Mid-term outcomes of patients with PCI prior to CABG in comparison to patients with primary CABG

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Abstract: The number of percutaneous coronary interventions (PCI) prior to coronary artery bypass grafting (CABG) increased drastically during the last decade. Patients are referred for CABG with more severe coronary pathology, which may influence postoperative outcome. Outcomes of 200 CABG patients, collected consecutively in an observational study, were compared (mean follow-up: 5 years). Group A (n = 100, mean age 63 years, 20 women) had prior PCI before CABG, and group B (n = 100, mean age 66, 20 women) underwent primary CABG. In group A, the mean number of administered stents was 2. Statistically significant results were obtained for the following preoperative criteria: previous myocardial infarction: 54 vs 34 ($P = 0.007$), distribution of CAD ($P < 0.0001$), unstable angina: 27 vs 5 ($P < 0.0001$). For intraoperative data, the total number of established bypasses was 2.43 ± 1.08 vs 2.08 ± 1.08 ($P = 0.017$), with the number of arterial bypass grafts being: 1.26 ± 0.82 vs 1.07 ± 0.54 ($P = 0.006$). Regarding the postoperative course, significant results could be demonstrated for: adrenaline dosage (0.83 vs 0.41 mg/h; [p is not significant (ns)]) administered in 67 group A vs 47 group B patients ($P = 0.006$), and noradrenaline dosage (0.82 vs 0.87 mg/h; ns) administered in 46 group A vs 63 group B patients ($P = 0.023$), CK/troponine I ($P = 0.002$; $P < 0.001$), postoperative resuscitation (6 vs 0; $P = 0.029$), intra aortic balloon pump 12 vs 1 ($P = 0.003$), and 30-day mortality (9% in group A vs 1% in group B; $P = 0.018$). Clopidogrel was administered in 35% of patients with prior PCI and in 19% of patients without prior PCI ($P = 0.016$). Patients with prior PCI presented for CABG with more severe CAD. Morbidity, mortality and reoperation rate during mid term were significantly higher in patients with prior PCI.

Keywords: CABG, CABG and PCI, CAD, outcome

Introduction

Acceptance of percutaneous coronary intervention (PCI) and use in patients with symptomatic coronary artery disease (CAD) has drastically expanded during the last decade,^{1,2} whereas rates of coronary artery bypass grafting (CABG) have a decreasing tendency.^{2,3}

Randomized trials conducted before and after the implementation of intracoronary stents have demonstrated comparable long-term survival between patients undergoing PCI and patients undergoing CABG for single and/or multivessel CAD.⁴⁻⁶ While PCI was offered only to patients with isolated coronary lesions and single-vessel disease years ago, it is now being used with higher frequency in a broader spectrum of patients. Consequently, a number of referred patients had received PCI, some multiple times, before undergoing CABG.

Results from medical literature comparing PCI and CABG have shown that initial PCI may lead to significantly higher rates of recurrent symptoms and redo operations

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than patients who undergo CABG initially.^{7–16} Although numerous studies have established that reoperative CABG is associated with increased perioperative and long-term mortality and morbidity,^{17–20} knowledge of outcomes in patients who undergo PCI prior to CABG is limited. The objective of this study was to compare the outcomes, including mortality, of group A and group B.

Patients and methods

Patient population

Two hundred patients who underwent CABG between January 2001 and June 2003 at the Department of Cardiac Surgery at the University Hospital of Munich were consecutively observed for our study. Group A included 100 patients (20 women) who had undergone PCI with the implantation of a minimum of one stent prior to CABG, at an average of 321.2 ± 501.8 days before surgery. Among them, in 70 patients, bare metal stents, and in 30 patients, drug-eluting stents were applied. The other 100 patients (group B, 20 women) were referred for primary CABG. Patients with organic valvular disease were excluded. Ethical committee approval was gained from the ethics committee of the University.

Demographic data for all patients, their classical cardiovascular risk factors and comorbidities are summarized in Table 1.

In group A, additional data regarding PCI and related vessels as well as number of stent implantation in total and per vessel were documented (Table 2).

Operative data including operative times, emergency status, number and kind of bypass grafts were recorded in all patients (Table 3). Data on postoperative course are displayed in Table 4. Aspects of intensive care stay, postoperative blood loss, application of blood products and catecholamines were registered. Postoperative outcomes, complications, and mortality after a follow up of 5 years were obtained. Comparison of all data in both groups was established.

Statistical data analysis was carried out with R statistical software (<http://www.r-project.org>). Differences between the two groups were statistically assessed using Fisher's exact test (for categorical variables) and the 2-sided Kolmogorov–Smirnov test (for continuous variables). The Kolmogorov–Smirnov test was applied because many variables did not fulfil, even approximately, the requirements of the *t*-test and the Mann–Whitney test. Multivariate analyses

Table 1 Patient's demographics, cardiovascular risk factors and comorbidities

Cardiovascular Risk Factors/Comorbidities	Group A (n = 100) [Number of Patients, Percentage]	Group B (n = 100) [Number of Patients, Percentage]	P-value
Male/Female sex	80 (80%)/20 (20%)	80 (80%)/20 (20%)	ns
Age [Years]	63 ± 10,5	65,9 ± 9,9	ns
Arterial hypertension	88 (88%)	83 (83%)	ns
Diabetes mellitus	31 (31%)	26 (26%)	ns
Hyperlipidemia	85 (85%)	84 (84%)	ns
Peripheral arterial occlusive disease	23 (23 %)	14 (14%)	ns
Body mass index [Mean in kg/m ²]	26,38 ± 3,87	27,2 ± 3,9	ns
Dialysis	6 (6%)	3 (3%)	ns
Transitory ischemic attack	4 (4%)	3 (3%)	ns
Distribution of NYHA Class	0: 3% I: 11% II: 40% III: 36% IV: 10%	0: 0% I: 3% II: 51% III: 41% IV: 5%	0,025
Mean NYHA Class	2,39 ± 0,92	2,48 ± 0,64	
Previous myocardial infarction	54 (54%)	34 (34%)	0,007
CAD	1: 11% 2: 23% 3: 67%	1: 27% 2: 40% 3: 33%	<0,0001
Number and percentage by vessel	Left Main Stem: 3%	0	ns
Left ventricular ejection fraction [Mean and SD]	60,04 ± 14,5	60,47 ± 14,1	ns
Unstable angina pectoris	27 (27%)	5 (5%)	<0,0001
LV aneurysm	5 (5%)	0	0,059

Table 2 Group A PCI details prior to CABG

PCI	1: 48%
(number of interventions as percentage)	2: 28%
	3: 14%
	4: 7%
	5: 3%
mean and SD	1.89 ± 1.08
Left main stem stenosis	3 (3%)
Target artery in n (%)	
LAD	56 (56%)
RCX	33 (33%)
RCA	47 (47%)
Multivessel PCI in n (%)	
RCX + RCA	10 (10%)
LAD + RCA	11 (11%)
LAD + RCX	4 (4%)
LAD + RCX + RCA	6 (6%)
Interval between PCI and CABG	321.2 ± 501.8
(mean time in days)	
Stents (mean number of implantations)	2.14 ± 1.32

Abbreviations: PCI, percutaneous coronary intervention; CABG, coronary artery bypass grafting; LAD, left anterior descending artery; RCA, right coronary artery; RCX, right circumflex artery; SD, standard deviation.

were performed using linear or logistic regression models, depending on the type of outcome. Statistical significance within multivariate models was assessed based on the Wald test. Overall survival was represented using Kaplan–Meier curves and the logrank test was used to compare the two groups. *P*-values <0.1 were marked with “ns” in Tables 1–4.

Table 3 Intraoperative data

Parameter	Group A	Group B	P-value
Intraaortic balloon pump	3 (3%)	0	ns
Intubation before surgery	2 (2%)	0	ns
Emergency	3 (3%)	0	ns
Time of operation	269.6 ± 102.5	252.7 ± 78.7	ns
(minutes, mean ± SD)			
Cardiopulmonary bypass	130.3 ± 66.8	109.3 ± 40.1	ns
time (minutes, mean ± SD)			
Aortic cross clamp time	72.5 ± 30	65.6 ± 24.4	ns
(minutes, mean ± SD)			
Time of reperfusion	40.8 ± 26.5	36.3 ± 19.8	ns
(minutes, mean ± SD)			
Lowest temperature (°C)	30.2 ± 3.3	30.9 ± 2.3	ns
Amount of Cell Saver®	616.1 ± 378	568.7 ± 333.1	ns
blood (mL, mean ± SD)			
Number of established	2.43 ± 1.08	2.08 ± 1.08	0.017
bypasses (mean ± SD)			
Number of arterial	1.26 ± 0.82	1.07 ± 0.54	0.006
bypass grafts (mean ± SD)			
Number of venous	1.17 ± 1.03	1.02 ± 0.94	ns
bypass grafts (mean ± SD)			
LITA to LAD	80 (80%)	89 (89%)	ns
Total RCA	21 (21%)	15 (15%)	ns

Abbreviations: LITA, left internal thoracic artery; LAD, left anterior descending artery; RCA, right coronary artery; SD, standard deviation; ns, not significant.

P-values <0.05 were considered significant, whereas *P*-values between 0.05 and 0.1 were considered as trends.

Results

Demographic data, classic cardiovascular risk profile and comorbidities are listed in Table 1. Therapy with aspirin, GPIIb/IIIa inhibitors, ACE inhibitors, AT-I receptor blockers, calcium channel blockers, diuretics, statins, and nitrates showed an almost equal distribution among groups. Clopidogrel was administered in 35% of the patients with prior PCI and in 19% of patients without prior PCI (*P* = 0.016).

Table 2 includes all data regarding PCI in patients of group A. Almost half of this group underwent one PCI (48%). LAD was stented in 56 patients (56%), followed by RCA in 47 patients (47%) and circumflex artery in 33%, either singly or in combination. Only 6 patients received stents in three coronary arteries. Mean duration between PCI and CABG was 321 days with a standard deviation of 502 days. This number may already show the big variance of evidence of clinical CAD symptoms.

Summarized intraoperative data for both groups are compared and listed in Table 3. All patients underwent CABG on cardiopulmonary bypass under cardioplegic hypothermic arrest. Time of extracorporeal circulation (130 vs 109 minutes; ns) revealed statistically nonsignificant differences between both groups.

The amount of blood saved by Cell Saver® was slightly lower in group B (616 ± 378 mL in group A vs 568 ± 333 mL in group B, respectively; ns). The mean number of established bypasses was 2.43 in group A vs 2.08 in group B (Fisher's exact test: *P* = 0.017). Establishment of arterial bypass grafts showed the same trend: 1.26 vs 1.07 (Fisher's exact test: *P* = 0.006). Left anterior descending artery was revascularized by the left anterior thoracic artery (LITA) in 80% of group A patients and 89% of group B patients (ns).

Most important for outcome evaluation are postoperative parameters. Ventilation time and stay on intensive care unit did not demonstrate significant differences. Contrary to our expectation, the registered amount of bleeding 48 hours postoperatively was slightly higher in group B (1345 ± 1113 mL vs 1055 ± 842 mL in group A; ns). A mean of 2.3 units of thrombocyte concentrate was used in 34 patients of group A and 1.6 units in 22 group B patients (ns). The amount of bleeding in both groups may be related to thrombocyte dosage, especially under the circumstance of being given preoperative antiplatelet medication. Neither the amount of erythrocyte concentrate nor the amount of fresh frozen plasmas was significantly

Table 4 Postoperative data

Parameter	Group A	Group B	P-value
Postoperative bleeding (mL, first 48 hours, mean $\pm$ SD)	1055.10 $\pm$ 842.5	1345.54 $\pm$ 1113.8	ns
Ventilation time (hours, mean $\pm$ SD)	34.86 $\pm$ 58.03	31.64 $\pm$ 91.95	ns
Stay on intensive care unit (hours, mean $\pm$ SD)	70.46 $\pm$ 71.59	76.07 $\pm$ 111.75	ns
Use of thrombocyte concentrate (mean $\pm$ SD)	Total: 34 patients 2.29 $\pm$ 1.45	Total: 22 patients 1.64 $\pm$ 0.85	0.083 ns
Use of erythrocyte concentrate (mean $\pm$ SD)	Total: 71 patients 4.00 $\pm$ 4.57	Total: 58 patients 3.62 $\pm$ 2.38	0.076 ns
Use of fresh frozen plasma (mean $\pm$ SD)	Total: 39 patients 6.92 $\pm$ 7.80	Total: 33 patients 5.09 $\pm$ 3.14	ns ns
Adrenaline dosage (mg/h, mean $\pm$ SD)	Total: 67 patients 0.83 $\pm$ 1.63	Total: 47 patients 0.41 $\pm$ 0.26	0.006 ns
Noradrenaline dosage (mg/h, mean $\pm$ SD)	Total: 46 patients 0.82 $\pm$ 1.24	Total: 63 patients 0.87 $\pm$ 2.11	0.023 ns
Hydrocortisone use (number of patients)	28	36	ns
Creatinine, highest level (mg/dL, mean $\pm$ SD)	1.95 $\pm$ 1.6	1.61 $\pm$ 1.4	0.054
Creatine kinase, highest level (U/L, mean $\pm$ SD)	876.52 $\pm$ 1815.83	414.29 $\pm$ 461.14	0.002
CK-MB, highest level (U/L, mean $\pm$ SD)	40.1 $\pm$ 53.14	24.8 $\pm$ 33.89	0.059
Troponin I, highest level (ng/mL, mean $\pm$ SD)	132.89 $\pm$ 206.10	106.54 $\pm$ 616.88	<0.001
Postoperative resuscitation (number of patients, percentage)	6 (6%)	0	0.029
Postoperative myocardial infarction	4 (4%)	0	ns
Neurological disorders	8 (8%)	4 (4%)	ns
Postoperative atrial fibrillation	25 (25%)	35 (35%)	ns
Postoperative acute renal failure	9 (9%)	3 (3%)	ns
Wound infection	2 (2%)	4 (4%)	ns
Intraaortic balloon pump	12 (12%)	1 (1%)	0.003
Rethoracotomy	8 (8%)	9 (9%)	ns
In-hospital stay (days; mean $\pm$ SD)	11.1 $\pm$ 10	12.4 $\pm$ 5.7	ns
30-day mortality	9 (9%)	1 (1%)	0.018

Abbreviations: SD, standard deviation; ns, not significant.

different between groups. The dosage of adrenaline was not significantly different in the two groups (0.83 vs 0.41 mg/h; ns). The difference in the mean was due to a small number of extreme values. However, the number of patients who received adrenaline was higher in group A than in group B (67 vs 47, respectively; $P = 0.006$). Mean noradrenaline dosage was approximately equal in the two groups (0.82 vs 0.87 mg/h; ns), and noradrenaline was administered in 46 group A vs 63 group B patients ($P = 0.023$). Six patients of group A were resuscitated postoperatively whereas no group B patient had to be resuscitated ($P = 0.003$). Myocardial markers CK and troponin I did show statistically significant differences between groups ($P < 0.002$; $P < 0.001$) with strongly higher levels in group A. Twelve group A patients needed

an intraaortic balloon pump implantation postoperatively, whereas only one was implanted in patients of group B ($P = 0.003$).

30-day mortality was 9% in group A and 1% in group B ($P = 0.018$). This difference may be caused by several factors that will be discussed below. The five-year mortality rate was 14% in group A vs 9% in group B (logrank-test: ns, Figure 1). Five group A patients necessitated redo-CABG vs none of group B (ns).

In multivariate regression analyses with age, NYHA class, ejection fraction, and CAD as adjustment factors, prior PCI was not a significant predictor of postoperative bleeding, ventilation time, stay on intensive care unit, use of thrombocyte concentrate, use of erythrocyte concentrate, or fresh frozen plasma.

Figure 1 Probability of survival among study groups.

Discussion

Trials comparing PCI and CABG have demonstrated that long-term survival of PCI and CABG is similar, but rates of symptom recurrence and repeated revascularization are significantly higher in patients with initial PCI.^{10,12,16} With dramatic increase in PCI rates during the last decade,^{1,2} it is a consequence that the number of patients referred for CABG with a prior PCI has increased.²¹

Kalaycioglu et al²² compared 40 patients who had undergone CABG with a prior PCI with a case-matched control group of 40 patients who underwent CABG with no prior PCI. In contrast to our study, the authors found that patients with a prior PCI were younger, had a higher preoperative ejection fraction, less extensive coronary disease, and had fewer bypass grafts. After finishing this study, we agree with their results of diminished freedom from death, angina, and repeat revascularization procedure in the 36 months after surgery in patients with a prior PCI. In another study, Barakate et al²³ compared 361 patients who underwent CABG after initially

successful PCI. Similar to the results from Hassan et al²⁴ they demonstrated that patients with a prior PCI had a higher pre-operative ejection fraction, less extensive coronary disease, and fewer bypass grafts, and were more likely to present with worsened symptoms. However, contrary to the findings from Hassan et al Barakate et al showed similar rates of in-hospital mortality between the 2 groups.

After analysis of our results, we found a stronger disease severity of CAD in group A ($P < 0.0001$). NYHA class distribution before surgery was significantly different among both groups ($P = 0.025$), 27 group A and 5 group B patients presented with unstable angina ($P < 0.0001$). Percentage of previous myocardial infarction differed significantly between both groups: 54% of group A and 34% of group B patients ($P = 0.007$). Clopidogrel was preoperatively administered in 35% of the patients with prior PCI and in 19% of patients without prior PCI ($P = 0.016$).

Regarding intraoperative data, statistically significant results were demonstrated for group A and group B: the

mean number of established bypasses was significantly different among groups (2.43 vs 2.08, $P = 0.017$). The mean number of arterial bypass grafts used for revascularisation showed significant results as well (1.26 vs 1.07, $P = 0.006$). Left anterior descending artery was revascularized by LITA in 80% of group A patients and 89% of group B patients (ns). In search of causes for this diverse percentage of LITA graft use, we conjecture that severity of LAD calcification and placement of intracoronary stents, as well as emergency status of surgery, may be primary reasons for using this procedure.

During the postoperative course, statistically significant results could be registered for: frequency of adrenaline ($P = 0.006$) and noradrenaline ($P = 0.023$) administration, CK ($P = 0.002$) and troponine I ($P < 0.001$), and postoperative resuscitation (6 vs 0; $P = 0.029$). Even during perioperative duration, data leads to the supposition that the severity of disease seems to be higher and further advanced in group A. Contrary to our supposition, the amount of postoperative drainage loss was higher in group B (1345 ml) than in group A (1055 ml; ns). We expected, especially after clinical intraoperative impression, a higher bleeding rate in group A, mostly due to antiplatelet therapy. Blood product utilization, especially of thrombocyte concentrates, was higher in group A (2.3 vs 1.6 units; ns) and may be one of the reasons for limited blood loss after surgery. Rethoracotomy rates were almost equal in both groups.

Twelve patients from group A received postoperatively an intraaortic balloon pump. Only one patient of group B necessitated an intraaortic balloon pump after surgical procedure ($P = 0.003$).

30-day mortality was significantly lower in group B (9% vs 1%; $P = 0.018$). Five-year mortality was decreased in group B (14% in group A vs 9% in group B; ns). 83 patients of group A and 89 patients of group B showed NYHA class one or two after five years (ns). 16 group A patients underwent re-PCI and stent implantation vs 13 of group B (ns). 5 patients of group A and no patient of group B underwent redo-CABG (ns).

The reasons for a higher mortality in short and mid term after CABG in patients with prior PCI are not clearly understood. In our cohort, these patients had more comorbidity and increased coronary disease pathology, and presented for surgery with advanced symptoms and greater urgency. It might be that patients who had initial PCI had a more aggressive atherosclerotic onset promoting a higher rate of restenosis and/or the development of new lesions at remote coronary sites.²⁵ In addition, the

application of successful PCI in these patients may have prevented the formation of protective collateral vessels, which resulted in the more acute presentation of patients who required repeated revascularization. Finally, prior PCI and, especially, stenting of a coronary lesion may have resulted in an anastomosis at a more distal site with a smaller luminal diameter and suboptimal runoff, thus compromising the degree to which the patient was completely revascularized.

Study limitations

This study has several limitations. It has been designed as a consecutive, observational, single center investigation. The number of enrolled patients limits the explanatory power of our study. Selection of patients of both groups may introduce an underlying bias.

We could not elaborate on the factors influencing the physicians' and patients' decisions to proceed with PCI or surgery first, noting that cardiothoracic surgeons do see patients usually after cardiologists have visited them. Even as we limited our analysis to short-term and mid-term outcomes, it is plausible that different methods of follow-up for patients treated at or outside our institution could have led to some misclassification bias. The number of enrolled patients limits the explanatory power of our investigation.

Disclosure

The authors report no conflicts of interest in this work.

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