Model to identify potentially preventable cerebral palsy of intrapartum origin

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Abstract
A six stage model was applied to a geographically defined population of 210 singleton children born at term who had a diagnosis of cerebral palsy at 5 years of age. Thirty five children were identified as those most likely to have cerebral palsy of intrapartum origin; in 26 of these there was evidence of suboptimal care.

It is suggested that this simple model should be tested on populations of children with cerebral palsy and the underlying principles used when considering the likely cause of cerebral palsy in individual children.

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There are some children with cerebral palsy in whom a different pattern of care during labour and delivery would have changed the outcome.1-4 It is important to identify these children for two reasons. First, it may be possible to implement changes in management which would prevent further similar cases. Second, some parents of disabled children resort to civil court action, seeking damages for alleged negligence around the time of birth. Correct identification of the children who seem to suffer the effects of suboptimal obstetric care is a part of this litigation process.

Based on a review of existing epidemiological evidence of the current understanding of the origin of cerebral palsy, we developed a model which can be used to identify those children with cerebral palsy who are most likely to have had a preventable intrapartum event.5-24 A sequence of six questions is asked. The response ‘No’ excludes the child from further questions. We believe that these children remaining at the end of the sequence are most likely to have had an obstetrically preventable cause of cerebral palsy. The model was tested on a geographically defined population of children with cerebral palsy.

Method
The model consisted of the six questions shown (figure). This applied to 210 children on a regional cerebral palsy register who were at least 5 years old and who were singletons, born at term between 1984 and 1987. Details of the register are discussed elsewhere.25

Results
The number of children excluded at each step in the model is shown in the figure.

QUESTIONS
(1) Postnatal cause Twenty seven children were identified as having a postnatal cause for cerebral palsy. The causes were meningitis in 12, trauma in six, cerebrovascular accidents in three, postoperative anoxia in three and miscellaneous other events such as near-drowning in three.

(2) Prenatal cause Forty two children had an identifiable prenatal origin to their cerebral palsy. These included cerebral malformations (n=10), cerebral vascular malformations (n=7), metabolic and chromosomal causes (n=7). Six had recognised syndromes, three had an intrauterine infection, and one child had a congenital ependymal tumour.

(3) Neonatal neurological dysfunction was present in 54 of the remaining 141 children. We thought it likely that among the 87 children excluded at this stage, some may have had an unrecognised prenatal origin for their cerebral palsy, but there was insufficient information available to support this.

(4) Depression at birth Forty six of the 54 babies with abnormal neonatal neurological signs had an Apgar score of 6 or less at one minute after birth. The remaining eight babies with abnormal neonatal neurological dysfunction had an Apgar score of more than 6 at one minute after birth. These eight babies had seizures during the first seven days of life but did not show any signs of altered levels of consciousness and there was no evidence of other organ ischaemia such as renal or myocardial dysfunction.

(5) Intrapartum fetal distress Of the 46 babies remaining at this stage, 35 had experienced an adverse event during labour or shortly before delivery. Twenty five mothers had an ominous cardiocograph reading in labour, including one with a ruptured uterus and three mothers who were delivered by emergency Caesarean section following failed forceps delivery. Three babies were delivered by emergency Caesarean section in the absence of labour. Of these, two were delivered because of an ominous prelabour cardiocograph and one was delivered at the time of fatal maternal sub-arachnoid haemorrhage. Of the seven remaining babies, three had evidence of fetal bradycardia on intermittent auscultation; one had a difficult rotational forceps delivery; one was a breech presentation diagnosed during labour at the time of delivery; in one there was a 20 minute delay in the delivery of the head of the infant at elective Caesarean section; and in the last case...
1. Has a postnatal cause for cerebral palsy been excluded?
   An event or illness which appears to have damaged the developing brain after the age of 28 days in a child previously considered to be neurologically normal.

2. Has a prenatal cause for cerebral palsy been excluded?
   An associated major congenital anomaly, strong family history of cerebral palsy, a chromosomal anomaly or evidence of brain malformation on neuroimaging.

3. Was there evidence of neurological dysfunction in the first week after birth?
   To include babies with seizures, tone changes and altered levels of consciousness.

4. Was the baby depressed at birth?
   To include babies with an Apgar score of 6 or less at one minute after birth.

5. Was there evidence of fetal distress or an obstetric emergency?
   To include an ominous cardiotocograph, definite evidence of clinical fetal distress (particularly bradycardia) on the obstetric record, or an acute emergency such as scar rupture.

6. Was there evidence of suboptimal intrapartum care?
   Using predefined standards of care.

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Attempts at delivery were made before full dilatation.

Eleven infants were excluded at this stage as no abnormality could be identified. Three mothers had continuous electronic fetal monitoring and in all cases this was normal. Two infants were delivered by elective Caesarean section, in one because there had been a previous Caesarean section and in the other because of a history of infertility in combination with unsuitability for induction of labour. The remaining six infants were not monitored but there was no evidence from intermittent auscultation that there had been any fetal heart rate abnormality during labour. In one case delivery was by Caesarean section following failure of a planned trial of forceps, but there was no evidence of fetal distress.

(6) Suboptimal intrapartum care
Thirty five babies remained in the model at this stage. Using predefined criteria for care, 26 of the 35 had evidence of suboptimal intrapartum care. In the nine babies excluded at this stage there was no evidence of a failure to respond to fetal distress within times specified in the predefined criteria for care.

Discussion
This model was developed using hypotheses about the origins of cerebral palsy derived from previous epidemiological studies. When applying the model, however, classification at each step in the model is based on the information available in the medical notes of each child. Misclassification seemed particularly likely to occur in three areas. First, although a postnatal cause of cerebral palsy may be readily identified from medical notes, cerebral palsy of prenatal origin is less easy to diagnose. It is likely, therefore, that a number of children are misclassified at stage 2. Most children have not had neuroimaging which might identify cerebral structural abnormalities either during the neonatal period or later childhood. We suggest that if more detailed investigations, including neuroimaging, were offered to children with cerebral palsy of uncertain origin, the size of the group of children with a
prenatal cause of cerebral palsy would be clearer.

Second, misclassification at stages 3 and 4 can occur if seizures, which are associated with hypoxic-ischaemic encephalopathy, are not distinguished from seizures due to other conditions. If, in the neonatal period neurological signs such as levels of consciousness, seizure activity, respiratory status, feeding and signs of other organ dysfunction were systematically recorded in a standard way, it would be easier to make this distinction both at the time of the events and when notes are reviewed retrospectively.

Third, in order to classify the children as accurately as possible at stages 5 and 6, detailed records of labour, including annotated cardiotocographs and other clinical particulars, are invaluable. These are not always available in medical and nursing records. Even in the absence of the currently accepted clinical signs of fetal distress, fetal compromise may still occur. Further research is needed on the reliability of clinical and physiological measures of fetal wellbeing.

One part of the model is not based on epidemiological evidence. The standards of care were derived from clinical consensus and many of these standards have never been subjected to rigorous testing or evaluation. This is an area of considerable debate both inside and outside the courts. We hope that the current interest in establishing standards of care based on evidence of effectiveness may help to clarify this part of the model.

As more information on the complexities of cerebral palsy becomes available, this model will need modification and refinement. In the meantime we would encourage others to test this model on other populations of children with cerebral palsy.