Risk factors for early sudden deaths and severe apparent life-threatening events

Anette Poets, Michael S Urschitz, Renate Steinfeldt, Christian F Poets

ABSTRACT

Objective To identify potential risk factors for unexpected sudden infant deaths (SID) and severe apparent life-threatening events (S-ALTE) within 24 h of birth.

Design Case-control study embedded in an epidemiological survey over a 2-year period.

Patients and methods Throughout 2009, every paediatric department in Germany was asked to report cases of unexplained SID or S-ALTE in term infants with a 10-min Apgar score ≥8 to the Surveillance Unit for Rare Pediatric Conditions. Throughout 2010, the inclusion criteria were extended to infants ≥35 week gestational age and those where an explanation for the deterioration had been found. For each unexplained case, hospitals were asked to fill in a questionnaire for 3 (near-)term controls with good postnatal adaptation at the age (in minutes) when the event had occurred in the case under study.

Results Of the 85 cases reported, 34 fulfilled the entry criteria; of these, two were near-term newborns and, in three cases, a cause had been identified for the event. For the 31 cases with unknown cause for the event (13 males; mean (SD) gestational age 38.9 (1.7) week), the authors gathered 93 controls (51 male infants; 38.9 (1.4) week). As significant risk factors for S-ALTE and SID, the authors could identify primipara (OR 6.22; 95% CI 2.11 to 18.32) and potentially asphyxiating position (OR 6.45; 95% CI 1.22 to 34.10).

Conclusions Close observation of newborns seems necessary, particularly in primipara, a potentially asphyxiating position should be avoided.

INTRODUCTION

In Germany, in 2009, we determined the incidence of unexpected sudden infant deaths (SID) and severe apparent life-threatening events (S-ALTE) in term infants (≥37 weeks’ gestational age) within 24 h of birth after a good postnatal adaptation (10-min Apgar score ≥8).1 Seventeen cases met inclusion criteria, corresponding to an incidence of 2.6 per 100 000 live births.

It was noticeable that more than half of the events occurred within 2 h of birth (mostly after vaginal delivery) and more than two-thirds of affected newborns were lying on the breast or abdomen of their mother or near to and facing her. It was also striking that most mothers were primipara, and that, in 7 of 17 cases, the mother had not recognised the deterioration of her baby despite her being present and awake.

In 2010, we continued the survey. As risk factors remain speculative without adequate controls, our aim was to obtain information on birth characteristics and postnatal practices for infants born in hospitals reporting a case, and to compare cases and controls concerning putative risk factors.

METHODS

As part of the Surveillance Unit for Rare Pediatric Conditions in Germany,2 all paediatric departments in Germany were sent monthly reporting cards between 1 January 2009 and 31 December 2009 asking them to notify the study centre of any case of unexplained SID or S-ALTE occurring within 24 h of birth in a full-term infant (≥37 week gestational age) after a good postnatal adaptation (10-min Apgar score ≥8). S-ALTE was defined as an acute state of cyanosis or pallor and unconsciousness, which was felt to require bagging, or intubation with or without cardiac compressions.

We continued the study in 2010 and extended the entry criteria to include near-term infants (≥35 week gestational age) and those where an explanation for the deterioration had been found in retrospect. We made this change because, during the first year of our study, we had been notified of four near-term infants who met inclusion criteria except for their premature birth and, also, of three newborns for whom an explanation for the event had been discovered later.

Hospitals reporting a case received a questionnaire covering the birth, the postnatal situation and the circumstances of the event. In addition, we asked for the infant’s hospital discharge letter, and for the autopsy protocol in the case of...
SID; all documents had to be anonymised before sending them to us. Having collected these data, we excluded cases not meeting inclusion criteria (eg, those who had only been stimulated).

Applying a matched pairs design, we asked every hospital reporting an unexplained case to fill in anonymised questionnaires for three (near-)term infants with a 10-min Apgar score ≥8 born within a few days after receipt of the questionnaires. These had to be filled in at the age (in min) at which the deterioration of the case had been recognised. Questions were on parity, mode of delivery, maternal use of sedatives in the past 24 h before delivery, whether the mother was awake or asleep, infants’ sex, sleeping position and sucking behaviour.

In comparing the information for cases and controls, we tried to verify putative risk factors for such events: primipara, a strenuous delivery, mother not supervising the infant because of being asleep or sedated and a potentially asphyxiating position of the infant, defined as lying prone on the breast or abdomen of its mother or close to her in a side position. As some health professionals reported that the cases had sucked excessively while being breastfed just before the event occurred, we also wanted to investigate the relationship between excessive sucking and such events, despite the lack of a clear definition for ‘excessive sucking’. Because of the predominance of the male sex in SID cases in the first year of life, we also wanted to check the data for each infant’s gender.

The study protocol, including an informed consent waiver, was approved by the Ethics Committee of Tuebingen University Hospital.

**Statistical analysis**

Descriptive statistics (numbers and percentages, means and SDs or medians and ranges) were used to summarise subject characteristics wherever appropriate. According to study design, unadjusted OR and their 95% CI were calculated using unconditional binary logistic regression analysis. The case-control status of the individual (ie, case or control) was used as dependent variable. The following risk factors were investigated for their association with the case-control status: primipara, mode of delivery, maternal use of sedatives in the last 24 h before delivery, sleeping mother, potentially asphyxiating position of the newborn, excessive sucking and gender. Due to the small sample size, multiple regression models to adjust for confounding were not built. A p value of <0.05 was considered statistically significant. All analyses were carried out with statistical software (IBM SPSS Statistics version 19).

**RESULTS**

Children’s hospitals notified us of 85 cases that had occurred in 2009 and 2010; in 2009, 17 of these had met inclusion criteria. In 2010, 17 additional infants met inclusion criteria; two of them were near-term infants, and in three, an explanation was found at a later stage (pulmonary hypertension of the newborn, pneumonia and a Norovirus infection, respectively). In 2010, we were notified of one additional event that had occurred at home; in this case we could not obtain detailed information about the circumstances of the event.

For each of the 31 unexplained cases of SID or S-ALTE (13 males; mean (SD) gestational age 38.9 (1.7) week), we received information on three controls (51 males; 38.9 (1.4) week). A comparison between cases and controls concerning possible risk factors for events is provided in table 1.

The median (25/75 percentile) age at which the case infant was found collapsed (the situation described for controls) was 90 (40/360) min in cases and 90 (59/285) min in controls.

**DISCUSSION**

The number of events reported to us in 2010 was comparable with that identified during the first year of this study. In a recently published nationwide survey in the UK, which was very similar to our study design, the incidence was slightly higher. This could be related to a better reporting of cases or to a less-intense postnatal surveillance, leading to a higher number of events.

In comparing cases with controls, we could confirm ‘asphyxiating position’ and ‘primipara’ as potential risk factors for SID or S-ALTE within 24 h of birth. While a potentially asphyxiating position is a well-known risk factor for SID in general and also supposed as a risk factor for such events in the first 24 h of life, primipara has only been reported as a risk factor for early SID. Beyond the first day of life, the risk for SID seems to increase with parity. In our previous publication, we proposed some explanations for why the above factors may predispose a term newborn to life-threatening events: ‘primipara’ because of an assumed inexperience of the mother in assessing her newborn baby and surveying it, and ‘asphyxiating position’ because of the risk of upper airway obstruction in a newborn lying prone on, or very near to, its mother. In this regard, a contributing factor could be maternal obesity; unfortunately, we did not obtain information on maternal body mass index. Another factor possibly leading to a dangerous situation for the infant could be maternal exhaustion or sedation around delivery. There was no difference in maternal use of sedatives, and labour duration could not be compared between cases and controls as more cases were born to primipara, leading to a
longer average labour duration in this group, and because we did not match cases and controls for mode of delivery.

In discussions of life-threatening events having occurred in the delivery room, health professionals have occasionally mentioned that they had noticed an excessive sucking of the infant shortly before the event had occurred. Therefore, we compared the frequency of excessive sucking in cases and controls but, contrary to our hypothesis, this factor was significantly more prevalent in controls. This might point to control infants being more vigorous, but may also simply be the result of recall bias, which inevitably is a caveat of our study anyway.

We did not analyse smoking as a potential risk factor for early SID/S-ALTE as the prevalence of smoking in cases (4%) was too low to be plausible, which may be related to the fact that this information was only collected several weeks after the event had occurred.

In 2010, we were notified of three cases where a preventable cause for the infant’s deterioration had been identified (although not asked for in 2009, the same number had been reported during that year).

The main limitation of this study is the relatively small number of cases and controls and, hence, the lack of statistical power. Some putative risk factors with elevated but non-significant OR (eg, mother asleep) could become significant in future larger studies. Another limitation was the selection of controls, which was left to the discretion of the hospital staff, leaving potential for selection bias. Although most questionnaires were filled in and nurses who have helped so much by taking part in this study, filling in our questionnaires and reporting their findings to us, and to Derek Stebbens, M.A, for improving the language of this manuscript.

Funding Britich Foundation.

Competing interests None.

Ethics approval Ethics Committee of Tuebingen University Hospital.

Provenance and peer review Not commissioned; externally peer reviewed.

Data sharing statement No additional data.

REFERENCES

Risk factors for early sudden deaths and severe apparent life-threatening events

Anette Poets, Michael S Urschitz, Renate Steinfeldt and Christian F Poets

Arch Dis Child Fetal Neonatal Ed 2012 97: F395-F397 originally published online January 31, 2012
doi: 10.1136/archdischild-2011-300752

Updated information and services can be found at:
http://fn.bmj.com/content/97/6/F395

These include:

References

This article cites 8 articles, 4 of which you can access for free at:
http://fn.bmj.com/content/97/6/F395#BIBL

Email alerting service

Receive free email alerts when new articles cite this article. Sign up in the box at the top right corner of the online article.

Topic Collections

Articles on similar topics can be found in the following collections

- Editor's choice (50)
- Child health (1515)
- Epidemiologic studies (929)
- Infant health (857)
- SIDS (13)

Notes

To request permissions go to:
http://group.bmj.com/group/rights-licensing/permissions

To order reprints go to:
http://journals.bmj.com/cgi/reprintform

To subscribe to BMJ go to:
http://group.bmj.com/subscribe/