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Influence of obstetric management on outcome of extremely preterm growth retarded infants

A H P Schaap, H Wolf, H W Bruinse, A L den Ouden, H Smolders-de Haas, I van Erbruggen, P E Treffers

Abstract

Aim—To describe the long term outcome of extremely preterm growth retarded infants in relation to obstetric management and various perinatal events.

Methods—A cohort study was undertaken in two tertiary care centres with different obstetric management. All infants with fetal growth retardation due to placental insufficiency and resulting in fetal distress at 26 to 32 weeks of gestation, were included for the years 1984–89. Main outcome measures were impairment, disability, or handicap at 2 years corrected age and at school age (4 1/2 to 10 1/2 years).

Results—One hundred and twenty five infants were followed up until 2 years corrected age in the outpatient department; 114 (90%) were assessed at school age. Impairments were found in 37% and disabilities or handicaps in 9% of the assessed infants, with no difference between centres. All disabled or handicapped children had already been identified by 2 years corrected age.

Conclusions—Disability or handicap were related to neonatal complications (intracerebral haemorrhage or bronchopulmonary dysplasia) and not to obstetric variables, thus making antenatal prediction impossible. The incidence of disability or handicap in these growth retarded infants was comparable with that of other preterm infants.

(Arch Dis Child 1997;77:F95–F99)

Keywords: growth retardation; disability; handicap; obstetric variables

Suspected early fetal growth retardation due to placental insufficiency, and causing fetal distress, poses a clinical dilemma. Deciding when to deliver such babies involves balancing the consequences of delivery and attendant neonatal mortality or long term morbidity against the risk of a fetus compromised by nutrient and oxygen deprivation, and the risk of intrauterine death.

We have already reported a comparison of perinatal mortality and short term morbidity in two university hospitals with different management strategies (active or more conservative) for this selected group. Overall survival was significantly greater at the centre with an active management strategy (centre B). This resulted from a number of intrauterine deaths at centre A, after a decision to abstain from active inter-
Table 1. Perinatal outcome of the original study population

<table>
<thead>
<tr>
<th></th>
<th>Conservative management (Centre A) No (%)</th>
<th>Active management (Centre B) No (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Total</td>
<td>107 (100)</td>
<td>95 (100)</td>
</tr>
<tr>
<td>Antenatal mortality</td>
<td>34 (32)</td>
<td></td>
</tr>
<tr>
<td>Total liveborn</td>
<td>73 (68)</td>
<td>95 (100)</td>
</tr>
<tr>
<td>Postnatal mortality</td>
<td>14 (13)</td>
<td>27 (28)</td>
</tr>
<tr>
<td>Survivors</td>
<td>59* (55)</td>
<td>68* (72)</td>
</tr>
</tbody>
</table>

* P < 0.05.

FOLLOW UP BY QUESTIONNAIRE

In 1994 parents were interviewed by questionnaire. They were asked to assess whether their child had limitations in walking, hand function, hearing, vision, speech–language and comprehension, and whether there was any respiratory impairment. The severity of functional limitation and activity restriction was judged on a five point scale.

The items were categorised according to the International Classification of Impairments, Disabilities, and Handicaps (ICIDH) of the World Health Organisation (WHO). We regarded a child as impaired if he or she had a disturbance at organ level, or disabled if the impairment or multiplicity of impairments caused loss of function or activity.

We regarded a child as handicapped if he or she had disabilities that caused a social disadvantage. We considered handicap minor if it did not seriously interfere with everyday life and did not require extensive caretaking, and major if it did interfere with everyday life and if it caused dependency or institutionalisation.

When multiple disturbances were present we assigned the child to the most severe category.

To determine whether the outcome changed with time, the group was divided into two periods, one with a follow up of more than 7 1⁄2 years and one of less than 7 1⁄2 years. We compared the results of scoring at the age of 2 with those of the questionnaire.

Approval for the study was given by the research ethics committees of the two university hospitals.

STATISTICS

Data were analysed by computer using BMDP statistical software (Los Angeles, USA). Differences between categories were tested for significance using the χ² test with Yates’ correction. Significance was considered at P < 0.05. The influence of gestational age, birthweight, sex of the infant, centre, intracerebral haemorrhage (ICH), respiratory distress syndrome (RDS), bronchopulmonary dysplasia (BPD), sepsis and period of follow up on the incidence of disability/handicap was analysed using logistic regression analysis to address the interaction between these factors.

Results

COMPARISON BETWEEN CENTRE A AND B

Table 1 shows the total study population (n=202), with 34 antenatal deaths at centre A. One hundred and twenty seven children were discharged home alive and were included in the outpatient department follow up programme. Two children were lost to follow up. One hundred and twenty five were followed up at least until the corrected age of 2 (101 in the perinatal centres and 24 by a local paediatrician). No infant died after discharge home. Thirteen out of 127 survivors were lost to follow up at the time of the questionnaire (six from centre A and seven from centre B): two children could not be traced and 11 families did not respond despite repeated requests. Thus 114 (90%) of the surviving children were completely assessed.

The overall outcome of assessed children is shown in table 2. The percentages of disabilities and handicaps in the two centres were similar, but the overall adverse outcome (mortality and disability or handicap) differed significantly: 52/107 (49%) in centre A compared with 33/95 (35%) in centre B (P < 0.05).

OUTCOME OF QUESTIONNAIRE

Table 3 shows the outcome of the questionnaire for the areas of assessment. The highest incidence of handicap was found in neuromotor function, mental development, and language and speech development. Disorders of neuromotor function, mental development, and/or language and speech development were often found in the same child.

All children but one with a disability or handicap at the time of the questionnaire had already been identified at follow up at the age of 2. The exception was a child who was considered disabled according to the questionnaire but was only classified as suspected of disability at the age of 2. The severity of the disorder changed with time in six children. In three infants, handicap was reclassified from major to minor; in the other three a minor handicap was reclassified as disability. Most of these changes were in the area of neuromotor
low up at 2 years) were analysed using logistic regression. The analysis showed that the risk for disability or handicap was increased in the first study period (RR 30.7 (2.4-396)) and in infants with ICH (RR 38.7 (2.1-717)) or BPD (RR 11.2 (1.5-83)). Centre, birthweight, gestational age, sex, RDS or sepsis did not contribute significantly to the model.

Figure 2 shows that birthweight relates to mortality and not to disability or handicap.

Discussion

The need for a functional classification of handicap is increasingly being recognised. To optimise the comparability of outcome we adhered to the ICIDH of the WHO, which relates to the consequences of disease. Although school performance can be categorised according to this, we have not reported this item because of the differences in age at follow up and because the need for special education increases proportionately with age. Ninety eight percent of the children were followed up at the corrected age of 2 and 90% responded to the questionnaire. Tyson et al suggested that untraced survivors may have the same likelihood of handicap as those evaluated. Wariyar reported significantly higher disability rates among those infants who were more difficult to trace, in a 100% follow up study. Of the 13 non-responders to the questionnaire, only one child was identified at outpatients as having a minor handicap. This made no difference to the disability or handicap rate. The reason for being lost to follow up in our population was not related to the health status of the infant.

The use of a questionnaire could possibly have generated a source of bias. Parents might overreport or underreport disabilities and handicaps in their children. However, other studies support the assumption that most parents accurately assess their child’s current level of functioning. Our results show a similar disability or handicap rate (9%) at follow up at 2 years, and in the questionnaire at school age, with a change in severity in six children. Therefore, we considered it justified to perform the logistic regression analysis in the 125 infants with a two year follow up examination.

Total mortality (antenatal and postnatal) was lower with active management. The assumption that antenatal selection could lower morbidity, apparent on short term follow up, was not evident on long term follow up. The disability and handicap rates were comparable between the centres and no difference was detected by univariate and multivariate analysis.

MULTIVARIATE ANALYSIS

Data from 125 children (including 11 non-responders to the questionnaire but with a follow up at 2 years) were analysed using logistic regression. The analysis showed that the risk for disability or handicap was increased in the first study period (RR 30.7 (2.4-396)) and in infants with ICH (RR 38.7 (2.1-717)) or BPD (RR 11.2 (1.5-83)). Centre, birthweight, gestational age, sex, RDS or sepsis did not contribute significantly to the model.

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Total mortality (antenatal and postnatal) was lower with active management. The assumption that antenatal selection could lower morbidity, apparent on short term follow up, was not evident on long term follow up. The disability and handicap rates were comparable between the centres and no difference was detected by univariate and multivariate analysis. This resulted in a significantly higher adverse outcome (total mortality and disability or handicap) in centre A, as a consequence of the intrauterine deaths that occurred with the deliberate non-intervention policy. The estimated handicap rate of 20% expected by centre A was not confirmed.

The total number of impairments by questionnaire was comparable between the centres. It is not clear yet whether the children with impairment suffered similar but less severe brain injuries than the disabled or handicapped
children. If so, the impairments could be in the area of cognitive and behavioural functions.13 15 25 21

No difference in total disability or handicap rate was detected on follow up at 2 years corrected age and at school age by questionnaire. This agrees with the results of Palfrey et al22 and Levy et al.23 Veen et al reported a similar rate of major handicap at 2 and 5 years of age, although their data did not refer to the same children and special education was included in the definition of handicap.

At 2 years of age one child was suspected but not confirmed as abnormal. The disabilities and handicaps on examination in the outpatient department and at follow up by questionnaire were attributed to the same children, but were less severe. The less severe outcome at an older age might have been due to the child’s development or because the investigation of certain abilities is easier at an older age, or a combination of these factors.2 24

Most children with disability or handicap had a combination of neuromotor function disorder, mental retardation, and language and speech abnormality, which has also been reported by other authors.3 25 26

In this group of children birthweight and gestational age were associated with mortality1 but not related to disability or handicap. Similar findings were reported by Veen et al,27 and Scherjon.28 On the other hand, Ehrenhaft et al concluded in a review that birthweight was an important factor, although birthweight alone was used to define the study populations.29

Some authors have found an increased incidence of neurodevelopmental handicaps among children with abnormal ultrasound images of the brain.30–33 Weisglas et al34 showed a significant relation between abnormal scan and BPD. Our results show an independent influence of BPD as well as ICH on disability/handicap.

In our selected study population disability or handicap were related to neonatal complications and not to gestational age and/or (estimated) birthweight. This explains the unconfirmed antepartum prediction of handicaps or disabilities in centre A.

The overall adverse outcome (mortality and disability or handicap) was comparable between both study periods. Notable is the lower disability or handicap rate in the second half of the study period. This concurs with a previous publication.35 This change with time was present in both centres.

As all disabled or handicapped children had already been identified by 2 years corrected age, the disability or handicap rate in the group ascertained in the second period will probably not change with advancing age. Michelsson36 has suggested that five year follow up findings remain valid at nine years of follow up.

Concerns have been expressed by many investigators that offering neonatal intensive care to extremely growth retarded infants might result in a higher prevalence of disabilities or handicaps due to an increase in survival.8 37–39

The lower disability or handicap rate in more recent years in our study is reassuring. During this time, ultrasound imaging of the brain became a definitive part of management in both centres. In some patients treatment was discontinued following sonographic detection of severe cerebral abnormalities. Discontinuation of treatment postnatally seemed to be effective in the prevention of disability or handicap, whereas antenatal prediction and selective non-intervention were not.

Published studies on long term follow up in early preterm infants deal with populations selected by birthweight, gestational age, or a combination of both. The specific selection in the present study—namely, growth retardation and fetal distress due to placental insufficiency, as well as the differences in duration of follow up and outcome definitions—make our results not strictly comparable with those of other studies.

The major handicap rate of survivors in a neonatal intensive care unit (NICU) based study is reported as being 10%.40 41 Dutch NICU based studies detected 12% at 1 year of age42 and 16% at 3.6 years of age.43 Contrary to expected, growth retarded infants as selected in our study seem to have a comparable risk of handicap as other preterm infants.

Whether this also holds for intellectual outcome remains to be seen. Hille et al3 reported a higher handicap rate at nine year follow up compared with that at five years, due to an increase in children who entered special education after the age of 5. Further follow up is needed to discover if the non-disabled or non-handicapped children in our population will eventually show a high rate of minor developmental problems.

Although we have selected our patients extremely carefully and have put considerable effort into a uniform classification of neonatal morbidity we could not control for all possible confounding factors. Some differences between the centres or changes during the period of the study may not have been accounted for.

We consider that randomisation between an intervention vs a selective non-intervention strategy (accepting fetal death) is unethical. A comparison between centres is therefore the next best solution.

Ninety one per cent of surviving infants in this selected population are not disabled or handicapped at school age. Gestational age and birthweight were associated with mortality but not with disability or handicap. Antenatal prediction and selective non-intervention were not effective in handicap and disability prevention. Disability or handicap is mainly related to neonatal complications (ICH and BPD).


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4 Pharush POD, Stevenson CJ, Cooke RWI, Stevenson RC. Clinical and subclinical deficits at 9 years in a geographically defined cohort of low birthweight infants. *Arch Dis Child* 1994;70:264-70.


