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Association Between Body Surface Area and Outcomes After Percutaneous Coronary Intervention

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Purpose

Obesity is a well-known risk factor for adverse cardiovascular events, but some studies suggest higher body mass index (BMI) is associated with better outcomes after ST-segment elevation myocardial infarction (STEMI). We sought to determine the effect of body surface area (BSA) on adverse events after primary percutaneous coronary intervention (PCI) for STEMI and how this relates to the reported obesity paradox theory.

Methods

We analyzed a prospective registry of patients with STEMI who underwent primary PCI at a tertiary care hospital from 2003 to 2009. Post-PCI complications and 1-year all-cause mortality were compared across BSA quartiles. Relationship with 1-year mortality was compared between BSA and BMI using logistic regression.

Results

Of 2,195 study patients (31.5% women), mean BSA and BMI were 2.0 ± 0.3 m² and 29.2 ± 6.2 kg/m², respectively. The 1-year all-cause mortality from the lowest to highest quartiles of BSA was 11.0%, 6.5%, 5.5% and 5.1%, P<0.001. Over a mean 5-year follow-up, there was a 76% relative risk reduction in death for each 1 m² increase in BSA. Higher BSA was associated with lower incidence of cardiogenic shock, acute renal failure, coronary dissection and vascular and bleeding complications post-PCI. In multivariate analysis, BSA remained strongly predictive of 1-year mortality (odds ratio 0.4 per m² of BSA, 95% confidence interval 0.15–0.9), but BMI showed no independent association with mortality (odds ratio 0.99, 95% confidence interval 0.95–1.04).

Conclusions

In STEMI patients undergoing primary PCI, high BSA is associated with lower mortality and complication rates. BMI is not independently associated with 1-year mortality after adjusting for BSA and sex. (J Patient-Centered Res Rev. 2015;2:9-16.)

Keywords

body mass index, body size, mortality, myocardial infarction, coronary angioplasty

Primary percutaneous coronary intervention (PCI) is the gold standard approach to reperfusion therapy in patients with ST-segment elevation myocardial infarction (STEMI) when there are no contraindications.1 There are modifiable procedural and clinical factors that affect the outcome of primary PCI,2-4 but there also are patient characteristics like sex5,6 and body mass index (BMI)7,8 that are associated with PCI outcomes. Despite multiple studies that have been done to determine the association between BMI and the outcome of PCI, there are still conflicting data in this regard.9,12 For example, Sarno et al. studied 1,707 patients who underwent PCI and followed them for one year.11 They found that BMI of >30 kg/m² was an independent factor for major cardiac events and stent thrombosis irrespective of stent type. On the other hand, a study of 6,560 patients with non-STEMI showed that patients with high BMI had better 30-day outcomes compared to patients with lower BMI, but the benefit was not present after 30 days and both groups had the same risk for primary endpoints (cardiovascular death, myocardial infarction and recurrent ischemia) when followed for one year.14

Obesity is a well-established risk factor for cardiovascular disease,15 hence findings of better prognosis in obese patients than in normal-weight patients who are treated for heart failure, coronary
disease and peripheral artery disease led to the term “obesity paradox.” This suggests that obesity is a marker for a different unidentified factor that confers favorable cardiovascular outcomes. A positive correlation has been shown between body surface area (BSA), BMI and coronary artery size. The calculations for BSA and BMI both use height and weight, which could lead to confounding between the two. BSA is reflective of body and organ sizes, whereas BMI is a measure of body fat. It is unclear how BSA relates to the “obesity paradox” debate. Particularly, the effect of BSA on outcomes after primary PCI in patients with STEMI has not been previously studied. We hypothesized that patients with high BSA who undergo primary PCI for STEMI will have a lower rate of adverse outcomes than those with low BSA. We also compared the strength of association between mortality after primary PCI and BSA to the strength of association with BMI.

METHODS
This was a nonrandomized, retrospective, registry-based study designed to test the association between BSA and clinical outcomes after PCI in patients presenting with STEMI. Local institutional review board approval was obtained prior to initiation of the study.

Patients and Variables
A consecutive population of all patients 18 years or older who presented at our tertiary care hospital with STEMI and were revascularized with primary PCI from January 2003 to December 2009 were included in the analysis. Data was extracted from the local National Cardiovascular Data Registry (NCDR) CathPCI Registry®. Hospital medical records were reviewed and relevant clinical information manually extracted for each patient. The National Death Index was queried to identify all out-of-hospital deaths.

Standard NCDR data definitions (www.ncdr.com/webncdr/cathpci) were used for all variables. The main independent variables were BMI, BSA and sex. BMI was calculated as weight (in kilograms)/height^2 (in meters). The commonly used Mosteller formula calculated BSA in m^2 as √(weight x height/3600), with weight in kilograms and height in meters. Outcome variables assessed were in-hospital death, post-PCI bleeding, coronary dissection, access-site vascular complications, cardiogenic shock and acute renal failure as well as death from any cause within one year. Adverse events were adjudicated by reviewing patients’ electronic hospital records.

Data Analysis
There are no universal cut-offs for BSA, hence the study sample was divided into quartiles of BSA and data compared between quartiles. Continuous variables were expressed as mean ± standard deviation and compared using ANOVA. Categorical variables were reported as percentages and compared using chi-square test. Linear association among BSA quartile outcomes was investigated using the linear applications of the ANOVA and chi-square tests. Logistic regression was used to compare outcomes across quartiles of BSA adjusting for baseline patient comorbidities. Overall survival was compared among quartiles of BSA using Kaplan-Meier analysis and the log-rank test. Cox proportional hazards regression was used to analyze long-term survival. In survival analysis, time was measured beginning from the index primary PCI. All P-values were two-sided.

RESULTS
There were 2,197 patients who underwent primary PCI for STEMI during the 7-year study period. Their baseline characteristics are shown in Table 1. Mean age was 61.5 ± 13.1 years, and there was a significant trend for BSA to decrease with increasing age. There was a strong association between sex and BSA. The proportion of women per BSA quartile decreased progressively from lowest to highest quartile. There was a strong correlation between BSA and BMI. Figure 1 shows that this correlation was strong in both male and female patients. Interestingly, however, at every level of BMI, BSA was higher in men than in women. The frequency of hypertension, smoking, prior myocardial infarction and prior PCI was similar across all quartiles of BSA. Cardiac symptomatology was similar between BSA quartiles, with about 82% arriving within 6 hours of symptom onset and 7% presenting with cardiogenic shock. Overall, 81% of patients had door-to-balloon time of <90 minutes and there was no difference in the arterial territory or number of lesions revascularized, or in the number of stents used. Mean stent diameter increased with increasing BSA. There was a significant correlation between BSA and stent
### Table 1. Baseline patient characteristics in all patients and according to quartile of body surface area

<table>
<thead>
<tr>
<th>Variable</th>
<th>All patients (N=2,197)</th>
<th>&lt;1.8 (n=437)</th>
<th>1.8 to 2 (n=590)</th>
<th>&gt;2 to 2.2 (n=586)</th>
<th>&gt;2.2 (n=584)</th>
<th>P*</th>
</tr>
</thead>
<tbody>
<tr>
<td>Age (years), mean ± SD</td>
<td>61.5 ± 13.1</td>
<td>69.5 ± 13.9</td>
<td>63.3 ± 12.8</td>
<td>59.3 ± 11.5</td>
<td>55.9 ± 10.9</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td>Female, %</td>
<td>31.2</td>
<td>37.7</td>
<td>34.6</td>
<td>17.6</td>
<td>9.1</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td>BMI (kg/m^2), mean ± SD</td>
<td>29.2 ± 6.1</td>
<td>24.2 ± 6.0</td>
<td>27.1 ± 3.5</td>
<td>29.8 ± 4.5</td>
<td>34.5 ± 5.4</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td>Smoker, %</td>
<td>44.8</td>
<td>37.7</td>
<td>46.9</td>
<td>46.9</td>
<td>45.9</td>
<td>0.11</td>
</tr>
<tr>
<td>Hypertension, %</td>
<td>57.9</td>
<td>61.1</td>
<td>55.5</td>
<td>56.0</td>
<td>59.9</td>
<td>0.16</td>
</tr>
<tr>
<td>Dyslipidemia, %</td>
<td>62.0</td>
<td>55.2</td>
<td>60.1</td>
<td>65.4</td>
<td>65.6</td>
<td>0.001</td>
</tr>
<tr>
<td>Diabetes, %</td>
<td>16.9</td>
<td>15.8</td>
<td>14.9</td>
<td>15.5</td>
<td>21.1</td>
<td>0.05</td>
</tr>
<tr>
<td>Previous MI, %</td>
<td>16.0</td>
<td>19.8</td>
<td>14.9</td>
<td>15.2</td>
<td>14.9</td>
<td>0.12</td>
</tr>
<tr>
<td>Previous PCI, %</td>
<td>21.4</td>
<td>22.5</td>
<td>20.7</td>
<td>22.0</td>
<td>20.5</td>
<td>0.83</td>
</tr>
<tr>
<td>Previous CABG, %</td>
<td>5.6</td>
<td>5.7</td>
<td>7.5</td>
<td>6.3</td>
<td>2.7</td>
<td>0.004</td>
</tr>
<tr>
<td>Symptom duration</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td>0.28</td>
</tr>
<tr>
<td>≤6 hours, %</td>
<td>82.0</td>
<td>76.9</td>
<td>83.5</td>
<td>81.4</td>
<td>84.8</td>
<td></td>
</tr>
<tr>
<td>6–12 hours, %</td>
<td>8.7</td>
<td>11.0</td>
<td>8.2</td>
<td>8.6</td>
<td>7.7</td>
<td></td>
</tr>
<tr>
<td>&gt;12 hours, %</td>
<td>9.3</td>
<td>12.1</td>
<td>8.2</td>
<td>10.1</td>
<td>7.5</td>
<td></td>
</tr>
<tr>
<td>Door-to-balloon time &lt;90 min, %</td>
<td>81.1</td>
<td>76.4</td>
<td>81.7</td>
<td>83.3</td>
<td>81.8</td>
<td>0.04</td>
</tr>
<tr>
<td>Transfer patient, %</td>
<td>49.6</td>
<td>49.5</td>
<td>43.4</td>
<td>50.7</td>
<td>54.8</td>
<td>0.01</td>
</tr>
<tr>
<td>Cardiogenic shock, %</td>
<td>7.1</td>
<td>8.2</td>
<td>6.6</td>
<td>6.7</td>
<td>7.0</td>
<td>0.74</td>
</tr>
<tr>
<td>LVEF ≤40, %</td>
<td>33.2</td>
<td>40.4</td>
<td>34.3</td>
<td>30.8</td>
<td>29.3</td>
<td>0.04</td>
</tr>
<tr>
<td>Arteries revascularized†</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Left main, %</td>
<td>0.5</td>
<td>0.7</td>
<td>0.2</td>
<td>0.7</td>
<td>0.7</td>
<td>0.55</td>
</tr>
<tr>
<td>Left anterior descending, %</td>
<td>38.3</td>
<td>41.0</td>
<td>38.0</td>
<td>35.8</td>
<td>39.0</td>
<td>0.39</td>
</tr>
<tr>
<td>Left circumflex, %</td>
<td>19.2</td>
<td>17.6</td>
<td>18.8</td>
<td>21.3</td>
<td>18.5</td>
<td>0.45</td>
</tr>
<tr>
<td>Right coronary, %</td>
<td>49.4</td>
<td>48.1</td>
<td>50.3</td>
<td>51.2</td>
<td>47.8</td>
<td>0.59</td>
</tr>
<tr>
<td>Lesions treated (n), mean ± SD</td>
<td>1.3 ± 0.6</td>
<td>1.3 ± 0.6</td>
<td>1.3 ± 0.6</td>
<td>1.4 ± 0.7</td>
<td>1.3 ± 0.5</td>
<td>0.16</td>
</tr>
<tr>
<td>Stents (n), mean ± SD</td>
<td>1.7 ± 1.0</td>
<td>1.6 ± 1.1</td>
<td>1.7 ± 1.2</td>
<td>1.7 ± 1.0</td>
<td>1.6 ± 0.9</td>
<td>0.53</td>
</tr>
<tr>
<td>Stent diameter (mm), mean ± SD</td>
<td>3.0 ± 0.5</td>
<td>2.9 ± 0.4</td>
<td>3.0 ± 0.4</td>
<td>3.1 ± 0.5</td>
<td>3.3 ± 0.3</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td>Bare-metal stent, %</td>
<td>46.6</td>
<td>49.0</td>
<td>44.7</td>
<td>45.1</td>
<td>48.3</td>
<td>0.39</td>
</tr>
<tr>
<td>Drug-eluting stent, %</td>
<td>41.1</td>
<td>34.6</td>
<td>43.4</td>
<td>43.9</td>
<td>41.1</td>
<td>0.01</td>
</tr>
<tr>
<td>Unknown stent type, %</td>
<td>12.2</td>
<td>16.5</td>
<td>11.9</td>
<td>11.1</td>
<td>10.6</td>
<td>0.02</td>
</tr>
</tbody>
</table>

*P*Two-tailed *P*-value calculated using chi-square test for categorical variables and ANOVA for continuous variables.

†Adds up to more than 100% because some patients had more than one artery revascularized.

BMI, body mass index; CABG, coronary artery bypass graft; LVEF, left ventricular ejection fraction; MI, myocardial infarction; PCI, percutaneous coronary intervention; SD, standard deviation.

Overall angiographic success, defined as residual stenosis of <30% with TIMI-3 flow, was 97.5%, and was similar across all BSA quartiles (P=0.69). There were some significant differences in post-PCI clinical outcomes between BSA quartiles (Figure 2). There was a significant inverse linear relationship between post-PCI complications and BSA quartile. The rate of adverse events decreased from lowest (≤1.8 m^2^) to highest BSA quartile (>2.2 m^2^). Although coronary dissection was generally rare, it had the highest relative increase, 27-fold from the highest BSA quartile to the
lowest quartile. There was approximately a doubling in the incidence of acute renal failure, vascular complications (mainly pseudoaneurysms), bleeding events and cardiogenic shock between the highest and lowest BSA quartiles. The incidence of repeat myocardial infarction prior to discharge was 0.8% overall, without any difference between BSA quartiles (P=0.16). The rates of emergency coronary artery bypass surgery were 2.7%, 2.5%, 1.9% and 2.6% in the lowest, second, third and highest quartiles, respectively (P=0.95).

These differences in post-PCI complications were accompanied by a significant decrease in mortality with increasing BSA. In-hospital mortality was 7.4%, 4.6%, 3.4% and 3.4% in the lowest, second, third and highest BSA quartiles, respectively (P=0.001). In patients with STEMI, the better prognosis associated with high BSA that is seen early after primary PCI persisted long term. Figure 3 shows Kaplan-Meier survival curves for the four quartiles of BSA. The 1-year overall survival rates were 88.3%, 93.7%, 94.5% and 94.9% in the lowest, second, third and highest BSA quartiles, respectively (P_{trend}<0.001). Over a mean follow-up period of 5 years (range: 2–9 years), the hazard ratios of all-cause death in the second, third and highest quartiles of BSA when compared to the lowest quartile were 0.63 (95% CI 0.46–0.87), 0.48 (95% CI 0.34–0.68) and 0.43 (95% CI 0.31–0.62). Considering BSA as a continuous variable, there was a 76% relative risk reduction in death for every 1 m\(^2\) increase in BSA (95% CI 61.0–85.0, P<0.001).

We compared the strengths of association of BSA with 1-year mortality and the association of BMI with 1-year mortality (Table 2). On univariate analysis, increases in both BSA and BMI were associated with decreases in 1-year mortality, but the association was stronger with BSA than BMI. We performed multivariate analysis adjusting for BSA, BMI and sex because all three variables affect each other and mortality. We also adjusted for age, symptom duration, shock, door-to-balloon time, coronary anatomy and comorbidities. BSA maintained a significant independent association with 1-year mortality, but BMI and sex were no longer associated with 1-year mortality.
DISCUSSION

In this study we divided 2,197 patients with STEMI who underwent primary PCI into quartiles of BSA and compared clinical outcomes between them. Going from the lowest (<1.8 m$^2$) to the highest quartile (>2.2 m$^2$) of BSA, there were significant reductions in post-PCI complications like coronary dissection, cardiogenic shock, acute renal failure, bleeding, vascular complications and in-hospital death. Long-term all-cause mortality also decreased with increasing BSA. BMI was associated with 1-year mortality only on univariate analysis. In the multivariate analysis, only BSA remained associated with 1-year mortality whereas BMI showed no independent effect.

Previous studies have investigated the association between BMI and PCI outcomes in patients with acute coronary syndromes and reported results that are in conflict with the obesity paradox.\textsuperscript{7,8,12,24} In a study of 4,972 patients who underwent PCI with drug-eluting stents, patients with higher BMI had a higher risk of stent thrombosis compared to patients with lower BMI; and after multivariate analysis, obesity was still an independent predictor for thrombotic events.\textsuperscript{12} Similarly, Ellis et al. evaluated the association of BMI and PCI outcomes on 3,571 patients and showed BMI of >35 was associated with higher mortality.\textsuperscript{24} Different mechanisms have been proposed to explain these findings. First, obese patients have higher levels of coagulation factors, including factors VII, VIII, plasminogen activator inhibitor-1 and fibrinogen, that make them relatively hypercoagulable.\textsuperscript{25,26} Periprocedural antithrombotic regimens may be inadequate in very obese patients due to their unusual weight.\textsuperscript{24,27} Other studies have paradoxically shown that higher BMI was associated with better outcomes in patients who undergo PCI, lending support to the obesity paradox.\textsuperscript{8-11} Obesity has long been associated with an increased risk of death from all causes, diabetes mellitus, hypertension, dyslipidemia and atherosclerosis.\textsuperscript{28,29} These well-documented adverse outcomes of obesity lead us to think that the observation

Table 2. Relative strengths of association of body surface area and body mass index with 1-year mortality

<table>
<thead>
<tr>
<th>Variable</th>
<th>Univariate</th>
<th></th>
<th></th>
<th></th>
<th>Multivariate adjusted*</th>
<th></th>
<th></th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Odds ratio</td>
<td>95% CI</td>
<td>$P$</td>
<td>Odds ratio</td>
<td>95% CI</td>
<td>$P$</td>
<td>Odds ratio</td>
</tr>
<tr>
<td>Body surface area (m$^2$)$^1$</td>
<td>0.24</td>
<td>0.13-0.45</td>
<td>&lt;0.0001</td>
<td>0.40</td>
<td>0.15-0.90</td>
<td>0.01</td>
<td>0.40</td>
</tr>
<tr>
<td>Body mass index (kg/m$^2$)$^1$</td>
<td>0.96</td>
<td>0.93-0.99</td>
<td>0.019</td>
<td>0.99</td>
<td>0.95-1.04</td>
<td>0.82</td>
<td>0.99</td>
</tr>
<tr>
<td>Male$^2$</td>
<td>0.51</td>
<td>0.36-0.71</td>
<td>&lt;0.0001</td>
<td>0.68</td>
<td>0.43-1.07</td>
<td>0.10</td>
<td>0.68</td>
</tr>
</tbody>
</table>

*Adjusted for listed variables plus age, symptom duration, shock, door-to-balloon time, coronary anatomy and comorbidities.

$^1$Odds ratios calculated per unit increase in body surface area or body mass index.

$^2$Compares male versus female.

CI, confidence interval.
of better post-PCI outcomes in obese patients is due to some other confounding factor. More recent studies have questioned the validity of an obesity paradox in patients treated with PCI. In the Swedish Coronary Angiography and Angioplasty Study, there was a biphasic relationship between BMI and mortality in which mortality decreased with increasing BMI up to 35 kg/m², after which mortality started to increase. A recent study of patients with STEMI from the NCDR registry also reported lower in-hospital mortality, bleeding and adverse cardiac events with increasing BMI, except for patients with BMI of >40 kg/m², who had worse prognosis. In this study, however, obese patients were more likely to be younger and have less extensive coronary artery disease. After adjustment for baseline differences, there was no significant difference in outcomes between normal-weight, overweight and mildly obese patients, putting into question the obesity paradox.

Our study also found that while mortality decreased with increasing BMI on univariate analysis, BMI had no effect on mortality on multivariate analysis that adjusted for baseline confounders. BSA, on the other hand, maintained a strong association with mortality both in unadjusted and adjusted analyses. The strong correlation we demonstrated between BSA and BMI show the confounding effect they have on each other. This leads us to propose that much of the previously observed association between BMI and prognosis after myocardial infarction or PCI can be attributed to the confounding effect of BSA. We suggest that BSA is the true independent predictor of outcomes after primary PCI in patients with STEMI.

Although there are no studies that directly test the effect of BSA on post-PCI outcomes, there have been indirect inferences in recent literature. In a large study based on the NCDR registry with an identical definition of bleeding to ours, the bleeding risk after PCI was not associated to BMI but was decreased with increase in BSA. Bleeding risk of women was double that of men (7.8% vs. 3.7%, P<0.01), which was attributable in part to lower BSA of women. Similarly, Peterson et al. evaluated 109,708 patients treated with PCI and reported a higher risk for stroke, vascular complications and in-hospital mortality in women than men, but there was no difference in mortality after adjusting for BSA. The authors concluded that BSA, rather than sex, is an independent risk factor for post-PCI mortality. There is a well-demonstrated association between high BSA and better outcomes after coronary artery bypass grafting (CABG). High BSA is associated with better graft patency, leading to lower incidence of adverse cardiac events after CABG. There is strong evidence showing that BSA and coronary artery diameter are inversely related with mortality rate in patients who undergo CABG. Much of the reports of higher mortality in women compared to men after CABG have been attributed to the smaller BSA of women.

The positive effect of BSA on coronary artery diameter, cardiac size and blood volume may explain the effect of BSA on post-PCI outcomes. In populations of patients both with and without established coronary artery disease, there is a significant increase in the diameter of coronary arteries with increasing BSA irrespective of sex. If stent diameter can be used as a surrogate for coronary artery diameter, this will be similar to our finding of a positive correlation between stent diameter and BSA. In the PCI population, large coronary diameter was associated with lower rates of restenosis, repeat revascularization, stent thrombosis, recurrent myocardial infarction and death. In the general population, BSA has correlated positively with other measures of heart size like left ventricular mass, ventricular volumes and blood volume. Larger arteries also prevent sheath/artery mismatch, which reduces access site complications. We theorize that prognosis is better in patients with high BSA after primary PCI following STEMI because they have more myocardial and vascular reserve that can compensate for the injury acutely as well as larger coronary arteries that lower the risk of cardiac events.

Limitations

Our study reflects the experience of a tertiary care center with a rare STEMI program that requires the catheterization laboratory staff and interventional cardiologist to remain in-house 24 hours/day, 7 days/week. Although our overall event rates may be different from other centers, we do not anticipate a different relationship between BSA and major outcomes. The use of a registry could introduce errors due to missing data and inaccurate data entry. We manually reviewed hospital records to crosscheck clinical
outcomes on every patient. Bleeding complications captured in our study were major ones associated with drop in hemoglobin of ≥3 g/dL, blood transfusion or procedural intervention to control bleeding. Less significant bleeding events, albeit important, were not captured. That said, our findings are valid and constitute an important consideration for clinicians when caring for patients with STEMI.

CONCLUSIONS

There is an inverse relationship between body surface area and adverse clinical events in patients with ST-segment elevation myocardial infarction who undergo primary percutaneous coronary intervention. Increase in BSA from the lowest to highest quartile was associated with decreasing rates of short- and long-term mortality as well as decreasing rates of coronary dissection, cardiogenic shock, acute renal failure, bleeding and vascular complications. Body mass index is not independently associated with 1-year mortality after adjusting for BSA and sex. Clinicians should pay attention to the increased risk of adverse outcomes in patients with BSA of <1.8 m², and consideration should be given to including BSA in risk adjustment models that compare clinical outcomes in patients undergoing primary PCI for STEMI.

Patient-Friendly Recap

• A heart attack (myocardial infarction) occurs when proper blood flow to the heart is stopped, usually due to blockage in one or more coronary arteries.
• Percutaneous coronary intervention (PCI) is a less invasive procedure used to unclog arteries and prevent further heart damage.
• Despite having an overall increased risk of heart disease, obese patients seem to do better following PCI.
• The authors report this “obesity paradox” can be attributed to increased body surface area, rather than body mass index (BMI) as previously reported.
• Patients with more body surface area tend to have wider arteries, which might explain why they experience fewer complications from PCI.

Acknowledgments

The authors gratefully acknowledge Brian Miller and Brian Schurrer of Aurora Sinai Medical Center for their help with figures.

Conflicts of Interest

None.

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