Weaning strategy with inhaled nitric oxide treatment in persistent pulmonary hypertension of the newborn

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Abstract

Aim—To determine if infants who had become dependent on inhaled nitric oxide treatment could be successfully weaned off it if FIO₂ was increased briefly during withdrawal.

Methods—Sixteen infants admitted for conditions associated with increased pulmonary vascular resistance responded well to inhaled nitric oxide treatment with a significant increase in PaO₂ (maximum inhaled nitric oxide given 25 ppm). Weaning from inhaled nitric oxide in 5 ppm decrements was initiated once the FIO₂ requirement was less than 0.5. When patients were stable on 5 ppm of inhaled nitric oxide, the gas was then discontinued. If a patient showed inhaled nitric oxide dependence—that is, oxygen saturation fell by more than 10% or below 85%—inhaled nitric oxide was reinstated at 5 ppm and the patient allowed to stabilise for 30 minutes. At this time, FIO₂ was increased by 0.40 and weaning from inhaled nitric oxide was attempted again.

Results—Nine infants were successfully weaned on the first attempt. The seven infants who failed the initial trial were all successfully weaned following the increase in FIO₂. After successful weaning, FIO₂ was returned to the pre-weaning level in mean 148(SD 51) minutes and inhaled nitric oxide was never reinstated.

Conclusion—Infants showing inhaled nitric oxide dependency can be successfully weaned by increasing FIO₂ transiently.

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Keywords: inhaled nitric oxide; weaning; ECMO.

A significant number of infants who respond favourably to inhaled nitric oxide treatment prove difficult to wean off it. This dependence necessitates prolonged use of inhaled nitric oxide or, on occasion, the use of alternative treatments, such as extracorporeal membrane oxygenation (ECMO).

Nitric oxide (NO), or endothelium derived relaxing factor, is an endogenously produced substance which helps regulate vascular smooth muscle tone in the circulation of many organs, including the perinatal lung.¹⁻⁷ Therapeutic inhalation of NO in various pulmonary diseases is currently under evaluation in several neonatal intensive care units. On the basis of preliminary reports, there is little doubt that inhaled nitric oxide is effective in raising systemic arterial oxygen concentration in several clinical situations where pulmonary vascular resistance is raised.⁶⁻⁹ This change in oxygenation is presumed to result from a reduction in pulmonary vascular resistance and diminution in right to left shunting. The clinical subtleties of the use of inhaled nitric oxide are as yet undefined.¹⁰⁻¹⁶ In particular, many

Table 1 Characteristics of study population

Case No	Sex	Birthweight (g)	Gestational age (weeks)	Diagnosis	Age at starting INO (hour)	pH at starting INO	PaCO ₂ at starting INO (mm Hg)	PaO ₂ at starting INO (postductal) (mm Hg)	PaO ₂ before INO weaning (postductal) (mm Hg)	Duration of INO (hour)
1*	F	3900	40	MAS	48	7.29	41	28	56	105
2*	M	2600	37	MAS	25	7.10	65	13	90	32
3*	F	3600	40	CHD	288	7.37	48	38	46	74
4*	F	657	26	HMD	34	7.15	67	28	79	49
5*	M	3230	40	Sepsis	28	7.30	48	22	59	132
6*	M	3100	40	CHD	48	7.27	56	21	43	146
7*	M	3153	42	MAS	45	7.35	35	39	51	92
8	M	3770	39	CDH	140	7.37	50	28	45	45
9	M	3900	39	1° PPHN	22	7.05	69	41	76	85
10	M	3011	40	MAS	13	7.36	55	36	62	128
11	F	3800	42	MAS	36	7.09	78	20	105	35
12	M	2600	40	1° PPHN	29	7.12	67	15	41	74
13	F	3100	41	MAS	9	6.89	87	18	68	44
14	F	3960	40	CHD	48	7.42	25	44	86	50
15	M	4165	41	MAS	29	7.44	28	59	73	105
16	M	3335	40	Sepsis	11	7.05	57	26	57	192

*Failed initial attempt at weaning; MAS meconium aspiration syndrome; CHD congenital heart disease; HMD hyaline membrane disease; CDH congenital diaphragmatic hernia; 1° PPHN=primary (idiopathic) persistent pulmonary hypertension of the newborn.

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Figure 1 In panel (A) measurements of mean blood pressure, heart rate, arterial oxygen saturation (SpO_2), FIO_2 and NO concentrations obtained from an infant in group I are plotted against time in minutes. Note stability of measurements as INO is withdrawn. In panel (B) the same measurements are plotted for an infant in group II. Acute deterioration in all variables followed the initial attempt at weaning. FIO_2 was increased and the weaning was successful. Note, in particular, how quickly FIO_2 was reduced following successful weaning.

observers have reported difficulty in discontinuing treatment with inhaled nitric oxide, in both clinical and experimental settings.^{9 17–20} We recently observed that seven of 16 infants who failed initial attempts at discontinuation of inhaled nitric oxide treatment were successfully and quickly weaned when the inhaled FIO_2 was increased briefly during the withdrawal of inhaled nitric oxide.

Methods

All newborn infants admitted to the neonatal intensive care unit between December 1994 and July 1995 with severe hypoxaemia and clinical and echocardiographic evidence of pulmonary hypertension were considered for this study. Infants whose hypoxaemia proved refractory to optimal ventilatory and cardiopulmonary support were offered treatment with inhaled nitric oxide; all parents opted for this treatment. The study was approved by the Institutional Review Board and parental informed consent was obtained for all patients.

Table 2 Clinical characteristics of group I infants compared with those of group II infants (mean (SD))

Criteria	Group I (n=9)	Group II (n=7)	P value
Birthweight	3.5 (0.5)	2.9 (1.0)	NS
Gestational age (weeks)	40.2 (1.0)	37.9 (5.4)	NS
PaO ₂ before INO (mm Hg)	31.9 (14.4)	27.0 (9.3)	NS
PaO ₂ before weaning NO (mm Hg)	68.1 (20)	60.1 (17.6)	NS
Duration of INO (hours)	84 (51)	90 (42)	NS

NS=not significant.

Twelve of these 16 infants had been hyper-ventilated at the referring hospital and met otherwise generally accepted criteria for extracorporeal membrane oxygenation (ECMO) treatment.²¹ The clinical characteristics of the study population are shown in table 1.

Nitric oxide was introduced into the afferent limb of the ventilator circuit via an adapter placed 18 inches upstream from the endotracheal tube. Nitric oxide treatment was initiated at a dose of 25 ppm. Continuous measurements were made of heart rate, systemic blood pressure, and pre- and post-ductal pulse oximetry before and after initiation of inhaled nitric oxide treatment. Frequent serial measurements of arterial pH, arterial carbon dioxide activity, and arterial oxygen activity were recorded as well. Data were collected and logged continuously using the V_T 1000 Neonatal Workstation (Vitaltrends Technology, Inc, Hackensack, NJ).

If an infant responded to inhaled nitric oxide with a significant increase in arterial saturation to above 85%, the treatment was continued at the same dose. If an infant failed to respond to inhaled nitric oxide, management with ECMO was started and the medication discontinued.

As the oxygenation improved, FIO_2 , ventilatory pressure and rate were decreased while the concentration of inhaled nitric oxide remained at therapeutic levels—greater than 5 ppm. Once the pre- and post-ductal oxygen saturation gradient was insignificant on modest FIO_2 (usually 0.40–0.45) and mean airway pressure

was less than 10 cm H₂O, weaning of inhaled nitric oxide started. Weaning was attempted in 5 ppm decrements every 4 hours, as tolerated by the infant. The FIO₂ was kept constant during the weaning phase. Heart rate, systemic blood pressure, and oxygen saturations were carefully monitored and arterial blood gases were obtained every 4 hours. If the infant tolerated weaning by 5 ppm for 4 hours, further weaning in 5 ppm decrements continued until a concentration of 5 ppm of inhaled nitric oxide was reached. Once the infant was stable on 5 ppm, inhaled nitric oxide treatment was discontinued. Heart rate, systemic blood pressure, and oxygen saturations continued to be closely monitored. If the saturation dropped by 10% or below 85% it was considered a weaning failure and inhaled nitric oxide was reinstated at 5 ppm. A second attempt at weaning from inhaled nitric oxide was made 30 minutes later when the infant's heart rate, blood pressure, and saturation were stable. However, before the second attempt, the FIO₂ was increased by 0.4. If the infant remained stable during this attempt at weaning, the FIO₂ was then decreased over the next few hours, as tolerated.

DATA ANALYSIS

Mean values for all variables for each infant were computed every 60 seconds and stored on

Figure 2 Changes in mean blood pressure, heart rate, and arterial oxygen saturation following weaning without oxygen (trial I) and with oxygen (trial II) for the seven infants who failed trial I.

the hard disc of the V_T 1000*. The data were subsequently analysed using commercial statistical software (Systat, Inc, Evanston, IL, USA). Depending on the response to weaning, infants were classified into two groups: group I infants were successfully weaned from inhaled nitric oxide without the need for additional oxygen; and group II infants did not tolerate weaning (as evidenced by decreasing oxygen saturation by 10% or less than 85%) and required reinstitution of inhaled nitric oxide. Group II infants subsequently underwent weaning while receiving higher concentrations of oxygen. Table 2 shows that there were no differences in birthweight, gestational age, duration of inhaled nitric oxide treatment, PaO₂ at time of introduction of inhaled nitric oxide, or PaO₂ at initial inhaled nitric oxide weaning between group I and II infants.

For purposes of analysis, the experimental protocol was divided into five phases: phase 1, baseline period; phase 2, weaning without incremental FIO₂; phase 3, restabilisation; phase 4, weaning from inhaled nitric oxide with supplemental oxygen; and phase 5, weaned from inhaled nitric oxide and back to the pre-weaning concentration of oxygen. The averages of all minute means of heart rate, blood pressure, and oxygen saturation for each phase of the study were computed. In addition to measurements of mean heart rate, mean blood pressure, and mean oxygen saturation during phases 1, 3, and 5, the maximum and minimum measurements during phases 2 and 4 (times of weaning) were also determined. Statistical comparisons between various phases within and between the two groups were made using paired *t* tests. Differences were considered significant at P < 0.05.

Results

Nine of the 16 patients were weaned successfully from inhaled nitric oxide without an increase in oxygen concentration. The other seven infants required reinstitution of inhaled nitric oxide because of unfavourable changes in oxygen saturation. The data from two infants, one from group I and another from group II, are shown in fig 1. These plots show the differences in physiological measurements during initial successful and unsuccessful weaning. All seven infants who failed initial weaning attempts were successfully weaned from inhaled nitric oxide when the FIO₂ was increased. Comparison of the means and standard deviations of the changes in blood pressure, heart rate, and oxygen saturation during weaning attempts in group II with and without increasing FIO₂ are shown in fig 2. In group II (n=7) the differences in all variables during the initial weaning attempt without oxygen and the second attempt with additional oxygen were highly significant. As evidence of successful weaning, measurements of the same variables at the onset of weaning (phase 1), and when the concentration of inhaled oxygen was returned to preweaning levels (phase 5), were not significantly different (n=16). The duration required for inhaled oxygen to be returned to

its preweaning level in group II infants (mean (SD)) was 148 (51) minutes.

Discussion

Endogenously produced NO is now recognised as an important modulator of fetal and neonatal pulmonary vascular tone.¹⁻⁷ It is produced by the normal human lung²² and can be found in exhaled gas.²⁴ Inhaled NO diffuses from the alveolar space through the alveolar wall and reaches the vascular smooth muscles of the small pulmonary arteries, where, in combination with endogenous NO, it enhances vasodilatation. Some NO traverses the endothelial cell and enters the vessel lumen, where it is rapidly inactivated by haemoglobin and, therefore, does not reach the systemic circulation.^{8 19 25} Therapeutic inhalation of NO (20 to 80 ppm) to supplement endogenous NO production seems to be an effective treatment for increased pulmonary vascular resistance due to a variety of underlying pathologies.

Early reports illustrate that there may be difficulty in weaning inhaled nitric oxide due to a "rebound" phenomenon—that is, a sudden deterioration in oxygenation after withdrawal of inhaled nitric oxide treatment. This rebound has been observed in both clinical and experimental settings.^{9 17-21} Heretofore, it has often been necessary to reinstate the inhaled nitric oxide treatment before subsequently repeating the weaning process. Occasionally, weaning has been delayed for as long as three weeks because of rebound hypoxaemia.¹⁷ The results of this study indicate that infants can be successfully weaned from inhaled nitric oxide by simply increasing the inhaled concentration of oxygen while inhaled nitric oxide is being withdrawn. In fact, early weaning from inhaled nitric oxide during the recovery phase of persistent pulmonary hypertension of the newborn was possible in all 16 infants described above. Despite major signs of deterioration during the initial attempt, all seven of the infants who failed the first attempt at weaning were subsequently weaned quickly with increased supplemental oxygen.

This study does not allow us to rule out the possibility that the second attempt at weaning was successful because of spontaneous improvement in the patients' underlying disease. However, the fact that weaning was successful within hours of the original weaning failure makes this explanation unlikely. Meanwhile, testing of this hypothesis by randomly assigning infants who fail initial attempts at weaning to treatment with and without incremental oxygen is recommended.

There are several possible explanations for this effect of supplemental oxygen. Nitric oxide and NO donor agents are known to inhibit endothelium dependent relaxation of arterial rings and endogenous NO generation from intact endothelial cells, without diminishing the sensitivity of the vascular smooth muscle cells to the relaxing effect of exogenous NO.²⁶ This observation suggests that inhaled nitric oxide inhibits the endothelial NO synthase enzyme by a direct feedback mechanism. A simple feedback mechanism could explain the

acute clinical deterioration of infants on withdrawal of inhaled nitric oxide treatment as endogenous NO might not be produced at a sufficient rate to replace the exogenous supply.

The fact that oxygen was able to prevent weaning failures could have been due to a direct vasodilatory effect of increased arterial oxygen activity in the alveolus and/or pulmonary circulation, at a time when synthesis of NO was recovering from earlier suppression. On the other hand, the effect of additional oxygen may be explained by enhanced NO production due to improved oxygenation. Dollberg *et al*²⁷ demonstrated that increasing oxygenation after initiation of ECMO does seem to be associated with increased NO synthesis, as evidenced by increased urinary nitrate and nitrate concentration in patients with persistent pulmonary hypertension of the newborn following institution of ECMO. Furthermore, it is well established that NO mediates pulmonary vascular tone via guanosine-3',5'-cyclic monophosphate (cGMP).²⁸⁻³⁰ In vivo animal studies have shown that inhibitors of cGMP-specific phosphodiesterase potentiate and substantially prolong the duration of the pulmonary vasodilating action of NO.²⁹ Animals pre-treated with one of these agents, Zaprinast, did not show the acute withdrawal reaction on discontinuation of inhaled nitric oxide treatment.³¹ This prevention of the rebound response, similar to what we observed with increased FIO₂, suggests that inhibition of cGMP-specific phosphodiesterase by oxygen may be another mechanism to account for the beneficial effects of oxygen during inhaled nitric oxide withdrawal. Finally, oxygen may be helpful in clearing N-nitro L-arginine, a competitive inhibitor of NO that increases pulmonary artery pressure and is cleared in the blood by oxidation.³² These data suggest that a second attempt at weaning infants from inhaled nitric oxide should be made with increased supplemental oxygen, whenever the overall clinical condition of the infant indicates weaning is appropriate.

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- Davidson D, Eldemerdash A. Endothelium-derived relaxing factor: Presence in pulmonary and systemic arteries of the newborn guinea pig. *Pediatr Res* 1990;27:128-32.
- Davidson D, Eldemerdash A. Endothelium-derived relaxing factor: evidence that it regulates pulmonary vascular resistance in the isolated newborn guinea pig lung. *Pediatr Res* 1991;29:538-42.
- Abman SH, Chatfield BA, Hall SL, Mc Murtry IF. Role of endothelium-derived relaxing factor during transition of pulmonary circulation at birth. *Am J Physiol* 1990;259:H1921-7.
- Shaul PW, Farrar MA, Zellers TM. Oxygen Modulates endothelium-derived relaxing factor production in fetal pulmonary arteries. *Am J Physiol* 1992;262:H355-64.
- Tiktinsky MH, Cummings JJ, Morin III FC. Acetyl choline increases pulmonary blood flow in intact fetuses via endothelium-dependent vasodilatation. *Am J Physiol* 1992;262:H406-11.
- Kinsella JP, Neish SR, Shaffer E, Abman SH. Low dose inhalational nitric oxide in persistent pulmonary hypertension of the newborn. *Lancet*. 1992;340:819-20.
- Roberts JD Jr, Chen T-Y, Kawai N, Wain J, Dupuy P, Shimouchi A, *et al*. Inhaled nitric oxide reverses pulmonary vasoconstriction in the hypoxic and acidotic newborn lamb. *Circ Res* 1993;72:246-54.

- 8 Pison U, Lopez FA, Heideimeyer CF, Rossaint R, Falke K. Inhaled nitric oxide selectively reverses hypoxic pulmonary vasoconstriction without impairing pulmonary gas exchange. *J Appl Physiol* 1993;74:1287-92.
- 9 Roberts JD, Polaner DM, Lang P, Zapol WM. Inhaled nitric oxide in persistent pulmonary hypertension of the newborn. *Lancet*. 1992;340:818-9.
- 10 Stephens RJ, Freeman G, Evans MJ, Park M. Early response of lungs to low levels of nitrogen dioxide. *Arch Environ Health* 1972;24:160-79.
- 11 Hugod C. Effect of exposure to 43 ppm nitric oxide and 3.6 nitrogen dioxide on rabbit lung. *Int Arch Occup Environ Health* 1979;42:156-67.
- 12 Hugod C. Ultrastructural changes of the rabbit lung after 5 ppm nitric oxide exposure. *Arch Environ Health* 1977;13-6.
- 13 Chang LY, Mercer RR, Stockstill BL, Miller FJ, Graham JA, Ospital JJ, et al. Effects of low levels of NO₂ on terminal bronchiole cells and its relative toxicity compared to O₃. *Toxicol Appl Pharmacol* 1988;96:451-64.
- 14 Dowell AR, Kilburn KH, Pratt PC. Short term exposure to nitrogen dioxide. Effects on pulmonary ultrastructure, compliance and the surfactant system. *Arch Intern Med* 1971;128:74-80.
- 15 Oda H, Kusumoto S, Nakajima T. Nitrosyl-hemoglobin formation in the blood of animals exposed to nitric oxide. *Arch Environ Health* 1975;30:453-65.
- 16 Chiodi H, Mohler JG. Effects of exposure of blood hemoglobin to nitric oxide. *Environ Res* 1985;37:355-63.
- 17 Goldman AP, Tasker RC, Haworth SG, Sigston PE, Macrae DJ. Four patterns of response to inhaled nitric oxide for persistent pulmonary hypertension of the newborn. *Pediatrics* 1996;98:706-13.
- 18 Kinsella JP, Niesh SR, Ivy DD, Shaffer E, Abman SH. Clinical responses to prolonged treatment of persistent pulmonary hypertension of the newborn with low dose inhaled nitric oxide. *J Pediatr*. 1993;123:103-8.
- 19 Frostell C, Fratacci MD, Wain JC, Jones R, Zapol WM. Inhaled nitric oxide. A selective pulmonary vasodilator reversing hypoxic pulmonary vasoconstriction. *Circulation*. 1991;83:2038-47.
- 20 Petros A. Down-regulation of endogenous nitric oxide production after prolonged administration. *Lancet* 1994;344:191.
- 21 Miller O, Tang S, Keech A, Celermajor D. Rebound pulmonary hypertension on withdrawal from inhaled nitric oxide. *Lancet* 1995;346:51-2.
- 22 Short BL. Clinical management of the neonatal ECMO patient. In: Arensman R, Cornish D, eds. *Extra-corporeal Life Support*. Oxford: Blackwell Scientific, 1993:195-206.
- 23 Palmer RM, Ashton DS, Moncada S. Vascular endothelial cells synthesize nitric oxide from L-arginine. *Nature* 1988;333:664-6.
- 24 Gustafsson LE, Leone AM, Persso MG. Endogenous nitric oxide is present in the exhaled air of rabbits, guinea pigs and humans. *Biochem Biophys Res Commun* 1991;181:852-7.
- 25 Fratacci MD, Frostell CG, Chen TY, Wain Jr, Robinson DR, Zapol WM. Inhaled nitric oxide: A selective pulmonary vasodilator of heparin-protamine vasoconstriction in the sheep. *Anesthesiol* 1991;75:990-9.
- 26 Ravichandran LV, Johns RA, and Rengasamy A. Direct and reversible inhibition of endothelial nitric oxide synthase by nitric oxide. *Am J Physiol* 1995;268:H2221-3.
- 27 Dollberg S, Warner BW, Myatt L. Urinary nitrite and nitrate concentrations in patients with idiopathic persistent pulmonary hypertension of the newborn and effect of extracorporeal membrane oxygenation. *Pediatr Res* 1995;37:31-4.
- 28 Rapoport RM, Murad F. Agonist-induced endothelium-dependent relaxation in rat thoracic aorta may be mediated through CGMP. *Circ Res* 1983;52:352-7.
- 29 Griffith TM, Edwards DH, Lewis MJ, Henderson AH. Evidence that cyclic guanosine monophosphate (CGMP) mediates endothelium-dependent relaxation. *Eur J Pharmacol* 1985;112:195-202.
- 30 MacLeod KM, Ng DDW, Harris KH. Evidence that CGMP is the mediator of endothelium-dependent inhibition of contractile responses of rat arteries to adrenoceptor stimulation. *Mol Pharmacol* 1987;32:59-64.
- 31 Ichinose F, Adrie C, Hurford WE, and Zapol WM. Prolonged pulmonary vasodilator action of inhaled nitric oxide by zaprinast in awake lambs. *J Appl Physiol* 1995;78:1288-95.
- 32 Fineman JR, Heymann MA, Soifer SJ. N-omega-nitro-L-arginine attenuates endothelium dependent pulmonary vasodilatation in lambs. *Am J Physiol* 1991;260: H1299-306.