Prophylaxis was more effective than rescue treatment. The onset of action is more rapid with animal derived surfactants. The animal derived surfactants comprise Curosurf, which is an extract of pig lung mince and given in a volume of 1.25 to 2.5 ml/kg, and Survanta, an extract of cow lung mince with three added lipids, given as a dose of 4 ml/kg. These surfactants contain the apoproteins SP-B and SP-C, and these are thought to enhance their properties. The synthetic surfactants comprise Exosurf, a mixture of the key phospholipid DPPC, hexadecanol, and tyloxapol, given in a volume of 5 ml/kg, and ALEC, a mixture of DPPC and phosphatidylglycerol, given as a dose of 1.2 ml, regardless of size.

Fifteen studies have compared different surfactants, seven of which were of suitable quality for meta-analysis. Six of these trials compared Survanta and Exosurf; the other trial compared Infasurf and Exosurf. The meta-analyses support a significant reduction in the risk of pneumothorax (0.69 CI 0.57 to 0.85), and showed a non-significant trend towards reduced mortality. Soll’s conclusion was that: “on clinical grounds, natural surfactant extracts were the more desirable choice.” The onset of action is more rapid with animal derived surfactants than with artificial surfactants. This means that the babies treated with these surfactants need to be carefully monitored and their ventilator settings adjusted appropriately. Concern has been expressed because rapid effects lead to temporary changes in cerebral blood flow velocity and EEG recordings. There is currently no evidence to suggest that this more rapid onset of action has any deleterious effects.

UPPER AND LOWER GESTATIONAL AGE LIMITS FOR SURFACTANT TREATMENT

Although the data are limited due to small numbers in trials, the data that can be extracted show benefit for the smallest of babies. Gestational or weight limits for
either giving or withholding surfactant are not helpful, and may deprive babies who could benefit from surfactant use.

**INDICATIONS FOR SURFACTANT USE IN OTHER CONDITIONS**

There have been several small trials on the use of surfactant outside the classic indication of RDS, such as sepsis, pulmonary haemorrhage, or meconium aspiration syndrome. All of these show some benefits. Most convincing to date is the trial using beractant (Survanta) in ventilated cases of meconium aspiration syndrome. Using a dose of 150 mg or 6 ml/kg, the trial showed improved oxygenation, and reduced air leaks, severity of pulmonary morbidity, and shorter inpatient stay among term infants.

### Inhaled nitric oxide

The discovery that endothelium derived relaxing factor (EDRF) was in fact a gas, NO, has revolutionised thinking about several diseases. The NO pathway seems to have a crucial role in the vasoactivity of the pulmonary vascular bed. PPHN may be in part due to a deficiency of or resistance to NO, and the endothelial cellular defects may represent a final common pathway for the diverse causes for PPHN.

Once suitable delivery systems were developed, inhaled NO was first tested in neonatal pilot studies and then full scale randomised controlled trials. The results of a large trial on the use of inhaled NO in full term infants with congenital diaphragmatic hernias and concluded that it did not reduce the need for ECMO or the incidence of death. Fifty three infants were enrolled into this trial; in spite of aggressive management, including ventilation, alkaloasis, surfactant, inhaled NO and ECMO, the mortality was almost 50% in both arms of the trial. The place of inhaled NO in treating premature babies who have pulmonary hypertension as a component of their RDS, also remains uncertain. One four way prospective trial comparing dexamethasone and inhaled NO, concluded that both or neither failed to show any benefit for any group.

### Prophylaxis against group B streptococcal infection

Group B streptococcus is the leading cause of serious neonatal infection. Infants who are infected with it can require prolonged hospital stay, and a third of the survivors sustain permanent sequelae. There is no doubt that selective intrapartum prophylaxis is effective, and this has been confirmed by a meta-analysis which showed a 30-fold reduction in group B streptococcal disease. More than a decade has passed since the first clinical trial showed the effectiveness of intrapartum antibiotic prophylaxis, but still prevention strategies have not been implemented widely or consistently, and the incidence of neonatal group B streptococcal infection has not declined.

The alternative strategies currently available and the difficult choices they present were recently thoroughly reviewed by Isaacs. The choice between inactivity (treating only symptomatic babies), universal screening, and treating on the basis of risk factors alone remains. Isaacs concluded that preventive measures may not be justified in terms of cost effectiveness when the incidence was below 0.6 per 1000, but there are no accurate incidence figures for most of the UK. In the Northern Region the incidence was recently estimated as at least 1 case per 1000 deliveries whereas in Oxford it was 0.5 per 1000. Confirmed cases of early onset group B streptococcal infection reported to the Public Health Laboratory Service this year suggest an incidence of exactly 0.6 per 1000, uncomfortably near a level at which screening ought to be considered for the UK population.

### Extra corporeal membrane oxygenation

ECMO involves oxygenating blood outside the body and providing cardiovascular support, using complex machinery resembling that used for cardiopulmonary bypass. ECMO can be used only in babies weighing more than 2 kg, and candidates for this treatment usually have PPHN or meconium aspiration syndrome. ECMO has now been used on over 11000 infants worldwide, with 80% survival reported. Traditional ECMO uses two large gauge catheters, usually one in the jugular vein and one in the carotid artery. This form of veno-arterial ECMO involves permanent sacrifice of one carotid artery, and more recently veno-venous ECMO has become more popular. While babies are on ECMO the ventilator is reduced to “rest” settings, allowing the lungs to recover without barotrauma. In general, about two weeks is the maximum time for which babies can safely be sustained on ECMO.

The UK collaborative ECMO trial enrolled 185 infants in two years; 30 of 93 infants allocated ECMO died com-
pared with 54 of 92 allocated conventional care. Two thirds of the cases were enrolled in the first 12 months of the trial, in 1993–94. Infants with congenital diaphragmatic hernia are an important subgroup for whom no benefit from ECMO has yet been shown, but small numbers of cases preclude a meta-analysis and some centres claim good results. Only four of the 35 infants with CDH in the UK trial survived, and all were in the ECMO arm of the trial. Concern about quality of survival remains. Forty five of 62 (73%) babies treated with ECMO in the UK trial seemed to be normal at one year of follow up. The international registry records that 17% of infants treated with ECMO revealed a 17% prevalence of major disability, with concern about difficult behaviour and academic failure in a higher percentage.  

Deafness seems to be a particular risk.

The ECMO trial began just about the time that the first babies were being treated with NO, and the numbers of neonates being offered ECMO each year in the UK is currently declining. There seems little doubt that the facility needs to be available in the UK, at a few specifically designated centres that can maintain levels of expertise because they are caring for enough cases each year. Because of the small number of ECMO centres, the difficulty for the future will be in identifying and referring appropriate cases in time. An oxygen index of 40 or above predicted 60% mortality in the ECMO trial, and if this index does not rapidly fall with NO and/or HFOV, too much time should not be wasted in considering ECMO. Ten of the 30 deaths in the ECMO arm of the UK trial occurred among the 15 infants who were allocated to ECMO, but did not actually receive it.

High frequency oscillation ventilation

High frequency oscillatory ventilation alternately subjects the lungs to positive and negative pressure at very fast rates, usually about 10 Hz (10 cycles per second). Special equipment is required to achieve effective ventilation at such high frequencies, and of the three oscillators available in the UK, only the Sensormedics 3100/S100A has been used in randomised controlled trials. The Sensormedics is a dedicated oscillator. The other available oscillators are the Draeger Babylog 8000 and the SLE HV2000 ventilator. Oscillators are powerful tools, and there is no doubt that in “rescue” mode HFOV can save infants with severe RDS who have failed to respond to conventional ventilation and surfactant. HFOV is particularly effective in hypercapnia. What is least certain is the role of HFOV as the primary mode of ventilation in RDS in very small babies who have received antenatal steroids and postnatal surfactant. The Provo trial17 randomly allocated 125 babies with RDS at less than 35 weeks of gestation who had received surfactant. Those who were ventilated with HFOV fared better than those ventilated conventionally in the short term, with more survivors without chronic lung disease at 30 days. Although there has been some concern about the high number of babies still ventilated at a month (half the HFOV group and all the conventionally ventilated group), the incidence of ultrasound abnormalities and retinopathy of prematurity was the same. Even this large study only enrolled 21 babies with a birthweight of less than 1 kg. HFOV is not the same as high frequency jet ventilation (HFJV). This involves the delivery of a jet of gas directly into the trachea, and lacks an active expiratory phase. Recent evaluations from the USA suggest an excess of cystic periventricular leukomalacia (PVL) in survivors ventilated this way. The increased risk of PVL is perhaps due to hypocapnia.  

A meta-analysis of trials of high frequency ventilation revealed a higher incidence of intraventricular haemorrhage and PVL which disappeared if the results of the large HFi trial were excluded.15 HFOV is currently reserved for rescue treatment in most UK neonatal units, although a large MRC sponsored trial of the use of HFOV from birth in infants 26–29 weeks of gestation (the UKOS trial) is actively recruiting.


32 Marlow N. High frequency ventilation and respiratory distress syndrome: do we have an answer? *Arch Dis Child* 1998;78:F1–F2.


