Regional Left Ventricular Systolic Function in Humans During Off-Pump Coronary Bypass Surgery

Philip M. Brown, Jr, MD; Victor B. Kim, MD; Bret J. Boyer, BS; Robert M. Lust, PhD; W. Randolph Chitwood, Jr, MD; Joseph R. Elbeery, MD

Background—Controversy exists as to whether off-pump CABG with local occlusion results in clinically significant myocardial ischemia during the occlusion period. This study was undertaken to delineate the effects of transient local coronary artery occlusion on regional systolic function.

Methods and Results—Eight consenting patients undergoing left internal mammary to left anterior descending coronary artery (LAD) bypass were instrumented with a left ventricular pressure catheter and 2 subepicardial cylindrical ultrasonic dimension transducers placed in the minor axis dimension in the region served by the LAD. A digital sonomicrometer was used to collect data before, during, and after coronary occlusion from which percent systolic shortening and pressure-dimension loops were derived. Measuring devices were removed immediately after the final time point. All patients tolerated the procedure well, and there were no complications. Average duration of local occlusion needed for CABG was 15.9 ± 4.4 minutes (range, 12 to 26 minutes). Local occlusion was associated with a decrease in peak systolic shortening from 5.8 ± 0.8% to 1.8 ± 0.8%. In all cases, function returned to baseline after restoration of flow. Pressure-dimension loops confirmed these findings and no evidence of diastolic creep. Linear repression analysis of degree of stenosis versus change in segmental shortening revealed a significant inverse correlation.

Conclusions—Local occlusion of the LAD resulted in a transient decrease in myocardial function during occlusion with complete recovery during reperfusion. This change was less significant with increasing degrees of coronary stenosis. These data suggest that local occlusion is not associated with permanent myocardial injury but that ischemic changes do occur that may be clinically significant, especially in patients with lesser degrees of coronary stenosis. (Circulation. 1999;100[suppl II]:II-125–II-127.)

Key Words: coronary disease ■ revascularization ■ surgery
minor axis plane to subtend the myocardial region supplied by the LAD. The crystals were secured with 5 to 0 prolene sutures to prevent dislodgement. Analog data (200 Hz) collected for 8 seconds were discreet time points during the procedure. These points were (1) before LAD occlusion (baseline), (2) just before LAD release (occlusion, 10 to 24 minutes), and (3) before incision closure (reperfusion, 7 to 10 minutes total). Analog recordings were converted to digital data points by a sonomicrometer (Sonometrics). The sonomicrometer has a range of measurement from 10 to 120 mm, with the smallest measurable change in distance being 0.024 mm. This is linear throughout the entire measurement range. At the conclusion of data collection, all monitoring devices were removed, and the patients’ wounds were closed as per standard practice.

Data Analysis and Statistical Comparison
Digital data were transferred into Microsoft Excel for further analysis. The cardiac cycle was defined by use of the first derivative of the pressure/time interval (dP/dt) as described previously, and pressure-dimension loops were generated by plotting all continuous measurements recorded for 8 seconds at each given time point. Percent systolic shortening (%SS) was calculated using this formula: 

\[
\text{%SS} = \frac{(\text{EDL} - \text{ESL})}{\text{EDL}} \times 100
\]

where EDL is end-diastolic length and ESL is end-systolic length.

Linear regression analysis was used to compare the decrease in percent systolic shortening to the degree of native coronary artery stenosis. Paired t tests and the Wilcoxon signed-rank test were used to compare values at different time points. Numerical data were expressed as mean±SD. Statistical significance was defined as \( P<0.05 \).

Results
All patients tolerated the procedure well, and no complications occurred. Patients in the study group had normal left ventricular function as assessed on ventriculography. LAD runoff was judged as good (n=2) or excellent (n=6) on the basis of preoperative angiographic interpretation by the surgeon. Stenosis of the LAD ranged from 70% stenosis to total occlusion (n=2). Collateral arterial supply was graded as minimal (n=5), moderate (n=1), or extensive (n=2). This was judged by the amount of back bleeding from the LAD during proximal occlusion after the arteriotomy was made. No patient developed significant ST-segment elevation during the operation, nor were any significant hemodynamic changes noted.

Duration of LAD occlusion was 15.9±4.4 minutes (range, 12 to 26 minutes). Regardless of occlusion length, there was a significant decline in systolic shortening in patients with nonoccluded LADs from 5.8±0.8% to 1.8±0.8% (n=6, \( P<0.02 \)). In all cases, dysfunction resolved completely after reperfusion (mean, 8.6±2.4; range, 6 to 13 minutes after

---

**Fig 1.** Correlation of percent LAD stenosis vs percent change in shortening fraction during local coronary occlusion.

<table>
<thead>
<tr>
<th>Patient</th>
<th>Age, y</th>
<th>Sex</th>
<th>LAD Stenosis, %</th>
<th>Collaterals</th>
<th>Occlusion Time, min</th>
<th>EDL, mm</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>75</td>
<td>F</td>
<td>70</td>
<td>Minimal</td>
<td>26</td>
<td>12.8</td>
</tr>
<tr>
<td>2</td>
<td>50</td>
<td>M</td>
<td>95</td>
<td>Minimal</td>
<td>17</td>
<td>15.0</td>
</tr>
<tr>
<td>3</td>
<td>59</td>
<td>F</td>
<td>90</td>
<td>Moderate</td>
<td>12</td>
<td>10.7</td>
</tr>
<tr>
<td>4</td>
<td>51</td>
<td>F</td>
<td>80</td>
<td>Minimal</td>
<td>15</td>
<td>11.4</td>
</tr>
<tr>
<td>5</td>
<td>59</td>
<td>M</td>
<td>100</td>
<td>Extensive</td>
<td>12</td>
<td>15.6</td>
</tr>
<tr>
<td>6</td>
<td>59</td>
<td>M</td>
<td>100</td>
<td>Extensive</td>
<td>15</td>
<td>15.6</td>
</tr>
<tr>
<td>7</td>
<td>42</td>
<td>M</td>
<td>80</td>
<td>Minimal</td>
<td>15</td>
<td>16.7</td>
</tr>
<tr>
<td>8</td>
<td>59</td>
<td>M</td>
<td>80</td>
<td>Minimal</td>
<td>15</td>
<td>19.7</td>
</tr>
</tbody>
</table>

EDL indicates end-diastolic length; C, control; I, ischemia; and R, reperfusion.
release of occlusion). Pressure-dimension loops (Figure 1) narrowed during ischemia but returned to baseline and revealed no diastolic creep as evidenced by return of end-diastolic length (the Table).

Linear repression analysis revealed a significant inverse correlation between degree of coronary stenosis and percent decrease in systolic shortening ($P<0.02$) (Figure 2).

Discussion

Detrimental effects of ischemia on regional myocardial function have been well documented. Previous studies have clearly shown that with the loss of active shortening during ischemia, regional systolic stress increases, contributing to regional ischemic dysfunction and facilitating passive elongation of ischemic myocardium. These regional myocardial geometric changes characterized by progressive increases in end-diastolic segment length and decreased end-diastolic wall thickness occur as a time- and load-dependent change identified as creep. Owen et al subsequently showed that significantly increased end-diastolic length persisted for hours after only a 15-minute period of regional ischemia in dogs and that late functional recovery correlated primarily with reversal diastolic creep.

The present study involved measurement of regional systolic function in chronically ischemic human myocardium that was then subjected to a period of acute ischemia during the time of local coronary occlusion necessary for off-pump bypass. A measurable decline in regional left ventricular systolic function was observed, as evidenced by a reduced percent systolic shortening. Pressure-dimension loops confirmed diminished shortening fraction but did not demonstrate evidence of the time-dependent stretch of myocardial segments or diastolic creep. Furthermore, these changes were more significant with lesser degrees of coronary artery stenosis and presence of established collaterals. After reperfusion of blood flow, systolic shortening returned to baseline within minutes. In contrast, previous animal studies have demonstrated prolonged myocardial dysfunction after comparable periods of regional ischemia.

The results of this investigation demonstrate an increased tolerance to ischemia in humans with coronary artery disease. This can be explained in part by the slowly progressive nature of coronary lesions caused by atherosclerosis, in contrast to the normal arteries present in animal studies. The slow disease progression allows development of collaterals and increased ischemia tolerance. This is supported by the inverse correlation noted between the degree of stenosis and decrease in systolic shortening in the present study. In addition, chronic coronary stenosis may induce a “preconditioned” state, rendering the myocardium more tolerant to acute occlusion.

The clinical significance of this investigation is 2-fold. First, myocardial ischemia of the duration necessary to perform a coronary anastomosis does not appear to result in any long-lasting change in myocardial function. Therefore, permanent injury to the myocardium is unlikely to occur. Second, local occlusion does result in transient myocardial dysfunction, the degree of which is greater with lesser coronary stenosis and collaterals. Although no patient in this study had hemodynamic compromise during coronary occlusion, this is a well-known albeit rare complication. These data provide evidence that other strategies, such as coronary shunting, rather than local occlusion might be advised in patients with only moderate coronary stenoses and large myocardial areas at risk.

Acknowledgment

We thank Susan Licata for help in the preparation of this manuscript.

References

Regional Left Ventricular Systolic Function in Humans During Off-Pump Coronary Bypass Surgery
Philip M. Brown, Jr, Victor B. Kim, Bret J. Boyer, Robert M. Lust, W. Randolph Chitwood, Jr and Joseph R. Elbeery

Circulation. 1999;100:II-125-II-127
doi: 10.1161/01.CIR.100.suppl_2.II-125

Circulation is published by the American Heart Association, 7272 Greenville Avenue, Dallas, TX 75231
Copyright © 1999 American Heart Association, Inc. All rights reserved.
Print ISSN: 0009-7322. Online ISSN: 1524-4539

The online version of this article, along with updated information and services, is located on the World Wide Web at:
http://circ.ahajournals.org/content/100/suppl_2/II-125