End-diastolic flow reversal limits the efficacy of pediatric intra-aortic balloon pump counterpulsation

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Background: Counterpulsation with an intra-aortic balloon pump (IABP) has not achieved the same success or clinical use in pediatric patients as in adults. In a pediatric animal model, IABP efficacy was investigated to determine whether IABP timing with a high-fidelity blood pressure signal may improve counterpulsation therapy versus a low-fidelity signal.

Methods: In Yorkshire piglets (n = 19; weight, 13.0 ± 0.5 kg) with coronary ligation–induced acute ischemic left ventricular failure, pediatric IABPs (5 or 7 mL) were placed in the descending thoracic aorta. Inflation and deflation were timed with traditional criteria from low-fidelity (fluid-filled) and high-fidelity (micromanometer) blood pressure signals during 1:1 support. Aortic, carotid, and coronary hemodynamics were measured with pressure and flow transducers. Myocardial oxygen consumption was calculated from coronary sinus and arterial blood samples. Left ventricular myocardial blood flow and end-organ blood flow were measured with microspheres.

Results: Despite significant suprasystolic diastolic augmentation and afterload reduction at heart rates of 105 ± 3 beats per minute, left ventricular myocardial blood flow, myocardial oxygen consumption, the myocardial oxygen supply/demand relationship, cardiac output, and end-organ blood flow did not change. Statistically significant end-diastolic coronary, carotid, and aortic flow reversal occurred with IABP deflation. Inflation and deflation timed with a high-fidelity versus low-fidelity signal did not attenuate systemic flow reversal or improve the myocardial oxygen supply/demand relationship.

Conclusions: Systemic end-diastolic flow reversal limited counterpulsation efficacy in a pediatric model of acute left ventricular failure. Adjustment of IABP inflation and deflation timing with traditional criteria and a high-fidelity blood pressure waveform did not improve IABP efficacy or attenuate flow reversal. End-diastolic flow reversal may limit the efficacy of IABP counterpulsation therapy in pediatric patients with traditional timing criteria. Investigation of alternative deflation timing strategies is warranted. (J Thorac Cardiovasc Surg 2014;147:1660-7)

Counterpulsation with an intra-aortic balloon pump (IABP) is the most common mechanical circulatory support strategy for a variety of cardiac pathologies in adults. Yet, IABP therapy in pediatric patients has not demonstrated the same degree of efficacy and has not gained widespread clinical use. Differences between adult and pediatric anatomy and physiology may limit the efficacy of IABP therapy in neonates, infants, and children.

In this study, we examined IABP counterpulsation in a pediatric model of acute ischemic left ventricular failure. We tested the hypothesis that the efficacy of pediatric IABP therapy is improved with a high-fidelity (micromanometer) rather than a traditional low-fidelity (fluid-filled) arterial blood pressure signal used to adjust IABP inflation and deflation timing.

METHODS

All animals received humane care and were handled in accordance with the Guide for the Care and Use of Laboratory Animals (National Research Council, 1996). Experimental procedures followed animal study protocols approved by the University of Louisville (Louisville, Ky) Institutional Animal Care and Use Committee.

Experimental Design

Piglets (n = 19; weight, 13.0 ± 0.5 kg) were instrumented surgically to determine aortic, carotid, and coronary artery hemodynamics. Myocardial oxygen consumption (MVO2) was calculated from coronary sinus and arterial blood gas data. Left ventricular myocardial blood flow and regional
end-organ blood flow were determined with neutron-activated 15-μm microspheres. Sequential coronary ligation was performed to induce acute ischemic left ventricular failure. A pediatric IABP was placed in the descending thoracic aorta.

In each animal, hemodynamic waveforms (15-second data epochs), blood gases, and end-organ blood flows were measured at steady state during the following experimental test conditions: (1) normal baseline (IABP off), (2) coronary ligation-induced left ventricular failure (IABP off), (3) IABP support with timing adjusted to the arterial blood pressure waveform acquired with a low-fidelity (<20-Hz response) fluid-filled catheter in the radial artery, and (4) IABP support with timing adjusted to the arterial pressure waveform acquired with a high-fidelity (<5-kHz response) micromanometer pressure transducer in the aortic root. Measurements were recorded during 1:1 counterpulsation support in which each aortic valve closure initiated rapid balloon inflation and each end diastole initiated rapid balloon deflation.

Of note, an aortic root pressure waveform transduced by a high-fidelity catheter results in a more precise waveform morphology with a more pronounced dicrotic notch, less signal gain, and less phase distortion. Consequently, a high-fidelity signal was presumed to enable more accurate inflation and deflation timing of the IABP.

Surgical Preparation

Animals were fasted, preanesthetized with intramuscular ketamine (30 mg/kg) and acepromazine (0.2 mg/kg), and anesthetized with isoflurane (1.5%-3%) and room air. A left lateral thoracotomy was performed, and the left and right carotid artery and jugular vein were exposed through neck incisions. A single-tip high-fidelity micromanometer catheter (2.5 Fr, SPR-524; Millar Instruments, Houston, Tex) was placed in the left atrium. A dual-tip high-fidelity micromanometer catheter (5 Fr, SPC-721; Millar Instruments) was advanced retrograde from the ascending aorta into the left ventricle. Transit-time ultrasonic flow probes (T205; Transonic, Ithaca, NY) were placed around the aortic root, left carotid artery, and left anterior descending (LAD) coronary artery to measure volumetric blood flows. Sampling catheters were introduced into the coronary sinus via the hemiazygous vein, pulmonary artery, and right carotid artery to sample blood and measure blood gases. An infusion catheter was introduced into the left atrium for serial injections of microspheres during each experimental test condition to determine end-organ blood flows throughout the body. A blood-sampling catheter was introduced into the right femoral artery for simultaneous withdrawal of a microsphere reference blood-flow sample during each test condition. A fluid-filled pressure-monitoring catheter (24-gauge Angiocath; Becton Dickinson, Sandy, Utah) was inserted into the right radial artery.

The animal was anticoagulated with a bolus injection of heparin (300 U/kg) and subsequent small boluses of heparin to maintain an activated clotting time of greater than 300 seconds. An IABP catheter (5 or 7 mL;Datascope, Fairfield, NJ) was inserted via a left femoral artery cut down and advanced until the tip was distal to the left subclavian artery, as confirmed by digital palpation. Balloon timing was manually adjusted with the arterial pressure waveform (fluid-filled radial artery catheter or aortic root Millar catheter) displayed on the IABP console (System 97;Datascope). Balloon inflation and deflation were timed to maximize aortic diastolic pressure augmentation and minimize aortic end-diastolic pressure (afterload).

Induction of Ischemic Left Ventricular Failure

Intravenous lidocaine (20-mg bolus, 20-mg/h infusion) and esmolol (5-mg/kg bolus, 50-μg/kg per minute infusion) were administered to prevent arrhythmia. Sequential coronary ligation of branches of the LAD induced acute ischemic left ventricular failure. Target cardiac dysfunction was achieved when 3 of 4 criteria were met: approximate reduction of (1) left ventricular cardiac output by 25%, (2) mean aortic pressure by 10 mm Hg, (3) mixed venous O2 saturation by 10%, and (4) an elevation of left atrial pressure (LAP) and/or left ventricular end-diastolic pressure by 5 mm Hg. Animals with life-threatening hypotension were supported with boluses of intravenous normal saline and continuous infusion of phenylephrine and/or epinephrine to effect.

Blood Gas Analysis

During each experimental test condition, blood samples were simultaneously withdrawn from the coronary sinus, pulmonary artery, and right carotid artery. Samples were processed with a blood gas analyzer (IRMAQ, Blood Analysis System; Diametrics Medical, Roseville, Minn) to measure hemoglobin content (\(\text{Hb} \text{g/dL}\)) and hemoglobin oxygen saturation (\(\% \text{O}_2 \text{Sat}\)). Oxygen content (\(\text{O}_2 \text{c}\)) was calculated for each blood sample:

\[
\text{O}_2 material per gram for each sample during each experimental test condition. Tissue and reference blood samples were sent to BioPAL, Inc (Worcester, Mass) for analysis.

End-Organ Blood Flow Measurements

During each experimental test condition, a different color (isotope label) of 15-μm neutron-activated microspheres (1 × 10⁶ microspheres in 0.4-mL suspension; Biopal, Worcester, Mass) was injected into the left atrium, followed by a 4-mL saline flush. The microsphere technique enabled the measurement of regional end-organ blood flow in vascular beds of interest, as previously described. During microsphere injection, a reference blood-flow sample was drawn from the femoral artery at a rate of 4 mL/min for 60 seconds with a calibrated syringe pump (Harvard Apparatus, Holliston, Mass). The withdrawal sample acted as a reference to determine organ-specific flows in mL/min per gram of tissue.

Necropsy and Microsphere Analysis

After completion of the experimental protocol, animals were euthanized with an increase in anesthetic depth and an intravenous bolus injection of supersaturated KCl. After euthanasia, the heart was harvested and weighed. The ventricles were dissected from the atria. The left ventricle was dissected from the right ventricle. The brain, lungs, kidneys, pancreas, liver, spleen, and adrenals were harvested and weighed.

Blood and reference blood samples were sent to BioPAL, Inc (Worcester, Mass) for radioactive assay and automated calculation of blood flow in mL/min per gram for each sample during each experimental test condition. Tissue and blood samples were bombarded with neutrons to transiently activate each

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**Abbreviations and Acronyms**

<table>
<thead>
<tr>
<th>Abbreviation</th>
<th>Description</th>
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<tbody>
<tr>
<td>AoF</td>
<td>aortic flow</td>
</tr>
<tr>
<td>AoP</td>
<td>aortic pressure</td>
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<tr>
<td>bpm</td>
<td>beats per minute</td>
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<tr>
<td>IABP</td>
<td>intra-aortic balloon pump</td>
</tr>
<tr>
<td>LAD</td>
<td>left anterior descending</td>
</tr>
<tr>
<td>LADF</td>
<td>LAD flow</td>
</tr>
<tr>
<td>LAP</td>
<td>left atrial pressure</td>
</tr>
<tr>
<td>LVP</td>
<td>left ventricular pressure</td>
</tr>
<tr>
<td>MVO₂</td>
<td>myocardial oxygen consumption</td>
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group of microspheres with a separate isotope label. The level of radioactivity detected in the sample was directly proportional to the number of microspheres present in the sample. The activity of microspheres in the reference blood sample was compared with the activity of microspheres that lodged in a tissue sample of interest. The ratio between the 2 activity counts was equal to the ratio between the calibrated rate of aortic withdrawal (known, 4 mL/min) and flow in the tissue of interest (unknown).

**Instrumentation, Data Reduction, and Statistics**

All transducers were precalibrated and postcalibrated against known physical standards to ensure measurement accuracy. Collected data were signal conditioned (1000× gain and 60-Hz low-pass filter) and analog-to-digital converted at a sampling rate of 400 Hz for digital analysis using our Good Laboratory Practices–compliant data acquisition system.6

Hemodynamic parameters were calculated on a beat-to-beat basis for each 15-second data set with the Hemodynamic Evaluation and Assessment Research Tool program7 developed in Matlab, version 6.5 (MathWorks, Natick, Mass). All analyzed beats in each data set (approximately 25-30 beats/15-second data set) were averaged to obtain a single representative mean value for each calculated parameter.

Pressure and flow waveforms were used to derive the following hemodynamic parameters: heart rate (HR), aortic pressure (AoP), left atrial pressure (LAP), left ventricular pressure (LVP), aortic flow (AoF), carotid artery flow, and LAD flow (LADF). Left ventricular myocardial blood flow, as

<table>
<thead>
<tr>
<th>Variable</th>
<th>HR, bpm</th>
<th>LADF\text{mean}, mL/min</th>
<th>LVCO, L/min</th>
<th>AoP\text{mean}, mm Hg</th>
<th>LAP\text{mean}, mm Hg</th>
<th>LVP\text{end-diastolic}, mm Hg</th>
<th>L kidney, mL/min per gram</th>
<th>Mixed venous O₂, %</th>
</tr>
</thead>
<tbody>
<tr>
<td>Baseline</td>
<td>107 ± 4</td>
<td>26 ± 4</td>
<td>2.30 ± 0.17</td>
<td>67 ± 4</td>
<td>9 ± 1</td>
<td>9 ± 1</td>
<td>1.69 ± 0.29</td>
<td>86 ± 1</td>
</tr>
<tr>
<td>LV failure</td>
<td>104 ± 3</td>
<td>16 ± 3*</td>
<td>1.67 ± 0.15</td>
<td>53 ± 3*</td>
<td>12 ± 1*</td>
<td>14 ± 1*</td>
<td>0.90 ± 0.13*</td>
<td>70 ± 4*</td>
</tr>
<tr>
<td>1:1 low-fidelity IABP</td>
<td>105 ± 3</td>
<td>18 ± 3*</td>
<td>1.78 ± 0.15</td>
<td>56 ± 3</td>
<td>11 ± 1</td>
<td>14 ± 1</td>
<td>0.99 ± 0.22</td>
<td>74 ± 4</td>
</tr>
<tr>
<td>1:1 high-fidelity IABP</td>
<td>107 ± 3</td>
<td>19 ± 3*</td>
<td>1.84 ± 0.15</td>
<td>56 ± 3</td>
<td>11 ± 1</td>
<td>12 ± 2</td>
<td>1.01 ± 0.15</td>
<td>70 ± 5</td>
</tr>
</tbody>
</table>

**FIGURE 1.** A, Aortic root pressure waveforms from the same animal demonstrated suprasystolic diastolic augmentation and afterload reduction during 1:2 (middle panel) and 1:1 (bottom panel) intra-aortic balloon pump (IABP) support compared with baseline (top panel). Arrows indicate supported beats. B, Diastolic augmentation of greater than 20 mm Hg was achieved when the IABP was timed with a low-fidelity (fluid-filled) catheter in the radial artery or a high-fidelity (micromanometer) catheter in the aorta. AoP, Aortic pressure; LV, left ventricular. *P < .0001 versus baseline; †P < .0001 versus LV failure.
determined by microspheres normalized to MVO$_2$ was used as an index of the oxygen supply/demand relationship. Prism, version 4.00 (GraphPad, La Jolla, Calif), was used to perform statistical analyses and to plot data. To verify the induction of a clinically relevant state of left ventricular failure, paired Student $t$ tests were used to compare values for LADF, AoF, AoP, mixed venous O$_2$ saturation, end-organ blood flows, LAP, and LVP at baseline and after coronary ligation. A 1-way repeated-measures analysis of variance with a Tukey posttest for comparison of means was performed to compare left ventricular failure and IABP timing modes (low and high fidelity) for each parameter. $P < .05$ (95% confidence) was considered statistically significant. All data were presented as mean $\pm$ SE.

**RESULTS**

**Model of Pediatric Left Ventricular Failure**

Of the 19 piglets, 10 completed the experimental protocol. In these animals, LAD ligation produced ischemic left ventricular failure with phenotypic similarities to clinical left ventricular failure (Table 1). Animals exhibited an approximate reduction in LADF$_{\text{mean}}$ by 40% (26 $\pm$ 4 to 16 $\pm$ 3 mL/min, $P < .01$), AoF by 30% (2.4 $\pm$ 0.2 to 1.7 $\pm$ 0.2 L/min, $P < .001$), AoP$_{\text{mean}}$ by 15 mm Hg (67 $\pm$ 4 to 53 $\pm$ 3 mm Hg, $P < .01$), mixed venous O$_2$ saturation by 15% (86% $\pm$ 1% to 70% $\pm$ 4%, $P < .001$), renal blood flow by 45% to 50% (left kidney: 1.69 $\pm$ 0.29 to 0.90 $\pm$ 0.13 mL/min per 100 g, $P < .05$; right kidney: 2.13 $\pm$ 0.45 to 1.02 $\pm$ 0.18 mL/min per 100 g, $P < .05$), an increased LAP by approximately 3 mm Hg (9 $\pm$ 1 to 12 $\pm$ 1, $P < .001$), and LVP$_{\text{end-diastolic}}$ by approximately 5 mm Hg (9 $\pm$ 1 to 14 $\pm$ 1, $P < .01$).

The other 9 animals died before completion of the study protocol, typically from intractable arrhythmia. Data from these animals were not included in any of the analyses.

**Diastolic Augmentation and Left Ventricular Afterload Reduction**

At heart rates of 105 $\pm$ 3 beats per minute (bpm; range, 83-125 bpm), it was possible to achieve statistically significant suprasystolic diastolic pressure augmentation during each native cardiac beat (Figure 1, A, bottom waveform). During 1:1 IABP support timed with a low-fidelity (fluid-filled) catheter in the radial artery or a high-fidelity (micromanometer) catheter in the aorta. B, Left ventricular myocardial blood flow (LVF) did not improve significantly during 1:1 IABP therapy timed with either a low-fidelity or a high-fidelity blood pressure signal. C, Myocardial oxygen consumption (MVO$_2$) did not improve during 1:1 IABP therapy timed with either a low-fidelity or a high-fidelity blood pressure signal. D, LVF normalized to MVO$_2$, an index of the myocardial oxygen supply/demand relationship, did not significantly improve during 1:1 IABP therapy timed with either a low-fidelity or a high-fidelity blood pressure signal. LV, Left ventricular. *$P < .01$ versus baseline; †$P < .05$ versus LV failure.
During 1:1 IABP support, aortic end-diastolic pressure decreased by approximately 8 mm Hg (left ventricular failure, 41 ± 2 mm Hg; low-fidelity signal, 34 ± 3 mm Hg [P < .0001]; high-fidelity signal, 33 ± 3 mm Hg [P < .0001]). The increase in left ventricular myocardial blood flow during IABP support was small and did not reach statistical significance with either a low- or high-fidelity blood pressure signal (Figure 2, B). Similarly, myocardial oxygen consumption did not change during 1:1 IABP support with either a low- or high-fidelity blood pressure signal (Figure 2, C). As a result, despite suprasystolic diastolic augmentation and left ventricular afterload reduction, the myocardial oxygen supply/demand relationship did not significantly improve with either a low-fidelity or a high-fidelity signal to adjust IABP inflation and deflation timing (Figure 2, D).

### Table 2. End-organ blood flow in a pediatric model of left ventricular failure during IABP support

<table>
<thead>
<tr>
<th>ANOVA</th>
<th>L cerebrum, mL/min per gram</th>
<th>R cerebrum, mL/min per gram</th>
<th>Cerebellum, mL/min per gram</th>
<th>Brainstem, mL/min per gram</th>
<th>L lung, mL/min per gram</th>
<th>R lung, mL/min per gram</th>
</tr>
</thead>
<tbody>
<tr>
<td>P value</td>
<td>Baseline</td>
<td>0.31 ± 0.08</td>
<td>0.28 ± 0.07</td>
<td>0.62 ± 0.08</td>
<td>0.42 ± 0.10</td>
<td>1.99 ± 0.48</td>
</tr>
<tr>
<td></td>
<td>Left ventricular failure</td>
<td>0.26 ± 0.05</td>
<td>0.24 ± 0.04</td>
<td>0.35 ± 0.03*</td>
<td>0.39 ± 0.08</td>
<td>0.96 ± 0.20*</td>
</tr>
<tr>
<td></td>
<td>1:1 low-fidelity IABP</td>
<td>0.36 ± 0.09</td>
<td>0.35 ± 0.07</td>
<td>0.38 ± 0.06</td>
<td>0.28 ± 0.08</td>
<td>1.14 ± 0.27</td>
</tr>
<tr>
<td></td>
<td>1:1 high-fidelity IABP</td>
<td>0.33 ± 0.07</td>
<td>0.30 ± 0.06</td>
<td>0.43 ± 0.06</td>
<td>0.38 ± 0.11</td>
<td>1.16 ± 0.24</td>
</tr>
</tbody>
</table>

ANOVA, Analysis of variance; L, left; R, right; IABP, intra-aortic balloon pump. *P < .05 versus baseline.
min \( [P < .0001] \). Mean flow reversal in the LAD increased by approximately 0.70 mL/min (left ventricular failure, \(-0.24 \pm 0.06\) mL/min; low-fidelity signal, \(-1.01 \pm 0.24\) mL/min \( [P < .0001] \); high-fidelity signal, \(-0.91 \pm 0.18\) mL/min \( [P < .0001] \)).

End-Organ Blood Flow

Cardiac output (LVO) did not significantly improve during 1:1 IABP support (left ventricular failure, \(1.67 \pm 0.15\) L/min; low-fidelity signal, \(1.78 \pm 0.15\) L/min; high-fidelity signal, \(1.84 \pm 0.15\) L/min). As a result, systemic end-organ blood flow did not significantly improve with 1:1 IABP support timed with a low- or a high-fidelity blood pressure signal (Table 2).

**DISCUSSION**

Pediatric patients with life-threatening heart failure present difficult clinical challenges. During the past 5 decades, the intra-aortic balloon pump (IABP) has successfully supported pediatric patients with congenital and acquired heart disease as a bridge to decision, cardiac transplantation, and recovery. Yet, mortality rates greater than 35%2,8 indicate a need to improve the efficacy of pediatric IABP therapy.

In this study, in a piglet model of acute left ventricular failure, 1:1 IABP counterpulsation was possible at high heart rates. However, brief systemic blood flow reversal coincident with IABP deflation may have limited the efficacy of IABP support. Despite significant suprasystolic diastolic augmentation and significant left ventricular afterload reduction during 1:1 IABP support, left ventricular myocardial blood flow, myocardial oxygen consumption, and the myocardial oxygen supply/demand relationship did not improve. Cardiac output and end-organ blood flow also did not significantly improve. A brief, but significant, flow reversal in conduit arteries may have limited systemic forward flow. Traditional IABP inflation and deflation timing with a high-fidelity trigger did not attenuate flow reversal or improve IABP efficacy. These findings suggest that end-diastolic flow reversal is an important and novel mechanism that may limit the efficacy of IABP counterpulsation in pediatric patients with traditional inflation and deflation criteria.

**Clinical Considerations**

The optimal timing of pediatric IABP inflation and deflation is uncertain.11,12 In any patient with an IABP, it is traditionally thought that inflation and deflation of the balloon must occur during diastole between closure of the aortic valve and the subsequent systole. In an adult patient, this timing paradigm typically is not difficult to achieve. However, pediatric patients have higher heart rates than adults (newborn, 120-160 bpm; infant/toddler, 90-130 bpm), and diastole is short. As heart rate increases, diastole becomes disproportionately shorter.13 As a result, inflation and deflation of the balloon must occur within a smaller window of time. From the current study, we speculate that the short duration of diastole during high heart rates may limit the hemodynamic benefits of IABP inflation. Correspondingly, rapid balloon deflation produces a sharp reduction in aortic end-diastolic blood pressure and brief flow reversal sufficient to limit net augmentation of coronary blood flow or improve the myocardial oxygen supply/demand relationship.

Indeed, in adult patients, balloon deflation may produce a steep reduction of central end-diastolic blood pressure that may lead to a “steal” phenomenon during which coronary, carotid,15,16 cerebral,17 and aortic15,18 blood flow reverse. In adults, flow reversal is brief, and net flow augmentation is still positive. However, in a pediatric patient with a high heart rate, forward flow from diastolic augmentation occurs over a shorter period, and rapid balloon deflation occurs more frequently. As a result, the ratio between flow reversal and augmented forward flow is greater and may limit maximum net forward flow.

Importantly, coronary flow reversal may be attenuated by adjusting inflation and deflation timing.14 In adults with a normal heart rate (60-80 bpm), sufficient time exists between each systole to allow a brief (up to 25-millisecond) deflation delay, which may reduce flow reversal. As a result, subtle variations in the timing of inflation and deflation may permit an incremental tradeoff between afterload and myocardial workload reduction versus diastolic augmentation and coronary perfusion. Unfortunately, in pediatric patients, high heart rates with a shorter diastole minimize the available window to delay deflation for this tradeoff.

Similarly, brief timing errors associated with a fluid-filled versus a high-fidelity signal12 or a high-fidelity signal versus

**TABLE 2. Continued**

<table>
<thead>
<tr>
<th>L kidney, mL/min</th>
<th>R kidney, mL/min</th>
<th>Pancreas, mL/min</th>
<th>Liver, mL/min</th>
<th>Spleen, mL/min</th>
<th>L adrenal, mL/min</th>
<th>R adrenal, mL/min</th>
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<tbody>
<tr>
<td>per gram</td>
<td>per gram</td>
<td>per gram</td>
<td>per gram</td>
<td>per gram</td>
<td>per gram</td>
<td>per gram</td>
</tr>
<tr>
<td>1.69 ± 0.31</td>
<td>2.13 ± 0.45</td>
<td>0.29 ± 0.08</td>
<td>0.52 ± 0.20</td>
<td>1.14 ± 0.22</td>
<td>0.88 ± 0.33</td>
<td>0.83 ± 0.35</td>
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<tr>
<td>0.90 ± 0.13*</td>
<td>1.02 ± 0.18*</td>
<td>0.22 ± 0.05</td>
<td>0.21 ± 0.06</td>
<td>0.41 ± 0.07*</td>
<td>0.41 ± 0.04</td>
<td>0.39 ± 0.08</td>
</tr>
<tr>
<td>0.99 ± 0.22</td>
<td>1.02 ± 0.20</td>
<td>0.20 ± 0.04</td>
<td>0.23 ± 0.05</td>
<td>0.50 ± 0.15</td>
<td>0.56 ± 0.11</td>
<td>0.52 ± 0.12</td>
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<tr>
<td>1.01 ± 0.15</td>
<td>1.20 ± 0.18</td>
<td>0.19 ± 0.04</td>
<td>0.36 ± 0.08</td>
<td>0.54 ± 0.09</td>
<td>0.62 ± 0.14</td>
<td>0.63 ± 0.13</td>
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echocardiographic timing\textsuperscript{19} may reduce counterpulsation efficacy in pediatric patients. A precise signal is required to accurately time inflation and deflation and maximize counterpulsation efficacy in pediatric patients. To this end, integration of a high-fidelity micromanometer into the catheter tip may eliminate timing error(s), especially compared with low-fidelity fluid-filled catheters, which introduce gain and phase distortion and tend to drift over time.\textsuperscript{20} Indeed, we\textsuperscript{19} observed that it was easier to identify inflation and deflation timing landmarks in the arterial pressure waveform acquired with a high-fidelity signal. Consequently, timing errors were unlikely responsible for end-diastolic flow reversal. In our study, adjustment of IABP timing with a high-fidelity blood pressure signal (that minimized timing error) did not significantly reduce end-diastolic flow reversal or improve hemodynamics or end-organ blood flow compared with a low-fidelity signal. This finding further supports the notion that balloon inflation and deflation with traditional timing criteria triggered by a high-fidelity arterial blood pressure signal are insufficient to improve counterpulsation therapy in pediatric patients. Instead, the development of alternative timing strategies are necessary to prevent end-diastolic flow reversal and improve IABP efficacy in pediatric patients.

Other Considerations

Counterpulsation therapy is most effective when the displaced blood volume of the balloon and the left ventricular stroke volume are similar.\textsuperscript{21,22} In this study, we used 5- and 7-mL IABPs, which are the clinically recommended sizes for an 8- to 18-kg infant.\textsuperscript{23} However, the stroke volume of a 12-kg piglet (and a 12-kg human) with left ventricular failure is approximately 15 to 17 mL, more than 3 times the displacement volume of these IABPs. This observation begs the question of whether a larger balloon (and a larger displaced blood volume) may produce greater diastolic pressure augmentation to significantly improve diastolic coronary blood flow. By the same token, rapid deflation of a balloon with a larger volume may produce a greater decrease in end-diastolic aortic blood pressure and further increase the volume of blood flow reversal. This hypothesis remains untested. However, from a technical standpoint, the size of peripheral arteries in pediatric patients may limit the size of the balloon catheter that may be safely inserted and preclude counterpulsation therapy with a larger-sized IABP.

Arterial compliance also plays an important role in counterpulsation therapy. It has been demonstrated in vitro,\textsuperscript{24} experimentally in animal studies,\textsuperscript{25} and clinically\textsuperscript{26} that the efficacy of counterpulsation therapy correlates inversely with arterial compliance. Specifically, stiffer arteries generate the greatest diastolic augmentation and reduction in end-diastolic blood pressure. As a result, it has been speculated that the high compliance of the pediatric aorta may limit IABP efficacy.\textsuperscript{27} This effect has not been rigorously defined. Moreover, there is evidence to the contrary that the compliance of the pediatric aorta\textsuperscript{28} is sufficient to produce diastolic augmentation, afterload reduction, and improved hemodynamics.\textsuperscript{27} Our results further support this point.

Future Investigations

Ongoing studies will determine whether alternative timing strategies, such as delayed balloon deflation, may improve the efficacy of pediatric counterpulsation. Similarly, the use of larger pediatric balloons with greater displacement volumes that are more comparable to native stroke volume may improve the efficacy of pediatric IABP counterpulsation.

CONCLUSIONS

An incomplete understanding of counterpulsation has limited the widespread application of IABPs in pediatric patients. In a piglet model of acute ischemic left ventricular failure, significant diastolic augmentation and afterload reduction were achieved with a pediatric IABP at high heart rates. However, significant aortic, carotid, and coronary blood flow reversal, coincident with balloon deflation during end diastole, reduced net systemic forward flow. IABP inflation and deflation timed with traditional criteria and a high-fidelity blood pressure signal did not attenuate flow reversal or improve IABP efficacy. These findings suggest that brief end-diastolic flow reversal is an important and novel mechanism that may limit the effectiveness of IABP counterpulsation in pediatric patients with traditional inflation and deflation criteria.

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