LETTERS TO THE EDITOR

A survey of neonatal resuscitation training provided to general professional trainees at neonatal units in England and Wales

EDITORS,—The Royal College of Paediatrics and Child Health produced recommendations on teaching and training junior doctors involved in neonatal resuscitation.1 These state that all new staff members should receive an induction programme on neonatal resuscitation that includes practice with mannequins. All should also be familiarised with resuscitation equipment on labour wards and be appraised before being permitted to operate alone. Regular updates on neonatal resuscitation training are also recommended and ideally all junior staff’s participation in resuscitation procedures should be formalised and documented.

We attempted to assess by telephone questionnaire the current position of neonatal resuscitation training in England and Wales, using a questionnaire of general professional trainees (GPTs). The questionnaire covered key areas within the college recommendations. An attempt within the questionnaire was made to assess the previous neonatal experience of the GPT respondents. A single GPT in each unit was interviewed by telephone. The survey was conducted from October 1999 to January 2000 (the latter part of most attachments). Two hundred and nine units were identified as units with paediatric Senior House Officer cover. One hundred and seventy seven GPTs successfully completed the questionnaire (85%). None contacted were unwilling to participate (some were busy while on duty and others did not answer within three attempts).

Most units (94%) provide a formal theoretical session on neonatal resuscitation during induction (table 1), 86% of units provided practical skills training at induction including endotracheal intubation. Training on practical skills of attaining vascular access appeared to be common in regional units (84% v 39%) along with higher chance of formal appraisal before being left alone for resuscitation (47% v 22%). District general hospitals provided more encouragement to attend courses (73% v 53%) but had fewer trainees with previous experience.

Resuscitation training includes induction with didactic teaching, practical skills training, and ongoing assessment of performance. In an ever changing situation of GPT education, our results need to be interpreted with caution but nevertheless, we believe, reflect current concerns about experience and education offered to trainees, which will hopefully be addressed by the Neonatal Life Support Course.

This study indicates that some, but not all, units meet the standards encompassed within the Royal College guidelines. In 1993 Barrie also recorded considerable variation in the delivery of such training in a Northwest area.1 This study reveals an identical problem right across England and Wales.

We hope the new neonatal life support will address many of the issues raised within the area of training of GPTs, but universal and mandatory attendance of this is necessary if standards of newborn resuscitation are not to vary unacceptably across England and Wales. We believe this aspect of care should be the subject of quality assurance when assessing the capacity of individual units to provide care in the future.

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Table 1 Results of the telephone survey of General Professional Trainees in England and Wales on neonatal resuscitation training provided to them in their neonatal units

<table>
<thead>
<tr>
<th>Questions</th>
<th>Total units (n=177)</th>
<th>Regional units (n=98)</th>
<th>DGH units (n=34)</th>
<th>Previously experienced (n=122)</th>
<th>Previously inexperienced (n=55)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Formal induction</td>
<td>167 (94)</td>
<td>18 (95)</td>
<td>149 (94)</td>
<td>50 (91)</td>
<td>117 (96)</td>
</tr>
<tr>
<td>Intubation skills</td>
<td>153 (86)</td>
<td>22 (85)</td>
<td>135 (85)</td>
<td>46 (84)</td>
<td>108 (78)</td>
</tr>
<tr>
<td>UVC insertion</td>
<td>77 (44)</td>
<td>16 (94)</td>
<td>61 (39)</td>
<td>26 (47)</td>
<td>35 (42)</td>
</tr>
<tr>
<td>Resuscitation</td>
<td>159 (90)</td>
<td>16 (84)</td>
<td>143 (91)</td>
<td>48 (87)</td>
<td>111 (91)</td>
</tr>
<tr>
<td>24 hr registrar cover</td>
<td>155 (88)</td>
<td>19 (100)</td>
<td>136 (86)</td>
<td>43 (78)</td>
<td>112 (92)</td>
</tr>
<tr>
<td>Awaiting postnatal unit successful</td>
<td>54 (31)</td>
<td>6 (32)</td>
<td>45 (29)</td>
<td>21 (38)</td>
<td>33 (27)</td>
</tr>
<tr>
<td>formal intubation</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Formal appraisal before left</td>
<td>44 (25)</td>
<td>9 (47)</td>
<td>35 (22)</td>
<td>16 (29)</td>
<td>18 (23)</td>
</tr>
<tr>
<td>supervised</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Ongoing training</td>
<td>80 (45)</td>
<td>11 (58)</td>
<td>69 (44)</td>
<td>29 (53)</td>
<td>51 (42)</td>
</tr>
<tr>
<td>Courses attended/encouraged</td>
<td>126 (71)</td>
<td>10 (53)</td>
<td>116 (73)</td>
<td>46 (84)</td>
<td>80 (66)</td>
</tr>
</tbody>
</table>


A rare cause of respiratory distress

EDITORS,—We would like to describe a case of respiratory distress associated with haemodynamic compromise. This presented as a clinical scenario, and the underlying cause is usually evident. Occasionally the cause is rare, and we report a case of respiratory distress caused by bilateral adrenal agenesis.

A baby girl was born at 37 weeks’ gestation to a diabetic mother. She remained well for the first six days, but on day seven she was transferred to our unit because of severe respiratory distress requiring intubation and ventilation. A chest radiograph showed a small pneumothorax and changes consistent with surfactant deficiency. Serum sodium concentration was 125 mmol/l, and serum potassium concentration 7 mmol/l. A septic screen and cardiac echocardiography were normal.

The combination of both fetal hyperinsulinaemia and glucocorticoid deficiency would considerably impair surfactant production and may explain the severity of our patient’s respiratory distress.

Both respiratory distress and electrolyte imbalances are common in the neonatal unit. Therefore when the diagnosis is not obvious, the search for causative factors must be broadened to ensure that rare but clinically important diagnoses are not missed.

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Disseminated candidiasis after steroid treatment for early neonatal hypotension

EDITOR,—We are concerned that intravenous steroids are increasingly being used for the treatment of early neonatal hypotension without proper evaluation. We recently treated three newborn premature infants who had received a six day reducing course of intravenous hydrocortisone (starting at 12.5 mg/kg/day in divided doses) from day 1 of age for the treatment of hypotension. They were otherwise stable, apart from one infant who had a patent ductus arteriosus. All developed severe systemic candidiasis and needed enterocolitis by 14 days of age, with only one infant surviving.

Reported adverse effects of steroid treatment for neonatal hypotension include fungal infection,1 hyperglycaemia, septal and ventricular hypertrophy,2 and hypertension.3 There are also well described short term complications of early steroid use for chronic lung disease including gastrointestinal haemorrhage and intestinal perforation.4 Long term adverse effects in these neonates include an increased risk of cerebral palsy and developmental delay.5 Any benefit of treating neo-

tnatal hypotension with intravenous steroids would have to be substantial in order to outweigh these risks.

It is not clear that steroids confer any advantage over appropriate inotropic treat-
ment of neonatal hypotension. A number of small studies have reported increases in blood pressure when steroids are used. The largest study of 40 infants randomised to receive inotropes or hydrocortisone6 showed 81% survival in the inotropic treated group compared with 45% survival in the hydrocortisone group with 100% success with dopamine treatment (at rates of 5–20 µg/kg/min). Inotropes allow more accurate titration of drug dose to response and are known to be effective in maintaining blood pressure. If dopamine is insufficient, adrenalin-
eline (epinephrine) or noradrenaline (nor-
ephinephrine) should be added for further inotropic support.

Further studies are needed to understand the role of steroids in the newborn premature infant before their use for the treatment of hypotension becomes universal, and at present any advantages of steroid treatment over escalation of inotropic support are outweighed by adverse effects. This is of par-


ticular importance in the light of the increasing evidence7 that the benefits of steroid treatment of chronic lung disease in newborn premature infants may not outweigh the adverse effects. Consideration must be given to limiting the use of steroids for neonatal hypotension to situations where other proven methods of cardiovascular support have failed.

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1 Bottas CM, Kurtal I, Young SM, et al. Dissemi-


2 Gainsmaier RE, Pohlandt F. Single-dose dexam-


3 Bourchier D, Weston PJ. Randomised control-

led trial of dopamine compared with hydro-

4 Soll RF, for the Vermont Oxford Network Ster-

oid Study Group. Early postnatal dexamethas-


Serum ferritin level in neonatal fulminant liver failure

EDITOR,—Elevated serum ferritin is a non-
specific indicator of severe hepatocyte in-
jury.7 Neonatal haemochromatosis (NH) is an uncommon disorder of neonatal iron stor-
ge, which often requires transplantation.

1 Serum ferritin levels are considerably elevated in NH,1 but differentiating NH from other forms of neonatal fulminant hepatic failure (FFH), in which liver trans-

plantation is contraindicated, can be very misleading.1 We reviewed the case records of all neo-
natal FFH (onset of liver failure less than 28 days of age) seen in this unit from 1990 to 1999 to determine whether causes of neonatal FFH can lead to an elevated serum ferritin level. Sixteen cases of neonatal FFH were seen, with a median onset of symptoms at 9 days (range 1–18). Eight had NH, confirmed either by histology showing paren-

chymal haemosiderin deposition in liver or buccal salivary tissue, and/or a positive family history.1 The remaining eight cases had the following causes: mitochondrial disorders (two); galactosaemia, Escherichia coli sepsis, Herpes simplex virus type 1 and 2 hepatitis (one each); undetermined (two). All had raised serum ferritin levels (range 1000–

217 000 µg/l). The median serum ferritin level for the patients with NH and the miscellaneous group were 15 000 µg/l (range 217 000). There was no significant difference between the mean serum ferritin level of the NH group and that of the miscel-

laneous group (40 916 v 5409 µg/l, p = 0.08).

This observation confirms that the serum ferritin level is raised as the result of severe hepatic injury in newborn infants, irrespective of the cause.1 In neonatal liver failure, other diagnostic methods, such as demonstration of extrahaemorrhagic iron—for ex-

ample, in lip salivary glands—or magnetic resonance imaging should be used to confirm or refute the diagnosis of NH.1

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2 Cotler SJ, Bronner MP, Press RD, et al. End-stage liver disease without hemochroma-
A rare cause of respiratory distress

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