| Original Research Article |
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| Pedicled and skeletonized single and bilateral internal mammary artery grafts and the incidence of sternal wound complications: Insights from the Arterial Revascularization Trial (ART) |
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39 Abbreviation

- 40 ASMD: absolute standardised mean difference
- 41 ART: Arterial Revascularization trial.
- 42 ATE: average treatment effect on the population
- 43 BIMA bilateral internal mammary artery
- 44 CABG: coronary artery bypass grafting
- 45 ESS: Effective sample size
- 46 IMA: internal mammary artery
- 47 SIMA: single internal mammary artery
- 48 S: skeletonized
- 49 P: pedicled

50 Structured Abstract

51 **Objective(s):** The question of whether skeletonized internal mammary artery (IMA) 52 harvesting reduces the incidence of sternal wound complications in comparison to the 53 pedicled technique, in the context of single or bilateral IMAs, remains controversial. 54 We studied the impact IMA harvesting strategy on sternal wound complication in the 55 Arterial Revascularization trial (ART).

Methods: Patients enrolled in the ART (n=3103) were randomised to coronary artery 56 bypass grafting with single or bilateral IMAs. Sternal wound complication rates were 57 examined according to the harvesting technique that was documented in 2056 58 patients. The IMA harvesting technique, based on surgeon preference, resulted in 4 59 groups: pedicled single IMA (P-SIMA, n=607), pedicled bilateral IMA (P-BIMA, n=459), 60 skeletonized single IMA (S-SIMA, n=512) and skeletonized bilateral IMA (S-BIMA, 61 n=478). Propensity Scores weighting was used to estimate the impact of the 62 63 harvesting technique on sternal wound complications.

Results: A total of 219 of 2056 patients (10.6%) experienced a sternal wound complication within 1 year from the index operation. Of those, only 25 (1.2%) patients required sternal wound reconstruction. P-BIMA (OR 1.80; 95%Cl 1.23 to 2.63) but not S-BIMA (OR 1.00; 95%Cl 0.65 to 1.53) or S-SIMA (OR 0.89; 95%Cl 0.57-1.38) was associated with a significantly increased risk of any sternal wound complications compared to P-SIMA.

Conclusions: The present ART sub-study suggests that, with a skeletonization
 technique, the risk of sternal wound complication with BIMA grafting is at a
 similar level to that after standard pedicled SIMA harvesting whilst skeletonized
 SIMA harvesting did not add any further benefit when compared to pedicled
 SIMA harvesting.

Central picture: Incidence of sternal wound complications according to internal
mammary artery harvesting strategies (P-SIMA: pedicled single internal mammary
artery; P-BIMA: pedicled bilateral internal mammary arteries; S-SIMA: skeletonized
SIMA; S-BIMA: skeletonized bilateral internal mammary arteries)

79 Central message

In the Arterial Revascularization trial, the risk of sternal wound complication with
bilateral internal mammary arteries was comparable to that after single pedicled
harvesting.

83 **Perspective Statement**

- By using skeletonized harvesting technique, the risk of sternal wound complication with bilateral internal mammary artery (IMA) grafting is at a similar level to that after standard pedicled single IMA harvesting also in patients at higher risk such as insulin dependent diabetes, females and those with increased body mass index.
- 88 Bilateral IMAs should not be denied on basis of increased risk of sternal wound
- 89 complication if sketetonized harvesting technique is used.

The long term patency of conduits is one of the most important determinants of longterm outcomes in coronary artery bypass grafting (CABG). The left internal mammary artery (IMA) is unanimously acknowledged as the best coronary conduit [1]. Although the right IMA has identical function and patency rates to the left IMA and despite accumulation of evidence on long term benefit by using bilateral IMAs (BIMA) over the past 20 years [2-4], the right IMA remains largely underutilized [5] mainly due to concerns over the potential for sternal wound complications [6].

There are two established techniques for harvesting the IMA: pedicled and 97 skeletonized. Harvesting the IMA(s) in a pedicled fashion can potentially lead to 98 significant sternal devascularisation [7,8]. As opposed to pedicled harvesting, 99 minimization of tissue mobilization during skeletonized IMA harvesting has been 100 101 shown to preserve substantial collateral flow to the sternum by sparing some of the sternal and intercostal branches that arise from the internal mammary artery as a 102 common trunk [7,8]. This finding may have potential clinical significance with respect 103 to reducing the risk of sternal wound complications by improving wound healing and, 104 in particular, when both left and right IMAs are used [9]. 105

However, the magnitude of the potential clinical benefit from skeletonized over pedicled IMA harvesting on sternal wound complications still remains to be determined [10,11]. Moreover, skeletonized IMA harvesting is a more technically demanding and time consuming technique and concerns remain over a perceived increased risk of injury to the IMAs during skeletonization that may affect early outcomes [12]. Consequently, in the absence of a general consensus, pedicled IMA harvesting, remains the generally preferred approach worldwide.

The Arterial Revascularization Trial (ART) is a randomized comparison of bilateral IMA (BIMA) versus single IMA (SIMA) grafting in CABG surgery [13] and is also one of the largest studies of contemporary CABG with a high proportion of patients undergoing skeletonized IMA harvesting. We studied the impact of IMA harvesting strategy on sternal wound complication by conducting an analysis of data collected prospectively in the Arterial Revascularization trial (ART).

119 Methods

120 This research adheres to the principles set forth in the Declaration of Helsinki (http://www.wma.net/en/30publications/10policies/b3/index.html). The ART has been 121 approved by the institutional review board of all participating centers and informed 122 123 consent was obtained from each participant. The protocol for ART has been published 124 [14] Briefly, ART is a two-arm, randomized multicentre trial, conducted in 28 hospitals in seven countries, with patients being randomized equally to SIMA or BIMA grafts. 125 Eligible patients were those with multivessel coronary artery disease (including urgent 126 patients but not evolving myocardial infarction) undergoing CABG, whereas those 127 requiring single grafts or redo CABG were excluded. Only surgeons with experience 128 of ≥50 BIMA operations were able to participate in the trial; standard methods for 129 anaesthesia and myocardial protection were used according to local practice. For the 130 purpose of the present analysis, patients were classified according to the "as 131 treated" principle in the following groups: pedicled single IMA (P-SIMA), 132 skeletonized single IMA (S-SIMA), pedicled bilateral IMA (P-BIMA) and 133 skeletonized BIMA (S-BIMA). IMA harvesting technique was based on surgeon 134 preference. This information was not recorded from the outset of the trial. Thus only 135 2056 out of 3102 patients were included in the analyses; among those 1022 and 136 1034 were initially allocated to BIMA and SIMA respectively. Crossover rate from 137

BIMA to SIMA was 115/1022(11.2%) and from SIMA to BIMA was 30/1034(2.9%). 138 Finally a total of 937 and 1119 patients received BIMA and SIMA respectively.

Outcomes definition 140

139

The primary end-point for these analyses was the incidence of any sternal wound 141 complication within 1 year after the index procedure, which included a broad definition 142 143 ranging from superficial sternal wound discharge to sternal wound reconstruction. We also investigated the impact of IMA harvesting strategy on the incidence of severe 144 sternal wound complications, defined as sternal wound infection requiring antibiotics 145 and/or sternal wound reconstruction. Adverse events including sternal wound 146 complications were adjudicated blind by a member of the Clinical Event Review 147 Committee. 148

149 Statistical analysis

For baseline characteristics, variables are summarised as mean for continuous 150 variables and percentage for categorical variables. The chi squared test was used to 151 test unadjusted association between treatment variable and outcomes. Multiple 152 imputation (m=3) was used to address missing data (165 patients). Rubin's method 153 [15] was used to combine results from each of *m* imputed data sets. 154

Inverse probability of treatment weighting for modelling causal effects was 155 used to for multiple treatments comparison [16]. One of the advantages of this 156 technique over standard pairwise propensity matching is the possibility of 157 simultaneous comparisons between multiple treatments. Moreover, all the 158 individuals in the study can be used for the outcomes evaluation whilst a large 159 number of subjects may not be used in a propensity matching in particular 160 when the sample size of treatment and control groups are similar. A 161 generalised boosted model was implemented to estimate multinomial propensity 162

scores (PS) adjusting for 14 pre-treatment covariates, and the propensity score was 163 assumed as the probability that an individual with pre-treatment characteristics X 164 receives treatment t (twang R package). The average treatment effect on the 165 population (ATE) was used to answer the question of how, on average, the outcome 166 of interest would change if everyone in the population of interest had been assigned 167 to a particular treatment relative to if they had all received another single treatment. 168 To estimate the ATE, we gave treated patients weight $w_i = 1/(1 - p(x_i))$, where $p(x_i)$ is 169 the propensity score, and reference patients $w_i = 1/p(x_i)$. P-SIMA was considered as 170 171 the reference group in all comparisons. The absolute standardised mean difference (ASMD) was used as a balance metric to summarize the difference between two 172 univariate distributions of a single pre-treatment variable. A value ≥0.20 (20%) was 173 considered as an indicator of imbalance [17]. Effective sample size (ESS) was 174 calculated to account for the potential loss in precision from weighting [16]. We then 175 estimated the treatment effect estimates with a weighted regression model that 176 contained only a treatment indicator. In addition, a combination of propensity score 177 weighting and covariate adjustment (double robust) was used to correct the effect of 178 IMA harvesting technique for residual imbalance and to estimate the effect size of 179 other covariates. Lastly, we estimated the treatment effect within subgroups 180 according to the presence of diabetes on insulin, gender and body mass index \geq 30. 181 R version 3.1.2 (2014-10-31) was used for all statistical analysis. 182

183 **Results**

184 Study population

Among 2056 patients included in the present analysis, 1022 and 1034 were initially allocated to BIMA and SIMA respectively. Crossover rate from BIMA to

SIMA was 115/1022(11.2%) and from SIMA to BIMA was 30/1034(2.9%). Finally a 187 total of 937 and 1119 patients received BIMA and SIMA respectively. IMA 188 harvesting groups compared were: 607 P-SIMA, 459 P-BIMA, 512 S-SIMA and 189 190 478 S-BIMA. The second IMA was initially attempted to be harvested but not used in 15 BIMA to SIMA crossovers. Of those, 5 were skeletonized and 10 were 191 pedicled. Reasons for the second IMA not to be used were: evidence of injury 192 during harvesting (n=4, all pedicled), unsatisfactory flow (n=5, 3 skeletonized, 2 193 pedicled) or unsatisfactory length or size (n=6, 2 skeletonized, 4 pedicled). 194 195 Overall, rate of injured/unsatisfactory second IMA was 5/483(1.0%) by using skeletonized technique and 10(2.1%) by using pedicled technique (P=0.22). 196 Among those 15 cases, only 1 patient who received pedicled harvesting, 197 experienced sternal wound complication. 198

Distribution of pre-treatment variables among IMA harvesting technique groups

Table 1 summarises the distribution of pre-treatment variables. Although the four groups were comparable for most of the pre-treatment variables, insulin dependent diabetes was more common in patients receiving S-BIMA than in patients receiving P-BIMA. In addition more women received either skeletonized SIMA or BIMA. Finally offpump surgery was more frequently performed in S-SIMA and S-BIMA groups compared to pedicled groups.

After multinomial propensity score estimation balance check showed that the groups were sufficiently similar (ASMDs <0.20) to support causal estimation of the treatment effects, although subjects receiving P-BIMA continued to have a slightly lower prevalence of diabetes on insulin.

210 Incidence of sternal wound complications

A total of 219 out of 2056 patients (10.7%) experienced a sternal wound complication within 1 year from the index operation. Of those, 75 (3.6%) patients had severe sternal wound complications including 50 (2.4%) with sternal wound infection requiring antibiotic therapy but not reconstruction and 25 (1.2%) who needed sternal wound reconstruction. Most sternal wound complications including those requiring reconstruction occurred during the first three months (Figure 1).

217 Effect of harvesting technique on sternal wound complication

218 Table 2 and Figure 2 show the incidence of any sternal wound complications according to IMA harvesting groups. P-BIMA patients had a higher incidence of any sternal 219 wound complication compared to the other groups. There were too few severe wound 220 221 complications to detect differences among the treatment groups. Table 3 summarises 222 the effect of IMA harvesting technique on the incidence of any sternal wound complications. PS weighted analysis showed that P-BIMA but not S-BIMA was 223 associated with a significantly increased risk (~ 2 times) of any sternal wound 224 complications when compared to P-SIMA. On the other hand, S-SIMA did not provide 225 any benefit on the incidence of any sternal wound complication when compared to P-226 227 SIMA. When the analysis was restricted to severe sternal wound complications only we were unable to demonstrate any significant impact of P-BIMA (OR 1.60; 95%CI 228 0.85-3.00), S-BIMA (OR 1.15;95%CI 0.58-2.28) and S-SIMA (OR 0.97; 95%CI 0.45-229 2.07) when compared to P-SIMA. 230

231 Subgroup analysis

Subgroup analysis (Table 3) suggested that the detrimental effect of P-BIMA on the incidence of any sternal wound complication might be exaggerated in the presence of diabetes on insulin **(OR 4.05; 95%CI 0.86-19.21)** although this analysis was largely

underpowered due to the very small number of patients on insulin (n=118). Of note,
P-BIMA remained significantly associated with a higher risk of any sternal wound
complication in patients not diabetic (OR 1.84; 95%Cl 1.18-2.85). Moreover P-BIMA
significantly increased the risk of any sternal wound complication in both obese and
non-obese patients.

In the situation of a single IMA, skeletonized SIMA did not add any significant benefit
in terms of sternal wound complication when compared to P-SIMA also among high
risk subgroups.

243 Independent risk factors for sternal wound complication

In a double robust analysis (Table 4 and Table 5), P-BIMA but not S-BIMA remained
independently associated with an increased risk of any sternal wound complication.
Insulin dependent diabetes, female gender, and higher BMI were independent risk
factors for any and severe sternal wound complications.

248 Mortality within 30 days and at 1 year

There were 31 (1.5%) deaths within 30 days and 55 (2.6%) deaths by 1 year followup. Mortality at 30 day and 1 year was comparable among IMA harvesting groups (Table 2). 30 day mortality among patients with and without sternal wound reconstruction was 0/25(0%) and 31/2031(1.5%). At 1 year, total deaths among patients with and without sternal wound reconstruction were 3/25(12%) and 52/2031 (2.7%).

255 Discussion

Despite increasing evidence from observational studies of the long term survival benefit of a second IMA [2,3], it remains largely underutilised being used in 4.1% of CABG in the USA [5], and around 10% in the UK and Australia [18]. Concern about

sternal wound complication is one of the main reasons limiting the use of more than
one IMA, as a severe sternal wound complication dramatically increases in-hospital
mortality as well as the expense of hospital stay [6].

The present post hoc analysis of the ART demonstrates that in the modern era of CABG surgery sternal wound complications still affect about 10% of patients. In particular, severe sternal wound infection requiring antibiotic therapy or sternal wound reconstruction still affects nearly 2% and 1% of the surgical population respectively. The anticipated impact of sternal wound complication on resource consumption and patient outcomes represents an important consideration in the utilisation of BIMA grafting and an argument in favour of skeletonized IMA over pedicled IMA harvesting.

269 The main finding of the present analysis is that BIMA harvesting can be safely 270 performed using the skeletonized technique without increasing the risk of sternal wound complications when compared to the standard approach using a pedicled 271 SIMA. Furthermore, skeletonized BIMA harvesting does not seem to significantly 272 increase the risk even in higher risk groups, such as diabetics on insulin, females and 273 the obese (BMI≥30). On the other hand, pedicled BIMA was associated with a nearly 274 2 fold increased risk of any sternal wound complication. The detrimental effect of 275 pedicled BIMA harvesting on sternal wound complication was relevant not only in high 276 risk cases such as those who were obese or who had insulin dependent diabetes but 277 also in the lowest risk CABG population who were not diabetic or obese, whilst 278 skeletonized BIMA harvesting did not significantly increase the risk of sternal wound 279 complications. 280

281 On the other hand, in the context of a single IMA graft, there was no evidence of the 282 superiority of skeletonized SIMA harvesting over pedicled SIMA harvesting in reducing 283 the risk of sternal wound complications.

284 Skeletonized harvesting has been proposed to minimise the risk of sternal wound 285 complication by preserving sternal perfusion especially in the context of BIMA usage 286 [6]. Kamiya et al. [7] showed better oxygen saturation and blood flow in the 287 microcirculation of sternal tissue when using skeletonized rather than pedicled IMA. 288 Similarly, Boodhwani et al. [8], using radionuclear perfusion scanning, demonstrated 289 that sternal perfusion was greater after skeletonized rather than pedicled harvesting.

However, whether skeletonized IMA harvesting should be considered the standard 290 291 approach with BIMA grafting and whether this approach also provides a significant 292 benefit in SIMA grafting still needs to be determined. The potential clinical superiority of skeletonized over pedicled harvesting on sternal wound complications has been 293 addressed only in a few studies with conflicting results reported [10-11]. Studies 294 published to date are remarkably underpowered to detect any clinical benefit on low 295 rate events such as sternal wound complications [11]. Moreover, skeletonized 296 harvesting is more technically demanding and time consuming and, in the absence of 297 general consensus, pedicled harvesting still remains the preferred approach 298 worldwide. 299

ART is one of the largest studies of contemporary CABG with a high proportion of patients undergoing skeletonized IMA harvesting [13]. To our knowledge, the present study is the largest analysis on the impact of IMA harvesting performed to date. We found that skeletonization while performing BIMA was safe as did not increase the risk of damage to harvested IMA. In fact, rate of injured/unsatisfactory

305 second IMA was 1.0% by using skeletonized technique and 2.1% by using pedicled technique thus supporting previous reports [22]. Moreover, mortality 306 rate at 30 days and 1 year was comparable among the two techniques. With 307 308 regard to sternal wound complications, skeletonized BIMA harvesting did not increase its risk when compared to pedicled SIMA and subgroup analysis suggested 309 a protective effect from skeletonized BIMA also among high risk subjects. On the other 310 hand, pedicled BIMA grafts seemed to increase the risk of sternal wound 311 complications also among low risk subgroups (ie not on insulin nor obese). We also 312 313 found no evidence that skeletonized SIMA harvesting added any protective effect when compared to a pedicled SIMA. 314

315 Limitations

316 The present analysis has some limitations. Despite propensity score adjustment, the present analysis was unable to address hidden biases due to unobserved differences 317 between treated and control patients before treatment. The present study was 318 underpowered to detect differences in severe sternal wound complications among 319 groups and most of sternal complications were clinically less relevant. 320 321 Fortunately, the low incidence of severe sternal wound complications would have required a much larger number of patients for analysis. Nevertheless, the difference 322 in the rate of severe wound problems between the 2 groups supports the intrinsic 323 benefit of the skeletonized technique of artery harvesting in terms of severe sternal 324 wound complications. Sparing of the communicating bifurcation of internal 325 mammary artery to the chest wall and preservation of pericardiacophrenic artery 326 branch has been reported to minimize the risk of sternal wound complication in 327 patients receiving pedicled BIMA [23]. In the present study we could not confirm 328

this hypothesis as data on technical aspects of harvesting technique were not
 reported.

331 Conclusion

In conclusion, the present ART sub-study suggests that, with a skeletonization technique, the risk of sternal wound complication with BIMA grafting is at a similar level to that after standard pedicled SIMA harvesting whilst skeletonized SIMA harvesting did not add any further benefit when compared to pedicled SIMA harvesting. Skeletonized BIMA harvesting seems to provide a protective effect also in those at higher risk such as insulin dependent diabetes, females and those with increased BMI.

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| | P-SIMA n=607 | S-SIMA n=512 | P-BIMA n=459 | S-BIMA n=478 | ASMD | Р | P-SIMA ESS=550 | S-SIMA ESS =454 | P-BIMA ESS =429 | S-BIMA ESS =430 | ASMD | Р |
|---------------------------|-----------------|-----------------|-----------------|-----------------|------|--------|--------------------------|---------------------------|--------------------|--------------------|------|------|
| | | l | Inweighted | | | | Weighted | | | - | | |
| Age, year (SD=9) | 64 | 65 | 63 | 64 | 0.23 | <0.001 | 64 | 64 | 64 | 64 | 0.06 | 0.36 |
| Female | 11% | 19% | 12% | 14% | 0.22 | <0.001 | 12% | 14% | 13% | 12% | 0.05 | 0.36 |
| BMI (SD=4) | 28.29 | 28.17 | 28.30 | 28.38 | 0.05 | 0.44 | 28.24 | 28.28 | 28.31 | 28.24 | 0.02 | 0.77 |
| Creatinine,mmol/I (SD=22) | 97.91 | 100.00 | 98.23 | 98.30 | 0.09 | 0.13 | 97.83 | 98.97 | 98.36 | 98.43 | 0.05 | 0.37 |
| NYHA III/IV | 26% | 19% | 28% | 22% | 0.20 | <0.001 | 24% | 22% | 24% | 21% | 0.06 | 0.37 |
| Diabetes orally treated | 19% | 19% | 19% | 19% | 0.02 | 0.70 | 18% | 19% | 19% | 19% | 0.02 | 0.73 |
| Diabetes on insulin | 5% | 6% | 3% | 8% | 0.21 | <0.001 | 5% | 6% | 3% | 6% | 0.13 | 0.02 |
| Smoker | 12% | 13% | 14% | 16% | 0.10 | 0.10 | 13% | 13% | 13% | 14% | 0.03 | 0.65 |
| COPD | 7% | 6% | 9% | 6% | 0.13 | 0.05 | 7% | 7% | 7% | 6% | 0.04 | 0.51 |
| PVD | 9% | 8% | 7% | 7% | 0.07 | 0.27 | 7% | 8% | 7% | 8% | 0.04 | 0.61 |
| Prior stroke | 3% | 4% | 2% | 3% | 0.09 | 0.16 | 3% | 3% | 2% | 3% | 0.09 | 0.10 |
| Prior MI | 42% | 44% | 39% | 35% | 0.19 | <0.001 | 41% | 41% | 42% | 39% | 0.06 | 0.38 |
| LVEF <.50 | 28% | 26% | 23% | 21% | 0.16 | 0.01 | 26% | 25% | 25% | 23% | 0.06 | 0.39 |
| Caucasian | 91% | 92% | 88% | 92% | 0.15 | 0.02 | 91% | 92% | 91% | 93% | 0.07 | 0.28 |
| On pump | 56% | 42% | 52% | 39% | 0.35 | 0.00 | 49% | 46% | 48% | 46% | 0.07 | 0.29 |

Table 1. Distribution of pre-treatment variables (as mean or percentage) before (unweighted) and after (weighted) propensity score

ASMD: absolute standardised mean difference; SD= standard deviation for all patients; P-SIMA: pedicled single internal mammary artery; P-BIMA: pedicled bilateral internal mammary arteries; S-SIMA: skeletonized SIMA; S-BIMA: skeletonized bilateral internal mammary arteries; ESS: effective sample size; BMI: body mass index; NYHA: New York Heart Association; COPD: chronic obstructive pulmonary disease; PVD: peripheral vascular disease; MI: myocardial infarction; LVEF: left ventricular ejection fraction

Table 2. Outcomes among treatment groups.

| | All SWC | | Severe SWC | 30 day | 1 year | |
|---|------------|------------|---------------|----------------|-----------|-----------|
| | (n=219) | | (n=150) | mortality | mortality | |
| | | | | (n= 31) | (n=55) | |
| | | All | SWC requiring | Sternal wound | | |
| | | (n=75) | antibiotics | reconstruction | | |
| | | | (n=50) | (n=25) | | |
| P-SIMA (n=607) | 58 (9.5%) | 20 (3.3 %) | 14(2.3%) | 6 (1.0%) | 8 (1.3%) | 13 (2.1%) |
| S-SIMA (n=512) | 41(8.0%) | 14 (2.7 %) | 12(2.3%) | 2(0.4%) | 8 (1.6%) | 15 (2.9%) |
| P-BIMA (n=459) | 74 (16.1%) | 24 (5.2 %) | 17(3.7%) | 7(1.5%) | 7 (1.5%) | 12 (2.6%) |
| S-BIMA (n=478) | 46(9.6%) | 17 (3.7 %) | 7(1.5%) | 10(2.1%) | 8 (1.7%) | 15 (3.1%) |
| χ^2 tests P P-SIMA as reference | | | | | | |
| S-SIMA | 0.39 | 0.60 | 1 | 0.30 | 0.80 | 0.44 |
| P-BIMA | 0.0014 | 0.12 | 0.19 | 0.57 | 0.79 | 0.68 |
| S-BIMA | 1 | 0.86 | 0.37 | 0.20 | 0.62 | 0.33 |

P-SIMA: pedicled single internal mammary artery; P-BIMA: pedicled bilateral internal mammary arteries; S-SIMA: skeletonized

SIMA; S-BIMA: skeletonized bilateral internal mammary arteries; SWC: sternal wound complication

Table 3. Propensity Score weighted effect (OR[95%CI]) of internal mammary artery harvesting on sternal wound complication.

| | Overall | Diabetes On insulin | Diabetes Orally treated | Not Diabetic | Female | Male | BMI≥30 | BMI<30 |
|-----------------------------|-------------------------|------------------------|----------------------------|---------------------|---------------------|---------------------|---------------------|---------------------|
| Comparison P-SIMA as ref | N=2056 | N=118 | N=386 | N=1552 | N=283 | N=1773 | N=631 | N=1425 |
| P-BIMA | 1.80 [1.23- 2.63] | 4.05 [0.86-19.21] | 1.41 [0.58-3.45] | 1.84 [1.18-2.85] | 1.08 [0.41-2.83] | 1.96 [1.30-2.98] | 2.07 [1.09-3.90] | 1.67 [1.03-2.68] |
| S-SIMA | 0.89 [0.57-1.38] | 1.35 [0.29-6.15] | 1.25 [0.49-3.19] | 0.75 [0.43-1.29] | 0.72 [0.27-1.90] | 0.91 [0.55-1.51] | 1.46 [0.73-2.90] | 1.09 [0.65-1.83] |
| S-BIMA | 1.00 [0.65-1.53] | 1.92 [0.48-7.73] | 1.54 [0.64-3.73] | 0.78 [0.46-1.34] | 1.59 [0.65-3.91] | 0.86 [0.52-1.42] | 0.83 [0.39-1.80] | 0.59 [0.32-1.09] |

Bold: P<0.05; OR: Odds ratio; CI: confidence interval; P-SIMA: pedicled single internal mammary artery; P-BIMA: pedicled bilateral

internal mammary arteries; S-SIMA: skeletonized SIMA; S-BIMA: skeletonized bilateral internal mammary arteries; BMI: Body Mass

index

Table 4. Results of double robust Propensity Score-weighted analysis on the incidence

of any sternal wound complication

| | OR | 95%CILL | 95%CI UL | Р |
|-------------------------|------|---------|----------|--------|
| P-BIMA vs P-SIMA | 1.85 | 1.25 | 2.74 | 0.002 |
| S-SIMA vs P-SIMA | 0.98 | 0.64 | 1.52 | 0.94 |
| S-BIMA vs P-SIMA | 0.87 | 0.55 | 1.36 | 0.53 |
| Age† | 1.00 | 0.99 | 1.02 | 0.77 |
| Female | 1.58 | 1.07 | 2.34 | 0.02 |
| BMI† | 1.08 | 1.04 | 1.13 | <0.001 |
| Creatinine† | 0.99 | 0.98 | 1.00 | 0.01 |
| NYHA III-IV | 1.01 | 0.70 | 1.45 | 0.96 |
| Diabetes orally treated | 1.20 | 0.82 | 1.74 | 0.34 |
| Diabetes on insulin | 2.17 | 1.29 | 3.66 | 0.003 |
| Smoking | 1.27 | 0.83 | 1.95 | 0.27 |
| COPD | 1.23 | 0.70 | 2.18 | 0.47 |
| PVD | 0.81 | 0.44 | 1.48 | 0.49 |
| Prior stroke | 1.67 | 0.80 | 3.50 | 0.17 |
| Prior MI | 0.94 | 0.68 | 1.30 | 0.70 |
| LVEF<.50 | 1.02 | 0.71 | 1.46 | 0.91 |
| Caucasian | 1.09 | 0.79 | 1.50 | 0.59 |

† used as continuous variable; Odds ratio; LLCI: confidence interval lower limit; CI UL:

confidence interval upper limit

P-SIMA: pedicled single internal mammary artery; P-BIMA: pedicled bilateral internal mammary arteries; S-SIMA: skeletonized SIMA; S-BIMA: skeletonized bilateral internal mammary arteries; BMI: body mass index; NYHA: New York Heart Association; COPD: chronic obstructive pulmonary disease; PVD: peripheral vascular disease; MI: myocardial infarction; LVEF: left ventricular ejection fraction

95%CI LL 95%CI UL Ρ OR P-BIMA vs P-SIMA 1.61 0.85 3.07 0.15 S-SIMA vs P-SIMA 1.14 0.56 2.31 0.71 S-BIMA vs P-SIMA 0.92 0.43 1.98 0.82 1.00 0.97 1.03 0.79 Age[†] Female 2.48 1.38 4.45 0.002 1.11 1.04 1.18 0.001 BMI† 0.76 1.00 0.99 1.01 Creatinine[†] NYHA III-IV 0.42 0.57 0.83 1.61 **Diabetes orally treated** 1.78 1.00 3.16 0.049 Diabetes on insulin 2.72 1.25 5.92 0.01 1.72 88.0 3.35 0.11 Smoking COPD 2.08 0.97 4.46 0.06 PVD 0.53 0.17 1.66 0.27 Prior stroke 1.74 0.62 4.90 0.29 Prior MI 0.54 0.92 1.59 0.77 LVEF<.50 1.03 0.56 1.87 0.93 Caucasian 1.26 0.73 2.18 0.40

Table 5. Results of double robust Propensity Score-weighted analysis on the

incidence of severe sternal wound complication

† used as continuous variable; Odds ratio; LLCI: confidence interval lower limit; CI UL:

confidence interval upper limit

P-SIMA: pedicled single internal mammary artery; P-BIMA: pedicled bilateral internal mammary arteries; S-SIMA: skeletonized SIMA; S-BIMA: skeletonized bilateral internal mammary arteries; BMI: body mass index; NYHA: New York Heart Association; COPD: chronic obstructive pulmonary disease; PVD: peripheral vascular disease; MI: myocardial infarction; LVEF: left ventricular ejection fraction

Figure Legends

Figure 1. Time from index operation to any and severe sternal wound complication Figure 2. Incidence of any sternal wound complication according to internal mammary artery harvesting strategies. (P-SIMA: pedicled single internal mammary artery; P-BIMA: pedicled bilateral internal mammary arteries; S-SIMA: skeletonized SIMA; S-BIMA: skeletonized bilateral internal mammary arteries)