TABLE 1. Surgical procedures

Associated procedures	No.	Mitral repair procedures	No.
CABG	3	Resection of posterior leaflet	4
TVR	1	Resection of anterior leaflet	1
		PTFE chordae implantation*	3
		Papillary muscle shortening	2

Preoperative and postoperative echocardiographic parameters			
Characteristics	Preoperative	Postoperative	1-yr follow-up
LA (mm)	55.4 ± 6.8	49.2 ± 5.4	41.4 ± 6.2†
LVED (mm)	65.2 ± 5.1	60.8 ± 5.5	51.2 ± 7.3†
LVES (mm)	43.5 ± 3.9	41.2 ± 6.1	36.1 ± 8.4

CABG, Coronary artery bypass graft; TVR, tricuspid valve repair; LA, left atrial; LVED, left ventricular end-diastolic; LVES, left ventricular end-systolic; PTFE, polytetrafluoroethylene.

*Chordal replacement with a 4.0 PTFE suture.

†Values differ significantly from preoperative values ($P < .1$) (t test for paired samples and Wilcoxon matched-pairs signed-ranks test).

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Inhaled nitric oxide does not improve systemic oxygenation after bidirectional superior cavopulmonary anastomosis

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Inhaled nitric oxide (NO) is a selective pulmonary vasodilator that decreases intrapulmonary shunt fraction.¹ We sought to determine whether inhaled NO would ameliorate hypoxemia after bidirectional superior cavopulmonary anastomosis (BCPA).

Methods

The research and ethics review board approved the study, and the parents of all children gave informed and written consent. We included patients with systemic oxygen saturations of 75% or less after BCPA without venous decompressing collaterals or pulmonary disease. Patients were mechanically ventilated and studied on the first postoperative day in the intensive care unit. From a stable baseline, patients received a 15-minute trial of inhaled NO (80 ppm).² During the trial of inhaled NO, ventilator parameters and FiO_2 were unchanged. In 5 patients, cyclic guanosine monophosphate (cGMP) levels were measured.² One patient first received inhaled NO in the cardiac catheterization laboratory because of persistent hypoxemia after coil embolization of decompressing venous collaterals. This same patient was readmitted 7 months after BCPA with refractory hypoxemia as the result of respiratory syncytial viral pneumonitis and was treated with 10 ppm NO.

Statistical Analysis

A paired Student t test with a Bonferroni correction for 3 comparisons was used. The difference between cGMP levels was compared by the nonparametric Wilcoxon signed-rank test.

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TABLE 1. Hemodynamic and blood gas response to inhaled nitric oxide after bidirectional superior cavopulmonary anastomosis (mean FiO_2 0.76 ± 0.2)

	Mean SVCp (mm Hg)	Mean CAp (mm Hg)	TPG (mm Hg)	Mean BP (mm Hg)	SVC sat (%)	Art sat (%)	Art Pao ₂ (mm Hg)	Art Paco ₂ (mm Hg)	Art pH	HR (beats/min)
Pre-NO	17 $\pm$ 3	7 $\pm$ 2	9 $\pm$ 5	65 $\pm$ 11	33 $\pm$ 11	63 $\pm$ 7	32 $\pm$ 4	38 $\pm$ 9	7.46 $\pm$ 0.08	143 $\pm$ 21
With NO	15 $\pm$ 3*	7 $\pm$ 2	7 $\pm$ 5	62 $\pm$ 9	34 $\pm$ 12	65 $\pm$ 5	32 $\pm$ 4	38 $\pm$ 8	7.46 $\pm$ 0.07	146 $\pm$ 20
Post-NO	16 $\pm$ 3†	8 $\pm$ 2	8 $\pm$ 4	63 $\pm$ 12	31 $\pm$ 7	64 $\pm$ 7	32 $\pm$ 4	38 $\pm$ 8	7.46 $\pm$ 0.06	146 $\pm$ 18

Art, Systemic arterial; BP, blood pressure; CAp, common atrial pressure; HR, heart rate; NO, nitric oxide; SVC, superior vena cava; SVCp, superior vena cava pressure; TPG, transpulmonary gradient. Values are mean $\pm$ SD.

* $P = .002$ compared with pre-NO.

† $P = .013$ compared with pre-NO.

TABLE 2. Cyclic guanosine monophosphate levels in response to inhaled nitric oxide after bidirectional superior cavopulmonary anastomosis

	SVC pmol/mL, median (range)	Artery pmol/mL, median (range)
Pre-NO	11.2 (9.4-14.0)	9.7 (9.2-13.2)
With NO	51.8 (30-124)*	54.8 (36-128)*

SVC, Superior vena cava; NO, nitric oxide.

* $P = .04$ pre-NO versus with NO.

Results

We studied 26 patients with a median age of 0.5 years (range 0.2-13 years) and a weight of 6.5 kg (range 3-41 kg) who had undergone BCPA. All patients were undergoing the second stage of cardiac palliation for a functionally single ventricle. The diagnoses were hypoplastic left heart syndrome in 19 patients, tricuspid atresia with transposed great arteries with aortic obstruction in 3 patients, unbalanced atrioventricular canal defects in 2 patients, Ebstein anomaly of the tricuspid valve in 1 patient, and double-inlet left ventricle in 1 patient.

The results of the trial of inhaled NO are displayed in Tables 1 and 2. There was no change in systemic oxygenation in response to inhaled NO. However, there was a significant decrease in superior vena caval pressure. Plasma cGMP increased significantly.

One patient received a trial of inhaled NO during cardiac catheterization to coil decompressing venous collaterals. After deployment of coils, the aortic oxygen saturation increased from 50% to 62% with an FiO_2 of 1.0. Inhaled NO changed aortic saturation minimally from 62% to 64%. Superior vena caval mean pressure decreased from 16 to 14 mm Hg. Pulmonary vein oxygen saturation and oxygen tension increased from 97% to 99% and 94 to 260 mm Hg, respectively. This patient was readmitted with respiratory syncytial viral pneumonitis, hypoxemia, and facial edema without cavopulmonary obstruction. The inhaled NO was administered at 10 ppm, systemic oxygen saturations increased from 48% to 87%, and transpulmonary gradient decreased from 12 to 4 mm Hg. Treatment was continued for 93 hours with gradual resolution of pneumonia.

The mean hemoglobin concentration was 13.9 ± 2 g/dL, and the methemoglobin concentration was 0.6 ± 1.4 g/dL. Nitrogen dioxide levels ranged from less than 1 to 4 ppm.

Discussion

We found that inhaled NO does not improve systemic oxygenation after BCPA in the absence of intrapulmonary shunting. The decrease in superior vena caval pressure and increase in plasma cGMP indicate that sufficient NO was delivered to activate guanylate cyclase and cause vasodilation. This suggests that pulmonary vasoconstriction does not limit pulmonary blood flow or determine systemic oxygen saturations after BCPA. In 1 patient with pulmonary venous desaturation, inhaled NO enhanced ventilation perfusion matching with substantial improvement in oxygenation. Previous studies of the effect of inhaled NO after BCPA are equivocal. In 4 patients, oxygenation deteriorated when NO was withdrawn, and 5 patients demonstrated a 5% increase in systemic oxygen saturation.^{3,4} Clinical details are insufficient to determine whether these patients had pulmonary venous desaturation.^{3,4} In contrast, we showed in 26 patients that inhaled NO does not improve systemic oxygenation in the absence of pulmonary venous desaturation after BCPA. These observations suggest that cardiac output and cerebral blood flow after BCPA are more important determinants of oxygenation than pulmonary vascular resistance. Indeed, recent studies demonstrated that a strategy of permissive hypercapnia improves oxygenation after BCPA.^{5,6} Although we report a "negative study," it has important implications both for the additional insight provided into the unique physiology of BCPA and the therapeutic economics. Inhaled NO has been transformed from an inexpensive medical grade gas to a costly pharmaceutical grade gas. Therefore, the use of inhaled NO should be justified fiscally and therapeutically. Our study suggests that after BCPA, strategies that increase cardiac output and cerebral blood flow (eg, hypercapnia) are more likely to improve oxygenation than inhaled NO unless there is concomitant pulmonary venous desaturation.

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Bonanno's catheter: A less invasive and cost-effective alternative for drainage of pleural effusion

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Pleural effusion is a common condition in both medical and surgical specialties. Pleural drainage is a method used to remove a collection of air, fluid, pus, or blood from the pleural space to restore normal lung expansion and function. In this context, although tube thoracostomy remains the gold standard, it is quite invasive, requiring significant blunt dissection, and therefore resulting in a great deal of patient discomfort. We suggest Bonanno's catheter as a less traumatic but equally effective alternative. Bonanno's catheter was first designed for suprapubic bladder drainage¹ and has been an important tool for the urologist because of its efficacy, simplicity, and wide range of applications. Its design is therefore well suited for drainage of other serous fluid collection, including pleural effusion. However, it is not suitable for frank hemothorax.

Technique

The patient is preferably positioned at 45° or sitting up, leaning slightly forward. The entry site can be clinically identified by means of percussion of the chest to elicit an area of stony dullness around the midaxillary line. After the skin has been prepared and the chest draped, local anesthetic is infiltrated up to the parietal pleura. The presence of effusion should be confirmed by aspiration with the same syringe as a guide. Bonanno's catheter is assembled as per the instruction kit, and a syringe is attached to one end

Figure 1. Bonanno's catheter and syringe.

(Figure 1). A 3- to 4-mm incision is made in the skin to avoid damage to the catheter tip, and the catheter is inserted gradually while applying constant suction on the syringe. As soon as serous fluid is aspirated, the trocar is held in a static position while advancing the cannula into the pleural cavity. The trocar is removed, and serous fluid should flow freely from the catheter, which can now be attached with the rubber tubing to a conventional underwater drainage system. The catheter is sutured to the skin, and a chest radiograph will confirm the position of the catheter (Figures 2 and 3).

Conclusion

Apart from standard tube thoracostomy, which can be quite invasive and painful,² patients with pleural effusion could also undergo

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