Reducing medication errors in the neonatal intensive care unit

J H Simpson, R Lynch, J Grant, L Alroomi

Background: Medication errors are common in the neonatal intensive care unit (NICU). Various strategies to reduce errors have been described in adult and paediatric patients but there are few published data on their effect in the NICU.

Aim: To describe the medication errors occurring within an NICU, and assess the impact of a combined risk management/clinical pharmacist led education programme on these errors.

Methods: Medication errors were identified prospectively over one year by critical incident reporting. Four months into the study, a pharmacist led education programme was instituted. This involved a daily, cot side, pharmacist led review of medication orders. Each new member of pharmacy, nursing, or medical staff was also required to successfully complete a series of dose calculations. In addition, a risk management approach was used to make several changes in practice during the study period.

Results: A total of 105 errors were identified, four serious, 45 potentially serious, and 56 minor. The four serious errors included two tenfold dose miscalculations. Most (71%) of the errors were due to poor prescribing. After the introduction of our interventions, monthly medication errors fell from a mean (SD) of 24.1 (1.7) per 1000 neonatal activity days to 5.1 (3.6) per 1000 days (p < 0.001) in the following three months. The subsequent change over of junior medical staff was associated with a significant increase in medication errors to 12.2 (3.6) per 1000 neonatal activity days (p = 0.037). However, the number remained significantly less than before our interventions (p < 0.001). Three serious errors occurred in the first four months compared with one in the second eight month period, the latter corresponding to the six monthly change over of junior medical staff.

Conclusions: Medication errors are common in NICUs. Fortunately, actual harm to an infant is rare. Interventions to reduce errors, particularly within the context of a risk management programme, are effective.
medication errors. Some of these changes coincided with the introduction of the pharmacist led review of medication orders—for example, the abolition of prescription writing during ward rounds—and others were introduced after this. These changes included writing the day as well as the date for medicines prescribed at unusual intervals (for example, gentamicin), provision of a single concentration of morphine for reconstitution on the NICU, and, where possible, reconstitution of intravenous medicines in the pharmacy. Insulin and heparin were stored in separate fridges to avoid confusion during reconstitution, and dedicated infusion pumps, preset for heparin, were used only for arterial line infusions. Throughout the study period, monthly neonatal activity days were monitored.

Statistical analysis
Mean (SD) medication errors per 1000 neonatal activity days were calculated. These were compared using Student’s t test. Statistical significance was taken at p < 0.05.

RESULTS
A total of 105 errors were identified, four serious, 45 potentially serious, and 56 minor. Most (75%) were reported by the clinical pharmacist, but all four serious errors were reported by medical or nursing staff involved in the error. Parenteral medicines were involved in 63 errors (60%), oral medicines in 41 errors (39%), and topical medicines in one error. Table 1 lists the parenteral medicines involved.

Four serious errors included two involving 10-fold dose miscalculations. In one case, an infant who received an opiate overdose required naloxone, and in the second an antibiotic overdose was given which required no specific treatment. The third serious error involved administration of an antibiotic to the wrong baby. There was no direct harm to the infant. The final serious error was a “near miss” situation involving the use of insulin rather than heparin to reconstitute a bag of “heparinised” saline. This error was identified before administration by a second member of staff checking the vial used for reconstitution.

Most (71%) of the medication errors were due to poor prescribing. The most common example of this was an incorrect dose (unexplained deviation of > 10% from the neonatal unit formulary), which occurred in 37 cases. An incorrect dose interval was found in 19 errors; this was particularly common with gentamicin. Other examples of poor prescribing included incomplete prescriptions (14) and incorrect units (5). Thirty errors (29%) were due to administration problems. In 16 cases, the medication was not documented as given, reflecting either poor communication between staff or more often poor documentation of administration. In eight errors, the medication was given incorrectly, either by the wrong route or too quickly, and in six cases administration was delayed by more than two hours, generally reflecting the intensity of work in the unit.

Following the introduction of our interventions after four months, medication errors fell from 24.1 (1.7) per 1000 neonatal activity days to 5.1 (3.6) per 1000 days (p < 0.001) in the following three months (table 2). The change over of junior medical staff in August was associated with a significant increase in medication errors to 12.2 (3.6) per 1000 neonatal activity days (p = 0.037). However, the number remained significantly less than before our interventions (p < 0.001; table 2). Three serious errors occurred in the first four months compared with one in the second eight month period, the latter corresponding to the six monthly change over of junior medical staff.

DISCUSSION
In common with other authors,11 we found that medication errors occur regularly in our NICU. Fortunately, actual harm to an infant is rare. The types of drugs used, many of which have a narrow therapeutic margin, and the ease of miscalculation during reconstitution from stock solutions based on adult concentrations mean that the potential for harm from these errors is great.12,13

We have shown that interventions to improve staff education and awareness of errors are effective in reducing such errors, although within the context of our overall risk management approach, it is difficult to quantify the proportion of errors reduced by any one change in practice.

Inexperience is a particular risk factor for medication errors.2 This is supported by our own findings and those of others that new staff are more likely to make errors.14 This is compounded by the fact that prescriptions are often written by the most junior doctors who may be unfamiliar with the medicine.2 The importance of the paediatric clinical pharmacist in monitoring drug treatment and preventing medication errors is well established.4,15 Fortescue et al17 recently showed that clinical pharmacist monitoring of orders might prevent 58% of all errors and 72% of potentially harmful errors, and that improved doctor-pharmacist communication might prevent 47.4% of errors.18 We found that close liaison with a ward based clinical pharmacist is an effective way of reducing errors. We effectively extended the role of the pharmacist to include real time staff feedback and education, both informally while reviewing medication orders and formally as a series of dose calculations. Calculation errors, particularly 10-fold errors, can be potentially fatal and any intervention to reduce these is important.12

Intensity of workload is recognised to be another risk factor for medication errors.2 We used neonatal activity days

### Table 1
<table>
<thead>
<tr>
<th>Drug involved</th>
<th>N (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Gentamicin/netilmicin</td>
<td>21 (33)</td>
</tr>
<tr>
<td>Benzyl penicillin</td>
<td>10 (16)</td>
</tr>
<tr>
<td>Vancomycin</td>
<td>9 (14)</td>
</tr>
<tr>
<td>Morphine</td>
<td>6 (10)</td>
</tr>
<tr>
<td>Insulin</td>
<td>4 (6)</td>
</tr>
<tr>
<td>Aminophylline</td>
<td>3 (5)</td>
</tr>
<tr>
<td>Immunisations</td>
<td>3 (5)</td>
</tr>
<tr>
<td>Tazocin</td>
<td>2 (3)</td>
</tr>
<tr>
<td>Other</td>
<td>5 (8)</td>
</tr>
</tbody>
</table>

### Table 2
<table>
<thead>
<tr>
<th>Month</th>
<th>Intensive care activity (days)</th>
<th>Special care activity (days)</th>
<th>Total activity (days)</th>
<th>Medication errors per 1000 neonatal activity days</th>
</tr>
</thead>
<tbody>
<tr>
<td>January</td>
<td>234</td>
<td>398</td>
<td>634</td>
<td>23.7</td>
</tr>
<tr>
<td>New junior medical staff started</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>February</td>
<td>234</td>
<td>435</td>
<td>669</td>
<td>23.9</td>
</tr>
<tr>
<td>March</td>
<td>190</td>
<td>528</td>
<td>718</td>
<td>26.5</td>
</tr>
<tr>
<td>April</td>
<td>223</td>
<td>359</td>
<td>582</td>
<td>22.3</td>
</tr>
<tr>
<td>Interventions started</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>May</td>
<td>278</td>
<td>283</td>
<td>561</td>
<td>8.9</td>
</tr>
<tr>
<td>June</td>
<td>205</td>
<td>419</td>
<td>624</td>
<td>4.8</td>
</tr>
<tr>
<td>July</td>
<td>189</td>
<td>428</td>
<td>617</td>
<td>1.6</td>
</tr>
<tr>
<td>New junior medical staff started</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>August</td>
<td>197</td>
<td>391</td>
<td>588</td>
<td>13.6</td>
</tr>
<tr>
<td>September</td>
<td>159</td>
<td>391</td>
<td>508</td>
<td>17.7</td>
</tr>
<tr>
<td>October</td>
<td>230</td>
<td>316</td>
<td>546</td>
<td>11.0</td>
</tr>
<tr>
<td>November</td>
<td>190</td>
<td>383</td>
<td>573</td>
<td>8.7</td>
</tr>
<tr>
<td>December</td>
<td>137</td>
<td>375</td>
<td>512</td>
<td>9.8</td>
</tr>
</tbody>
</table>
as a surrogate marker of workload to correct for the potential confounding effect of variations in activity. We accept, however, that a description of medication errors per number of prescriptions written would provide more accurate information on the impact of workload.

Longitudinal monitoring enabled us to identify sources of recurrent error and facilitate several changes in our practice. This technique of continuous critical incident monitoring and quality improvement through system change has been described previously in an NICU. In this prospective study of 284 medication related critical incidents, 46 were followed by system changes. Knowledge acquired in this way, even from apparently minor errors, is an important resource in the prevention of future errors.

Prescribers are human and therefore mistakes will be made. Recognition of these errors is the first step in their prevention. It is important that prescribers are aware of their own vulnerability and that we all learn from our own mistakes and those of others. Increased awareness of error is an important preventive tool. In addition, we feel that ward based input from a clinical pharmacist, particularly within the context of a risk management programme, is important in the NICU both to monitor medication orders and to provide education to frequently changing members of staff.

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