Abstract

Objective—To assess the effect of endothelin type A (ET\textsubscript{A}) receptor antagonism in infants with pulmonary hypertension following corrective surgery for congenital heart disease.

Design—Open label, preliminary study.

Setting—Tertiary paediatric cardiothoracic surgical centre.

Patients—Three infants (aged 3 weeks, 7 weeks, and 8 months) with postoperative pulmonary hypertension unresponsive to conventional treatment, including inhaled nitric oxide.

Interventions—Patients received incremental intravenous infusions (0.1 to 0.3 mg/kg/h) of the ET\textsubscript{A} receptor antagonist BQ-123.

Main outcome measures—The response to BQ-123 administration was determined using continuous invasive monitoring of cardiorespiratory variables.

Results—BQ-123 infusion caused a reduction in the ratio of pulmonary to systemic pressures (0.62 (0.01) to 0.52 (0.03), mean (SEM)) with an accompanying decrease in right ventricular stroke work index (4.6 (0.4) to 2.5 (0.3) g/m) and a tendency for the cardiac index to rise (2.1 (0.2) to 2.7 (0.6) l/min/kg/m\textsuperscript{2}). This was associated with a well tolerated fall in the arterial partial pressure of oxygen (16.5 (4.1) to 12.4 (3.3) kPa) and mean systemic arterial pressure (57 (3) to 39 (3) mm Hg).

Conclusions—ET\textsubscript{A} receptor antagonism in infants with postoperative pulmonary hypertension after corrective surgery for congenital heart disease led to significant improvement in pulmonary haemodynamic indices. However, these benefits were associated with reductions in systemic blood pressure and arterial oxygen saturation, the latter consistent with a ventilation-perfusion mismatch. On the basis of these results, studies in pulmonary hypertension will need to proceed with caution.

(Heart 1999;82:505–508)

Keywords: endothelin-1; pulmonary hypertension; receptor antagonism; congenital heart disease

Postoperative pulmonary hypertension is a common clinical problem following successful surgical correction of congenital heart defects and may lead to significant morbidity and mortality.\textsuperscript{1} Its occurrence relates both to pre-existing pulmonary hypertension and the acute effects of surgery and cardiopulmonary bypass. Dysfunction of the pulmonary vascular endothelium appears to be a major contributing factor for the development of pulmonary hypertension in this group of patients.

Endothelin-1 is an extremely potent endothelium derived vasoconstrictor peptide\textsuperscript{2} which is released and cleared in the pulmonary circulation.\textsuperscript{3} Plasma concentrations of endothelin-1 are increased in subjects going to high altitude,\textsuperscript{4} in patients with chronic heart failure,\textsuperscript{5} and in patients with pulmonary hypertension.\textsuperscript{6} Moreover, in these conditions, the degree of pulmonary hypertension and pulmonary vascular resistance correlates closely with plasma endothelin-1 concentrations.\textsuperscript{4–6} Children with pulmonary hypertension\textsuperscript{7} and persistent pulmonary hypertension of the newborn\textsuperscript{8} also have raised plasma endothelin-1 concentrations that correlate with disease severity,\textsuperscript{9} and may play a role in its pathogenesis.\textsuperscript{10}

Studies in animal models of pulmonary hypertension have reported reversal of pulmonary hypertension with endothelin receptor antagonists\textsuperscript{11–13} and endothelin converting enzyme inhibition.\textsuperscript{11–13} Indeed, in a sheep model of pulmonary hypertension induced by aortopulmonary shunting in utero, endothelin antagonism eliminated the postoperative increase in pulmonary vascular resistance following cardiopulmonary bypass.\textsuperscript{14} Reddy and colleagues\textsuperscript{14} concluded that endothelin antagonism warrants further study in children at risk of pulmonary hypertension after surgical repair with cardiopulmonary bypass. There have been no published clinical studies to date assessing the therapeutic benefits of endothelin antagonism in postoperative pulmonary hypertension.

We report our preliminary experience with the therapeutic use of the endothelin type A (ET\textsubscript{A}) receptor antagonist, BQ-123, in three infants with postoperative pulmonary hypertension following corrective surgery for congenital heart disease.

Methods

Written informed parental consent was obtained for each child and the study was approved by the local research ethics committee.

BQ-123 (American Peptide Company, Sunnyvale, California, USA) was given under a Department of Health (UK) Doctors and Dentists Exemption Certificate.
The three infants (aged 3 weeks, 7 weeks, and 8 months) were anaesthetised according to our standard protocol. Phenoxybenzamine (1 mg/kg) was given before establishing cardiopulmonary bypass. Corrective surgery was performed after inducing systemic hypothermia and cold crystalloid cardioplegic arrest. A thermodilution pulmonary artery flow catheter (3 F; Baxter Health Care, Thetford, UK) and a left atrial line were inserted before discontinuing cardiopulmonary bypass.

After chest closure, the infants were returned to the intensive care unit and maintained on a standard regimen of vecuronium (0.1 mg/kg/h), fentanyl (5.0 µg/kg/h), and midazolam (0.1 mg/kg/h). Following rewarming, the ratio of pulmonary to systemic arterial pressure (P/S ratio) was determined using invasive monitoring. Infants were entered into the study if they did not have a residual left to right shunt on echocardiography and had a P/S ratio greater than 0.5 which did not respond to standard treatment, including inhaled nitric oxide (10 ppm increasing to 20, 30, and 40 ppm for 30 minutes at each dose). During the study period, the amount of sedation and inotropic support was maintained constant and atrial pressures kept stable using packed red blood cells or human albumin solution. Following stabilisation for three hours, BQ-123 was dissolved in 0.9% saline and given intra-venously at 0.1, 0.2, and 0.3 mg/kg/h, for 30 minutes at each dose.

Data are presented as mean (SEM). Haemodynamic variables were measured in triplicate at each time point and the mean taken. Recognising the small sample size and inherent variation between haemodynamic variables, non-parametric analyses (Wilcoxon rank sum) were used to compare variables before, during, and after BQ-123 infusion. Statistical significance was assumed at the 5% level.

**Results**

Characteristics of the patients are shown in table 1.

Baseline left and right atrial pressures were 9.6 (1.2) and 7.3 (0.3) mm Hg, respectively, and did not change during or after BQ-123 infusion. However, the P/S ratio fell in all patients during BQ-123 administration (p < 0.001; fig 1) and returned to baseline about 90 minutes after discontinuation of the infusion. Concomitant with the changes in the P/S ratio, the systemic arterial pressure fell (fig 1), although the fall was proportionately less than for the pulmonary arterial pressure and was well tolerated. Right ventricular stroke work index mirrored the changes in P/S ratio and fell significantly in response to BQ-123 infusion, from 4.56 (0.31) to 2.90 (1.90) g.m/m² (p < 0.001). The cardiac index tended to increase and left ventricular stroke work index fell from 8.0 (0.9) to 6.2 (0.9) g.m/m² (p < 0.001) but there were no changes in heart rate, acid–base balance, or urine output.

Despite haemodynamic improvements, the arterial partial pressure of oxygen fell in all three infants during BQ-123 infusion, from 16.5 (4.1) to 12.4 (3.3) kPa. Because of the

<table>
<thead>
<tr>
<th>Patient</th>
<th>Age (weeks)</th>
<th>Sex</th>
<th>Weight (g)</th>
<th>Diagnosis</th>
<th>Procedure</th>
<th>Inotropes and vasodilators</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>3</td>
<td>Female</td>
<td>3100</td>
<td>Anomalous aortic origin of left pulmonary artery, patent foramen ovale, persistent arterial duct</td>
<td>Mobilisation of left pulmonary artery with formation of pulmonary artery bifurcation, closure of patent foramen ovale, division of persistent arterial duct</td>
<td>Dopamine 3 µg/kg/min</td>
</tr>
<tr>
<td>2</td>
<td>7</td>
<td>Male</td>
<td>3500</td>
<td>Obstructed partial anomalous pulmonary venous drainage, left pulmonary artery stenosis, atrial septal defect</td>
<td>Right pulmonary vein to right atrium anastomosis, atrial diverting patch, left pulmonary artery patch</td>
<td>Dobutamine 20 µg/kg/min, Glyceryl trinitrate 2 mg/kg/min</td>
</tr>
<tr>
<td>3</td>
<td>25</td>
<td>Female</td>
<td>6100</td>
<td>Ventricular septal defect, persistent arterial duct (Aicardi and Poland syndromes)</td>
<td>Patch closure of ventricular septal defect, closure of persistent arterial duct</td>
<td>Dobutamine 20 µg/kg/min, Dopamine 3 µg/kg/min</td>
</tr>
</tbody>
</table>

![Figure 1](http://heart.bmj.com/)

**Figure 1** Effect of BQ-123 (0.1 to 0.3 mg/kg/min) on cardiac index (•), pulmonary/systemic ratio (○), heart rate (△), and mean systemic (□) and pulmonary (●) arterial pressure in infants with postoperative pulmonary hypertension following corrective surgery for congenital heart disease. (Error bars are SEM, n = 3.)
Endothelin antagonism in pulmonary hypertension after congenital heart surgery

A degree of systemic hypotension and impaired oxygenation is the inevitable consequence of effective systemic vasodilatation and these are the limiting factors in the clinical use of conventional agents. Intrapulmonary ventilation/perfusion matching is dependent upon local hypoxic vasoconstrictive reflexes and so is impaired by pulmonary vasodilatation. Although they were initially unresponsive to inhaled nitric oxide, improved oxygenation was seen in the two patients who received inhaled nitric oxide (20 ppm) after ETA receptor antagonism. This effect may be related to the recently described improvement in pulmonary vascular responsiveness to nitric oxide following ET\textsubscript{\alpha} receptor antagonism. The mechanisms of this effect remain to be established, but in an animal model of pulmonary hypertension, ET\textsubscript{\alpha} receptor antagonism was associated with both an improvement in endothelin-dependent vasodilatation and an increase in pulmonary vascular smooth muscle sensitivity to nitric oxide.

This first preliminary report of the use of ET\textsubscript{\alpha} receptor antagonism in infants with postoperative pulmonary hypertension following corrective surgery for congenital heart disease suggests an improvement in the pulmonary to systemic ratio and right ventricular stroke-work index. However, these benefits were counterbalanced by potentially adverse reductions in arterial oxygenation and systemic blood pressure. These findings suggest that endothelin antagonism, particularly in combination with inhaled nitric oxide, may represent a valuable new approach to the treatment of refractory postoperative pulmonary hypertension which merits further but cautious investigation.

We thank Dr M J Godman for funding the study through the Calderwood Research Endowment Fund. DEN was the recipient of a British Heart Foundation Junior Research Fellowship (FS/95/009). DJW is supported by a Research Leave Fellowship from the Wellcome Trust (WT 052633/0).


Early therapeutic experience with the endothelin antagonist BQ-123 in pulmonary hypertension after congenital heart surgery

B Prendergast, D E Newby, L E Wilson, D J Webb and P S Mankad

Heart 1999 82: 505-508
doi: 10.1136/hrt.82.4.505

Updated information and services can be found at:
http://heart.bmj.com/content/82/4/505

These include:

References
This article cites 22 articles, 10 of which you can access for free at:
http://heart.bmj.com/content/82/4/505#BIBL

Email alerting service
Receive free email alerts when new articles cite this article. Sign up in the box at the top right corner of the online article.

Topic Collections
Articles on similar topics can be found in the following collections

- Hypertension (3006)
- Drugs: cardiovascular system (8842)
- Congenital heart disease (762)
- Interventional cardiology (2933)

Notes

To request permissions go to:
http://group.bmj.com/group/rights-licensing/permissions

To order reprints go to:
http://journals.bmj.com/cgi/reprintform

To subscribe to BMJ go to:
http://group.bmj.com/subscribe/