥의 이식 실패(vein graft failure, VGF)에 의해 제한된다. VGF와 관련된 요인들을 이해하면 환자의 임상 결과를 향상시킬 수 있을 것이다.

방법 및 결과
PREVENT IV(Project of Ex Vivo Vein Graft Engineering via Transfection IV) 연구에 참여한 1,828명의 환자를 대상으로 분석을 진행하였다. 이 연구에서는 프로토콜상 의무적으로 CAGB 12-18개월 이후 추적 혈관조영술을 시행하거나, 그 이전에 임상증상이 발생하게 되는 경우 추적 혈관조영술을 시행하게 되어 있었다. 임상 결과는 환자 및 이식관 수준의 VGF(≥75% 혈착 또는 폐색)가 포함되었다. 변수는 빠른 거짓 선택 평가 방법론(Fast False Selection Rate methodology)을 사용하여 선택하였다. 연구자들은 일반화 추정 방정식을 포함 또는 제외하는 로지스틱 회귀 분석을 사용하여 환자 및 이식관 수준의 모델에서 변수들과 VGF 사이의 관계를 조사하였다. CAGB 12-18개월 후 782/1,828명(42.8%)의 환자에서 VGF가 발생하였고, 1,096/4,343(25.2%)의 이식관에서 VGF가 발생하였다. 인구 통계학적 및 임상적 특성 등은 VGF가 발생한 환자나 그렇지 않은 환자에서 차이가 없었다. 다만, VGF 환자들은 외과적 수술 시간이 더 길었고, 목표 혈관의 상태가 좋지 않았던 경우가 많았으며, 이식 복재정맥의 길이가 더욱 짧은 경우가 많았고, 내시경 정맥 수확을 한 경우가 많았다. 단변량 조정 후, 긴 수술 시간(10분 증가 시마다 OR, 1.05; 95% CI, 1.03-1.07), 내 시경 정맥 수확(OR, 1.41; 95% CI, 1.16-1.71), 적절치 않은 목 표 혈관 상태(OR, 1.43; 95% CI, 1.11-1.84)와 clopidogrel 또는 ticlopidine의 수술 후 사용(OR, 1.35; 95% CI, 1.07-1.69)은 환 자 수준 VGF와 관련 있었다. 이식관 수준 VGF 모델의 예측 가능성을 12.1-63.6%였다.

결론
VGF는 CAGB 후 흔히 발생하며, 환자 및 수술적 요인과 관련이 있다. 이러한 연구 결과는 VGF 위험인자들 간의 환자들을 식별하는 데 도움이 되고, VGF를 줄이기 위한 중재적 방법을 개발하는 데 중요한 정보를 제공해 줄 것으로 보인다.