Haemodynamic changes during high frequency oscillation for respiratory distress syndrome

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Abstract
In a crossover trial left ventricular output (LVO), cerebral blood flow velocity (CBFV), and resistance index (RI) of the anterior cerebral artery were compared using Doppler ultrasonography, in eight preterm infants with respiratory distress syndrome (RDS) during conventional mechanical ventilation and high frequency oscillation. LVO was 14% to 18% lower with high frequency oscillation. There were no significant changes in CBFV. On the first day of life there was a trend towards lower RI on high frequency oscillation; the fall in LVO on high frequency oscillation was not related to lung hyperinflation.

Changes in ventilation type (from conventional mechanical ventilation to high frequency oscillation, or vice versa) can induce significant LVO changes in preterm infants with RDS.

(Arch Dis Child 1996; 74: F172-F176)

Keywords: high frequency oscillation, cardiac output, cerebral blood flow, Doppler ultrasonography.

High frequency oscillation is an effective way of supporting gas exchange in critically ill neonates with respiratory distress syndrome (RDS).\(^1\)\(^2\) Cerebral damage during high frequency oscillation is a controversial issue. While Clarke et al\(^1\) and Ogawa et al\(^2\) reported no increased risk of intracranial haemorrhage, both HIFI\(^1\) and HIPPO\(^2\) trials showed that severe intracranial haemorrhage was more common during high frequency oscillation than during conventional mechanical ventilation. In premature baboons with RDS\(^3\) or in surfactant deficient adult rabbits,\(^4\) high frequency oscillation did not affect the cerebral blood flow. Animal studies of cardiac output during high frequency oscillation showed that increasing mean airway pressure (Paw) could decrease cardiac output,\(^5\) but that strategies which avoid lung hyperinflation prevented a fall.\(^5\)\(^6\)

To our knowledge, no study on cardiac output or on left ventricular output (LVO) and cerebral blood flow velocity (CBFV) has been reported in preterm infants with RDS during high frequency oscillation. Based on animal studies,\(^5\)\(^6\) our hypothesis was that high frequency oscillation, used with a cautious strategy of lung volume recruitment, would not impair LVO or CBFV in premature infants with RDS.

Methods
Ten preterm infants with a clinical and radiological diagnosis of RDS were studied. None had presented with significant perinatal asphyxia (defined as an Apgar score <4 at 5 minutes and a cord blood pH <7-10), and none had a congenital heart or brain malformation or an intracranial haemorrhage at the time of the study. Their median birthweight was 1395 g (range 650–2650 g) and their gestational age 31 weeks (26–36 weeks). Parental informed consent was obtained in all cases and the protocol was approved by the research ethics committee of the Centre Hospitalier Universitaire Vaudois.

The study comprised two phases. Initially, all infants were on a time-cycled, pressure-limited ventilator (Bear Cub BP 2001) with a positive end expiratory pressure (PEEP) of 4–6 cm H\(_2\)O and an inspiratory time of 0.4–0.7 seconds. \(\text{FiO}_2\), peak inspiratory pressure, and breath rate were set to maintain \(\text{PaO}_2\) between 60 and 80 mm Hg (8 and 10.7 kPa) and \(\text{PaCO}_2\) between 40 and 50 mm Hg (5.3 and 6.7 kPa). All infants received continuous morphine infusion and were paralysed with pancuronium bromide. After initial resuscitation and stabilization of blood pressure with intravenous albumin 20% (median 24 ml/kg (range 4–35)) or dobutamine infusion (two infants), a chest radiograph was taken. The endotracheal tube was suctioned and 5 ml/kg of synthetic surfactant (Exosurf) was instilled over 20 minutes via the sideport on the special endotracheal tube adapter, without interrupting mechanical ventilation. Five minutes after the end of surfactant administration, ventilation settings, mean arterial blood pressure (MAP), arterial gases, CBFV and LVO were measured. The infants were then immediately transferred to high frequency oscillation. Five 1 second sustained inflations at 10 cm H\(_2\)O over the Paw delivered by conventional mechanical ventilation were administered. After five minutes, the same variables were measured. These paired measurements constituted phase 1 of the study (T1).

High frequency oscillation was obtained using an OHF-1 high frequency oscillatory ventilator (Dufour Cie, Villeneuve d’Asq, France) which generates sinusoidal pressure swings with a motor driven piston at an inspiratory:expiratory ratio of 1:1. The frequency was set at 15 Hz. Oscillatory amplitude was adjusted to achieve adequate \(\text{PaCO}_2\). The initial Paw on high frequency oscillation was 2 cm H\(_2\)O higher than that delivered by conventional mechanical ventilation. A chest radiograph was obtained a few hours later with the same Paw and lung inflation was assessed. Hyperinflation was defined as flattening of the diaphragm borders below the eighth posterior thoracic rib.
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AAD02 was computed according to the following formula:

$$713 \cdot FIO_2 - PaCO_2 - PaO_2 \text{ (mmHg)}.$$  

During recovery from RDS, infants were transferred from high frequency oscillation to conventional mechanical ventilation. Ventilator settings on conventional mechanical ventilation were as described initially. At this stage, all infants were receiving continuous morphine and dobutamine infusion. Three out of eight infants were paralysed with pancuronium MABP. Ventilation settings, bromide, arterial gases, CBFV and LVO were again measured during high frequency oscillation and conventional mechanical ventilation. These paired measurements constituted phase 2 of the study (T2).

Haemodynamic measurement techniques

Doppler ultrasound recordings were made using the duplex Doppler Acuson 128 with a 7-0 MHz transducer. All infants were kept supine with the head in anatomical position during measurements. Each variable was analysed on line, but the recorded values were read on photographic prints only after the second set of measurements. All measurements were performed by one author (BL). The anterior cerebral artery was insonated anterior to the corpus callosum in parasagittal section through the anterior fontanelle. Optimal Doppler signal was obtained by simultaneously listening to the audio signal and observing the sonogram in real time. The CBFV were measured at an angle as close to zero as possible. Proucelot’s resistance index (RI=(S-D)/S) was calculated as the mean of five waveforms, where S and D are the maximum and the minimum blood flow velocities during one cardiac cycle.

LVO was calculated according to the following formula:

$$\text{LVO} = \text{aortic flow velocity integral} \cdot \text{aortic root area} \cdot \text{heart rate/birth weight (ml/min/kg)}^8$$

Aortic flow velocity curve was recorded from the suprasternal window using continuous wave Doppler. The beam was aligned in the ascending aorta. Maximal Doppler signal was obtained by simultaneously listening and observing the signal. The aortic flow velocity was integrated by tracing the area of the Doppler curve. Aortic flow velocity integral was averaged over five consecutive beats. The aortic root diameter was recorded from the parasternal long axis window at the level of the bases of the aortic leaflets in early or mid systole, from inner edge to inner edge. The diameter was averaged over five consecutive measurements. The aortic root area was calculated as: $$\pi \cdot (0.5 \text{ diameter})^2$$. The ductus was imaged from the suprasternal and the parasternal short axis windows using colour flow and Doppler.

To estimate intra-observer variability, 14 premature infants with a median gestational age of 29 weeks (range 26–36) and a birthweight of 1340 g (710–2560) were also studied during high frequency oscillation and conventional mechanical ventilation. All had RDS and were stable at the time of measurements. At 20 minute intervals, 10 infants were measured 13 times during high frequency oscillation and nine infants 12 times during conventional mechanical ventilation. No significant differences between the paired haemodynamic measurements (S, D, and LVO) were found either during high frequency oscillation or conventional mechanical ventilation (all P>0.40 by paired Wilcoxon rank sum test). The coefficients of repeatability\(^8\) for LVO were 56 ml/kg/minute during high frequency oscillation and 60 ml/kg/minute during conventional mechanical ventilation, and for median LVO 270 ml/kg/minute and 267 ml/kg/minute, respectively. They compare favourably with those reported by Fenton et al\(^8\) in infants whose cardiac output was determined by thermodilution and Doppler technique. For the anterior cerebral artery blood flow velocity, an analysis of variance showed that the SDs (expressed as percentage of mean replicate measurements) caused by intra-observer variability were 1.8 cm/second (7%) for S and 1.1 cm/second (18%) for D, respectively, during high frequency oscillation; during conventional mechanical ventilation, these were 2.1 cm/second (8%) for S and 0.6 cm/second (9%) for D, respectively. These levels of reproducibility agree with those reported by Winberg et al\(^8\).

Statistical analysis

Comparisons were made using the paired Wilcoxon rank sum test. A P value of <0.05 was considered significant. As we did not know in advance the magnitude of the haemodynamic changes, we recruited eight patients to detect with a power of 80% at a 0.05 significance level, a variation in CBFV or LVO as large as 1 SD of the mean change. Results are shown as median and range.

Results

Of the 10 infants recruited, six were studied at both T1 and T2, two only at T1, and two only at T2 leading to eight paired measurements at both study times. Postnatal age at T1 and T2 was 5 hours (3–12) and 40 hours (21–73), respectively. Time interval between measurements during both ventilation modes was 16 minutes (11–26) at T1 and 12 minutes (10–23) at T2. All patients tolerated surfactant administration and the ventilation mode changes well. No treatment changes were made during the paired measurements. Seven patients had a patent ductus arteriosus with left to right shunt at T1 and two at T2.

Table 1 shows the ventilation requirements and the arterial blood gases at T1 and T2. FIO\(_2\) and AAD02 were significantly higher with conventional mechanical ventilation than with high frequency oscillation at both study times. No significant changes in blood gases occurred between the two ventilation modes. However, at T1, PaCO\(_2\) was slightly higher on high.
frequency oscillation than on conventional mechanical ventilation.

Table 2 shows the haemodynamic effects of ventilation changes at T1 and T2. The only significant haemodynamic changes were a decrease in LVO between conventional mechanical ventilation and high frequency oscillation at T1, and an increase in LVO between high frequency oscillation and conventional mechanical ventilation at T2 (fig 1) in all patients. At T1, the decrease in LVO was 33 ml/min/kg (10–73). At T2, the increase in LVO was 29 ml/min/kg (11–76).

At T1 six out of eight patients showed a decrease in resistance index; one an increase and one no change at 1–00. The changes, however, did not reach significance (P=0.14). Using the data presented by Levene et al.12 and van Bel et al.,13 we corrected the RI for PaCO2 variations and their changes approached significance (P=0.0912 and P=0.0613 respectively). At T1 no significant change in lung inflation was detected radiologically between conventional mechanical ventilation and high frequency oscillation (nine posterior ribs (7–9) with conventional mechanical ventilation and 9 (8–10) with high frequency oscillation; P=0.4). At T2 all patients presented a slight degree of lung hyperinflation during high frequency oscillation (nine posterior ribs (9–10)) before being transferred to conventional mechanical ventilation.

Discussion

Although the number of subjects is small, our study shows that compared with conventional mechanical ventilation, high frequency oscillation significantly decreased LVO without affecting blood pressure and heart rate in premature infants with RDS. Furthermore, high frequency oscillation significantly improved oxygenation which agrees with previously reported findings.24 To our knowledge, no study has ever reported on LVO during high frequency oscillation in preterm infants. Animal studies on cardiac output during high frequency oscillation suggest that this does not affect cardiac output as long as lung hyperinflation is avoided.56 However, a study in cats showed that an increase in Paw during high frequency oscillation adversely affected cardiac output, but that it was better tolerated when lung compliance was reduced.7 A study of cardiac output during high frequency oscillation and conventional mechanical ventilation in one teenager and three toddlers with various very severe lung pathologies,14 showed no adverse effect of high frequency oscillation on cardiac output, although the Paw was higher than during conventional mechanical ventilation. However, only a small number of patients with heterogenous pathologies were studied, and their very low lung compliances might have dampened the negative side effect of Paw on cardiac output.
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The mechanisms by which PaO₂ affects cardiovascular function have been well studied. An increase in PaO₂ causes various degrees of increase in alveolar volume, depending on the pulmonary compliance. Alveolar expansion can increase pulmonary vascular resistance and therefore diminish venous return. The right ventricular stroke volume is decreased, hence the reduction in cardiac output.

At T1 and at T2, all the patients presented a lower LVO with high frequency oscillation than with conventional mechanical ventilation and all the Paw were lower during conventional mechanical ventilation. All our patients had adequate intravascular volumes at the times of measurements as judged by blood pressure, heart rate, and clinical examination. Furthermore, two out of eight patients were receiving dobutamine infusion at T1 and all eight patients at T2. Dobutamine is a potent inotropic agent in premature infants with RDS. We therefore postulate that the negative effect on LVO during high frequency oscillation was due to an increase in pulmonary vascular resistance, leading to a decrease in right ventricular stroke volume.

Human adult and animal studies showed that a rise in PaCO₂ resulted in a net increase in cardiac output. In preterm infants a study suggested that a 1 KPa increase in PaCO₂ led to a 10% decrease in stroke volume. However, for our group as a whole, the decrease in LVO between conventional mechanical ventilation and high frequency oscillation at T1 was too important to be related only to the slight PaCO₂ increase.

Although we followed a 'high volume' strategy with a potential risk of alveolar overdistention, our patients had similar lung inflation with high frequency oscillation and conventional mechanical ventilation at T1. We therefore believe that the drop in LVO with high frequency oscillation at T1 was not attributable to hyperinflation. At T2, however, all patients presented various degrees of lung hyperinflation with high frequency oscillation and this increased lung volume probably adversely affected LVO.

Animal studies comparing cardiac output and organ blood flow between high frequency oscillation and conventional mechanical ventilation at similar or different Paw did not show any significant change in cerebral blood flow. Only one study evaluated the effects of high frequency oscillation on CBFV in premature infants. It showed that this was significantly lower in the anterior cerebral artery with high frequency oscillation than with conventional mechanical ventilation. However, the significantly lower PaCO₂ during high frequency oscillation might have induced the observed CBFV changes. We could not show any significant change in CBFV with high frequency oscillation compared with conventional mechanical ventilation. As we measured the CBFV over a five cardiac cycle period, any cycling might have limited the value of our CBFV comparison. However, as both systolic and diastolic velocities seem to vary to the same degree, cycling variability may not be reflected in the RI. At T1, six of eight patients showed a decrease in RI with high frequency oscillation. After correcting for PaCO₂ changes, the overall variation in RI was close to significance and a type 2 error cannot be excluded. This drop in RI with high frequency oscillation might be the result of a cerebral vasodilatation in response to a falling LVO and a reduced cerebral perfusion pressure, as shown in ventilated newborn piglets.

By analogy we would have expected a similar but opposite change in RI at T2 during transfer from high frequency oscillation to conventional mechanical ventilation. We have no definite explanation for the lack of such change at T2.

In our patients we showed that the LVO during high frequency oscillation was 14% to 18% lower than during conventional mechanical ventilation. Systemic vasodilation must have occurred at T1 during high frequency oscillation to maintain blood pressure at the same level as that with conventional mechanical ventilation. This increase in systemic vascular resistance might have decreased blood flow to organs other than the brain as shown by Mirro et al in piglets during conventional mechanical ventilation at variable Paw. At T2, the rise in LVO during transfer from high frequency oscillation to conventional mechanical ventilation might have led to an increase in organ blood flow. Changes in organ blood flow are critical in premature infants. They were related to periventricular haemorrhage or necrotising enterocolitis. The correlations between high frequency oscillation, LVO, CBFV and organ blood flow merit further evaluation.

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Arch Dis Child Fetal Neonatal Ed 1996 74: F172-F176
doi: 10.1136/fn.74.3.F172

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