Effects of cardiopulmonary bypass surgery on coronary flow in children assessed with transthoracic Doppler echocardiography

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Effects of cardiopulmonary bypass surgery on coronary flow in children assessed with transthoracic Doppler echocardiography. Am J Physiol Heart Circ Physiol 293: H1138–H1143, 2007. First published May 4, 2007; doi:10.1152/ajpheart.00025.2007.—Perturbation of coronary blood flow (CF) is an important contributor to myocardium-related complications. The study was primarily designed to assess the impact of cardiopulmonary bypass (CPB) surgery on CF by aid of transthoracic Doppler echocardiography. Changes in CF after off-pump coarcction surgery were also studied. All ultrasounds were performed before and 1 days after surgery. Eighteen children underwent CPB surgery of ventricular left-to-right shunts at the mean age of 6 mo, while off-pump surgery (aortic coarctectomy) was undertaken at the mean age of 10 days in 12 children. After CPB surgery, both left anterior descending coronary artery mean diameter and basal CF increased from 1.7 ± 0.3 to 2.1 ± 0.4 mm (P = 0.001) and 27 ± 10 to 47 ± 15 ml/min (P = 0.0001), respectively. These two coronary variables decreased after off-pump coarctectomy: left anterior descending coronary artery mean diameter from 1.8 ± 0.1 to 1.7 ± 0.1 mm (P = 0.06), and CF from 44 ± 12 to 25 ± 8 ml/min (P = 0.001). The findings are in keeping with the hypothesis that the previously reported impairment of coronary flow reserve after CPB surgery could be due to increase in basal coronary flow after CPB. Off-pump coarctectomy seems to have little impact on CF, as the postsurgical decline in flow in these patients seems to relate to the reduction in cardiac pressure afterload.

congenital heart disease; coronary blood flow; cardiopulmonary bypass surgery

PERTURBATION OF CORONARY BLOOD flow (CF) is an important contributor to myocardium-related complications, such as contractile dysfunction and arrhythmias (14), which sometimes appear during the first week after cardiac surgery on cardiopulmonary bypass (CPB). Coronary flow reserve (CFR) after CPB surgery has been reported to be decreased (7), but the precise pathophysiology remains debatable.

Congenital heart diseases cause several hemodynamic and functional changes that are likely to affect CF. Reduced arterial oxygen saturation, myocardial hypertrophy, increased heart rate, and volume and pressure overload contribute to myocardial oxygen deprivation and CF (8, 17, 18, 22). Oxygen demand regulates coronary flow. The major determinants of the oxygen demand of the myocardium are heart rate, contractility, and wall stress. Wall stress is related to ventricular pressure, chamber diameter, and wall thickness (10). Myocardial oxygen demand is described as rate-pressure product (RPP), the product of heart rate and mean systolic blood pressure (2). The maximal ability of coronary circulation to increase in response to increased cardiac metabolic demand is referred to as CFR. CFR is commonly expressed as the ratio of maximal coronary flow (e.g., by adenosine infusion) to basal flow (5, 15, 16). CPB, together with cardiac surgery, may affect coronary flow.

Myocardial ischemia-reperfusion leads to coronary endothelial dysfunction due to decreased endothelium-dependent relaxation (26). It has been shown in experimental models that endothelial dysfunction after ischemia-reperfusion persists for at least 4–6 wk (21, 32). Moreover, CPB seems to directly affect the arterial smooth muscle cells (40). The aim of this paper was to assess the effects of CPB on the coronary flow in children.

SUBJECTS AND METHODS

Study Population

Eighteen infants (mean age 6 ± 4 mo) referred to our center for CPB surgery due to ventricular or common atrioventricular septal defect and 12 neonates (mean age 10 ± 8 days) with aortic coarctation referred for coarctectomy (off-pump cardiac surgery) were enrolled. The exclusion criteria were as follows: clinical signs of infectious illness or a CRP value before surgery >0.8 mg/l, heart failure, and preoperative therapy with vasoactive drugs. Transthoracic Doppler echocardiography (TTDE) was performed on all children 1 day pre- and 5 ± 1 days postoperatively. The exclusion criteria were as follows: clinical signs of infectious illness or a CRP value before surgery >0.8 mg/l, heart failure, and preoperative therapy with vasoactive drugs. For shunt lesion, age-matched controls with no structural heart disease were investigated (n = 19, mean age 6.5 ± 5 mo). For coarctation group, earlier obtained CF data from 55 neonates with mean age of 5 days were used.

Written consent was obtained from the guardians of all children enrolled in the study. The study was approved by the ethics committee for human research at the Lund University.

TTDE

TTDE examination was performed using Sequoia C512 (Acuson, Mountain View, CA) with 7- to 10-MHz transducer. Standard M- and B-mode and Doppler echocardiographic studies were performed for determining the anatomy and function of the heart. The examination was done by one investigator (EHA).

For CF and flow velocity measurements, the following adjustments were made in the ultrasound device: space time in high frame rate (T1), wall filter was set at two-thirds (F2), the color gain was adjusted to minimize color flow signal scatter (gate 3), and color Doppler mix was on (20, 35). Pulsed Doppler of 4.5 MHz, sweep rate of 100 mm/s, and velocity range of 15–60 cm/s were used. Measurements were

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corrected for the angle between the Doppler beam and the coronary flow direction. True velocity was defined as the measured velocity divided by the cosine of the angle between the Doppler beam and the direction of blood.

The bifurcation of the left main coronary artery was imaged from the standard parasternal short-axis view of the great arteries. The internal dimension of the left anterior descending coronary artery (LAD) was measured at the R-wave with the calipers applied to the inner borders. The velocity scale was decreased to the minimum range and then gradually increased until color signals were optimized within the vessel lumen. After good coronary flow signals were detected, the pulsed Doppler sample volume was placed within the LAD artery 2–3 mm distal to the bifurcation of the left main coronary artery, and the sample volume was adjusted to 0.5–1.0 mm. A sample volume that gave the best quality envelope and pure sound throughout the cardiac cycle was chosen.

All images were saved on magneto-optic disk and reviewed offline in slow motion and single-frame advance mode for analysis. The diameter of the aortic ring was measured in a long-axis view by M-mode and used for cardiac output calculation. Left ventricular (LV) mass (LVM) was calculated from M-mode in accordance with the American Society of Echocardiography’s recommendations (35a). LV fractional shortening was computed from the standard formula (29).

Arterial blood pressure was measured with an automatic oscillometer cuff sphygmomanometer (Dynamap, Critikon, Tampa, FL). The RPP was calculated by multiplying heart rate with a mean systolic blood pressure (1). The analysis package of the ultrasound unit was used for RPP and LV external work divided by the cosine of the angle between the Doppler beam and the flow direction. True velocity was defined as the measured velocity divided by the vessel lumen.

Table 1. Hemodynamic and echocardiographic data before and 5 ± 1 days after surgery with CPB in patients with shunt and controls

<table>
<thead>
<tr>
<th></th>
<th>Controls (n = 19)</th>
<th>Presurgery (n = 18)</th>
<th>Post Surgery (n = 18)</th>
<th>P</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Age (range), mo</strong></td>
<td>6.5 (3–16)</td>
<td>6 (3–18)</td>
<td>6 (3–18)</td>
<td></td>
</tr>
<tr>
<td><strong>Systolic BP, mmHg</strong></td>
<td>89 (7)</td>
<td>95 (10.0)</td>
<td>98 (5.0)</td>
<td>0.46</td>
</tr>
<tr>
<td><strong>Diastolic BP, mmHg</strong></td>
<td>50 (6)</td>
<td>53 (8.0)</td>
<td>54 (8.0)</td>
<td>0.9</td>
</tr>
<tr>
<td><strong>HR, beats/min</strong></td>
<td>132 (10)</td>
<td>139 (15)</td>
<td>129 (17)</td>
<td>0.02</td>
</tr>
<tr>
<td><strong>RPP, mmHg/min</strong></td>
<td>11,940 (1,690)</td>
<td>13,278 (1,127)</td>
<td>12,672 (1,050)</td>
<td>0.25</td>
</tr>
<tr>
<td><strong>CO, ml·min⁻¹·kg⁻¹</strong></td>
<td>446 (133)</td>
<td>422 (222)</td>
<td>423 (191.3)</td>
<td>0.9</td>
</tr>
<tr>
<td><strong>FS, %</strong></td>
<td>38 (4)</td>
<td>46 (9)</td>
<td>42 (9)</td>
<td>0.6</td>
</tr>
<tr>
<td><strong>LVM, g</strong></td>
<td>21 (5.6)</td>
<td>25 (10)</td>
<td>25 (10)</td>
<td>0.9</td>
</tr>
<tr>
<td><strong>Aortic cross clamp time, min</strong></td>
<td>20.2 (6.0)</td>
<td>21.8 (5.1)</td>
<td>21.6 (5.0)</td>
<td>0.58</td>
</tr>
<tr>
<td><strong>CPB duration, min</strong></td>
<td>98.8 (30.6)</td>
<td>58.9 (23.5)</td>
<td>58.9 (23.5)</td>
<td></td>
</tr>
<tr>
<td><strong>Hb, g%</strong></td>
<td>12 (1)</td>
<td>13.1 (0.9)</td>
<td>13.1 (0.9)</td>
<td>0.0001</td>
</tr>
<tr>
<td><strong>LAD diameter, mm</strong></td>
<td>1.5 (0.2)</td>
<td>1.7 (0.3)</td>
<td>2.1 (0.4)</td>
<td>0.001</td>
</tr>
<tr>
<td><strong>LAD VTId+s, cm</strong></td>
<td>10 (3)</td>
<td>9 (3.0)</td>
<td>14 (5.0)</td>
<td>0.05</td>
</tr>
<tr>
<td><strong>LAD PFVd, cm/s</strong></td>
<td>37 (5)</td>
<td>36 (10)</td>
<td>49 (16)</td>
<td>0.05</td>
</tr>
<tr>
<td><strong>LAD BF, ml/min</strong></td>
<td>17 (5)</td>
<td>27 (10)</td>
<td>47 (15)</td>
<td>0.0001</td>
</tr>
</tbody>
</table>

Values are means (SD); n, no. of subjects. BP, blood pressure; HR, heart rate; RPP, rate pressure product; CO, cardiac output; FS, fractional shortening; LVM, left ventricular mass; CPB, cardiopulmonary bypass; LAD, left anterior descending artery; VTId+s, velocity time integral in diastole and systole; PFVd, diastolic peak flow velocity; BF, blood flow. P values are for presurgery vs. postsurgery.

**CPB and Cardioplegia**

Open-heart surgery was performed under mild to moderate hypothermic CPB (body temperature 28–32°C). For myocardial protection, cold (+4°C) hyperkalemic (K⁺ = 0.07 mmol/ml) blood cardioplegic solution was used. All patients had similar protocol for cardioplegia with anterograde perfusion of the coronary arteries to start with and then retrograde filling through the coronary sinuses every 20 min during the CPB.

**Postoperative Treatment**

All children after CPB surgery received opioid derivatives in the form of intravenous ketobemidone hydrochloride during the first 2–3 days and then followed by oxicodon orally between 3 and 5 days postoperatively. However, during the postoperative coronary flow studies, these medications had already been stopped. None of our patients was on angiotensin I-converting enzyme inhibitors, lanoxine, phenylepherine, beta-blockers, or any other vasoactive medications, such as nitrate or dopamine, during the intensive care management after day 1 postoperatively. Patients treated by cardiac catheterization were not on any of the above medications.

**Intraobserver Variability**

In 10 children, two registrations of LAD flow velocities were performed 15 min apart by the same observer (EHA). The paired data were analyzed regarding peak flow diastolic and systolic velocities, VTI, and LAD blood flow. The analyses of the Doppler tracings were performed offline.

**Statistics**

Paired Student’s t-test was used for comparison between patient’s data before and after surgery. Stepwise multiple regression analysis was used to compare CF with LVM, RPP, and LV fractional shortening. Simple regression analysis was used to calculate the correlation of the changes of CF with diastolic blood pressure changes. All statistical analyses were performed using StatView (SAS Institute 5.0) as a statistical software package. A P value of <0.05 was considered statistically significant. Results are presented as means ± SD. The intraobserver variability was statistically measured according to the British Standards Institution (2).
RESULTS

Infants With Ventricular Left to Right Shunt

Before surgery. Measurable coronary flow signals in LAD were obtained in 18 of 19 patients. The mean diameter of LAD in patients was 1.7 ± 0.3 mm, and in their age-matched healthy controls 1.5 ± 0.2 mm. VTIs+d was 9 ± 3 and 10 ± 3 cm and PFVd was 36 ± 10 and 37 ± 5 cm/s in patients and controls, respectively. Mean CF in LAD was 27 ± 10 ml/min in patients and 17 ± 5 ml/min in controls.

After CPB surgery. Mean LVM, LV fractional shortening, cardiac output, RPP, and systolic and diastolic blood pressure were similar before and after surgery. Heart rate was significantly higher before surgery (139 vs. 129 beats/min, \( P = 0.02 \), Table 1). CF did not correlate to LVM before or after surgery. It correlated to RPP after surgery (\( r = 0.5, P < 0.04 \)), but not before surgery (\( r = 0.1, P = 0.7 \), Fig. 1). The mean (range) aortic cross clamp time was 59 (33–121) min, and CPB time was 98 (64–193) min. There was no correlation between the change in CF before and after surgery and the duration of the CPB. In stepwise multiple-regression analyses, there were no correlations between CF and RPP, LVM, and LV fractional shortening after surgery, and neither was there before surgery.

Myocardial oxygen consumption increased after surgery from 65 ± 23 to 71 ± 27 ml O\(_2\)·min\(^{-1}\)·100 g\(^{-1}\) (\( P < 0.02 \)). There was no correlation between VTId+s or CF and myocardial oxygen consumption. The mean LAD diameter increased from 1.7 ± 0.3 to 2.1 ± 0.4 mm (\( P = 0.001 \)) with no correlation to CF. LAD VTId+d increased from 9 ± 3 to 14 ± 5 cm (\( P = 0.05 \)), and PFVd from 36 ± 10 to 49 ± 16 cm/s (\( P < 0.05 \)). A transthoracic Doppler signal of coronary flow velocity in LAD after surgery with CPB is presented in Fig. 2. The CF increased from 27 ± 10 to 47 ± 15 ml/min (\( P = 0.0001 \), Fig. 3). Hemoglobin concentration was higher 5 ± 1 days postoperatively (12 ± 1 vs. 13 ± 0.9 g/100 ml, \( P < 0.0001 \)).

Neonates With Coarctation of Aorta

Before surgery. Coronary flow recording in LAD was successful in all patients with coarctation of aorta. The LAD mean diameter in patients was 1.8 ± 0.1 mm, and in their age-matched controls it was 1.2 ± 0.1 mm. VTIs+d was 11 ± 4 and 6 ± 2 cm and PFVd was 42 ± 14 and 26 ± 8 cm/s in patients and controls, respectively. The LAD mean CF in patients was 44 ± 12 ml/min, and in age-matched neonates 8 ± 4 ml/min. There was a significant correlation (\( r = 0.76, P = 0.004 \)) between CF and LVM before, but not after, surgery.

After off-pump surgery. Mean LVM, LV fractional shortening, cardiac output, and heart rate before and after surgery were similar. The mean systolic and diastolic blood pressures were significantly lower after surgery: 102 and 80 mmHg, \( P = 0.0003 \), and 55 and 42 mmHg, \( P = 0.005 \), respectively. RPP was higher before than after surgery (\( P = 0.06 \) (Table 2). Before surgery, there was a weak correlation between CF and RPP (\( r = 0.53, P = 0.1 \)), but not at all at 5 days postoperatively (\( r = 0.025, P = 0.9 \)). In stepwise multiple regressions, there were significant correlations between CF and RPP, LVM, and fractional shortening before (\( r = 0.79, P = 0.01 \) for all), but these disappeared after surgery.

Myocardial oxygen consumption decreased after surgery from 18 ± 4 to 15 ± 3 ml O\(_2\)·min\(^{-1}\)·100 g\(^{-1}\), \( P < 0.001 \). There was no correlation between CF and VTId+s and myo-
cardiac oxygen consumption. LAD mean diameter was 1.8 ± 0.1 mm before surgery and 1.7 ± 0.1 mm after surgery (P = 0.06). LAD VTIs+d decreased from 11 ± 4 cm to 8 ± 2 cm (P < 0.05), PFVd from 42 ± 14 to 30 ± 10 cm/s (P = 0.008), and LAD CF from 44 ± 12 to 25 ± 8 ml/min (P = 0.001, Fig. 4).

Intraobserver Variability

The following coefficients of variation (COV) were obtained: 4.6% for LAD’s diameter, 11.7% for PFVd, 7% for PFVs, 9% for VTIs and VTId, and 3.3% for LAD blood flow. The COV for heart rate was 7%.

DISCUSSION

To our knowledge, this study is the first to study, by aid of TTDE, the CF in relation to CPB surgery. Our study shows that CF in LAD in children with left-to-right shunt is increased for at least 5 days after CPB surgery. The discrepancy between the relatively small increase in myocardial oxygen consumption (from 65 to 71 ml O2/min) and the 2.5-fold increase in coronary flow suggests that other factors are mainly responsible for the postsurgical rise in flow after CPB surgery. Because there is a certain maximal dilatory state that can be reached, it is conceivable to assume that an increase in CF may lead to impaired CFR (38). Therefore, our findings support earlier studies by positron emission tomography (PET) and intracoronary Doppler guide wire (IDGW) showing a drop in CFR after CPB surgery (7, 13). In contrast, surgical off-pump correction of aortic coarctation resulted in decrease of both myocardial oxygen consumption and coronary flow, which implies either that off-pump surgery has little or no effect on the coronary flow or that the removal of pressure overload after coarctectomy counteracts possible detrimental effects of off-pump surgery on coronary flow. Importantly, the intraobserver COV for flow velocities and also for flow volume were fairly low, being comparable with earlier reports (2, 31).

TTDE is noninvasive, safe, widely available, and inexpensive. TTDE assessment of CF and flow velocity has been shown to correlate well with IDGW, PET, and coronary angiography measurements (35, 19). The changes in the diameter of coronary arteries affect CF dramatically, because the area of the coronary artery lumen is related to the artery’s radius squared. In the present study, the employed ultrasound probes (7 and 10 MHz) have an axial resolution of 0.1 mm, thus rendering the comparison in diameter measurements accurate. Measurements of coronary artery diameter by transthoracic echocardiography correlate well with measurements with quantitative coronary angiography (24, 27).

Table 2. Hemodynamic and echocardiographic data in patients with coarctation of aorta before and 5 ± 1 days after non-CPB surgery and in controls

<table>
<thead>
<tr>
<th></th>
<th>Controls* (n = 55)</th>
<th></th>
<th>Patients (n = 12)</th>
<th></th>
<th></th>
<th></th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td></td>
<td>Pre surgery</td>
<td>Post surgery</td>
<td>P</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Age (range), days</td>
<td>5 (0.5–30)</td>
<td>10 (2–50)</td>
<td>10 (2–50)</td>
<td>0.0003</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Systolic BP, mmHg</td>
<td>79 (12)</td>
<td>102 (11)</td>
<td>80 (7.0)</td>
<td>0.005</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Diastolic BP, mmHg</td>
<td>43 (10)</td>
<td>55 (8.0)</td>
<td>42 (4.0)</td>
<td>0.005</td>
<td></td>
<td></td>
</tr>
<tr>
<td>RPP, mmHg/min</td>
<td>9,724 (2,042)</td>
<td>14,302 (1,310)</td>
<td>12,045 (1,298)</td>
<td>0.06</td>
<td></td>
<td></td>
</tr>
<tr>
<td>HR, beats/min</td>
<td>123 (17)</td>
<td>148 (6.5)</td>
<td>147 (10.6)</td>
<td>0.8</td>
<td></td>
<td></td>
</tr>
<tr>
<td>CO, ml·min⁻¹·kg⁻¹</td>
<td>39 (5.1)</td>
<td>47 (10.2)</td>
<td>46 (9.5)</td>
<td>0.6</td>
<td></td>
<td></td>
</tr>
<tr>
<td>FS, %</td>
<td>12 (3)</td>
<td>14 (5.5)</td>
<td>14 (5.5)</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>LVM, g</td>
<td>1.2 (0.1)</td>
<td>1.8 (0.1)</td>
<td>1.7 (0.1)</td>
<td>0.06</td>
<td></td>
<td></td>
</tr>
<tr>
<td>LAD diameter, mm</td>
<td>6 (2.0)</td>
<td>11 (4)</td>
<td>8 (2)</td>
<td>0.05</td>
<td></td>
<td></td>
</tr>
<tr>
<td>LAD VTId+s, cm</td>
<td>26 (8.0)</td>
<td>42 (14)</td>
<td>30 (10)</td>
<td>0.008</td>
<td></td>
<td></td>
</tr>
<tr>
<td>LAD BF, ml/min</td>
<td>8 (4.0)</td>
<td>44 (12)</td>
<td>25 ± 8</td>
<td>0.001</td>
<td></td>
<td></td>
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</table>

Values are means (SD); n, no. of subjects. P values are for presurgery vs. postsurgery. *From Ref. 30.

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In healthy children (4–53 mo of age), there is a linear correlation of LAD’s CF and PFVd with LVM (30). During the first month of life, the PFVd, VTIs+t, and CF in LAD increase. These changes are related to the rapid body growth during this period of life. The physiological improvement in LV relaxation and compliance that occur during the first month of life could be an explanatory factor for the increase in coronary flow parameters with age (25). In our study, flow measurements in the LAD in age-matched normal controls were used as normal values for coronary flow (31).

The hypertrophic myocardium due to coarctation implies increased oxygen demand and blood flow. After coarctectomy, with subsequent decrease in cardiac pressure afterload, the LAD diameter, VTId+s, PFVd, and CF all decreased, but they were still higher on day 5 than in control neonates. The increase in CF before surgery in coarctation patients is related to coarctation-induced elevation in blood pressure, which leads to high myocardial oxygen demand and consequent increase of coronary flow. The CF decreases after coarctectomy, as expected, because of diminished cardiac pressure afterload due to decreased myocardial oxygen demand.

The postoperative decrease of CF along with the normalization of myocardial oxygen demand per unit of myocardial tissue result in increase of CFR in these patients (39).

During the immediate postoperative period, there are factors affecting coronary flow, such as an inflammatory response. Inflammation is associated with several factors with adverse influence on arterial endothelial and smooth muscle cells. Complement system is activated, proinflammatory cytokines are released, and oxidative stress and disturbances in calcium homeostasis lead to ischemia-reperfusion (36). Christen and colleagues (3) suggested that hemolysis was the main cause of early oxidative stress. TNF-α upregulates inducible nitric oxide (NO) synthase activity with secondary increased release of NO, thereby contributing to the reduction in the intrinsic tone of coronary microcirculation (26). Intense upregulation of inducible NO synthase leads to high levels of intravascular NO, which can damage endothelial cells through its oxidative potential (36). Prolonged ischemia-reperfusion causes an increase in oxygen-derived free radicals such as superoxide anions. Superoxide reacts with NO, forming peroxynitrite, which is highly damaging for endothelial cells (11, 34). CPB with cardioplectic arrest results in vasomotor dysfunction (23, 37, 38).

Previous studies by PET in neonates and infants operated with CPB have shown low CFR values (7, 42). As suggested by the present study, one possible explanation could reside in the increased basal coronary flow at 5 days after CPB surgery. In controls, CBW was 17 ml/min, assuming a normal CFR of ~4, the maximal obtainable coronary flow in the patients would be 68 ml/min, and we measured 47 ml/min. Our findings suggest that use of CPB during cardiac surgery and the type of congenital heart disease are important factors affecting the postoperative resting CF in children. It has been reported that, in patients with transposition of great arteries after arterial switch operation, the coronary flow as assessed by IDGW is increased and CFR is decreased, even 5 yr after surgery (9). This could be due to surgical technique per se, but eventual long-lasting effects of CPB surgery cannot be ruled out.

**Study Limitations**

Intravenous adenosine infusion to measure CFR could not be done, because the consents from the parents were not obtained. Longer postoperative follow up would have been indicated, but this could not be done, as the patients were discharged by day 7–8 after surgery.

In conclusion, coronary flow is increased after CPB surgery, a finding that could explain, in part, the previously reported decrease of CFR. Off-pump surgery as such, at least in children with aortic coarctation, seems to have little or no impact on coronary flow as the postsurgical decline in flow in this category of patients relates to the reduction in cardiac pressure afterload. TTDE renders possible serial coronary flow studies in children after cardiac surgery.

**ACKNOWLEDGMENTS**

We thank Annica Maxedius, registered research nurse, for assistance in patient recruitment, and Dr. Markku Saraste for technical advice in the registrations.

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