Cumulative mortality in children aged 1 to 6 years born in Western Australia from 1980–89

Louisa M Alessandri, Helen M Chambers, Carol Garfield, Sharon Vukovich, Anne W Read

Abstract

Purpose—To investigate cumulative mortality for children aged 1–6 years born in Western Australia from 1980 to 1989. Study design—Births and deaths were ascertained from a linked total population database supplemented by information from postmortem records. Deaths were classified according to the underlying cause, and mortality rates, including factor specific rates, were calculated. Trends were investigated and comparisons were made using relative risks with 95% confidence intervals.

Results—Cumulative mortality was 2.2/1000 infant survivors, with a significant decrease during the years studied. Mortality was almost four times higher for Indigenous children, with no decrease. Accidents comprised 45.6% of all deaths, birth defects 17.3%, cancer and leukaemias 12.5%, and infections 11.0%. Low birth weight, preterm birth, and young maternal age significantly increased the risk of death in both Indigenous and non-Indigenous children; single marital status was also a significant risk factor for non-Indigenous children.

Conclusion—High quality data and appropriate classification systems are essential to enable effective monitoring of childhood deaths and the planning of preventive programmes. Further decreases in mortality rates might be dependent on ensuring that resources are directed towards improving social and economic conditions for Indigenous and other disadvantaged families.

Keywords: mortality; Western Australia; cause of death; epidemiology

Recent literature on childhood mortality in industrialised countries focuses on accidents and/or injuries as the major causes of childhood deaths.1–4 There is little that deals with the overall patterns of childhood mortality and, in particular, we have been unable to find any investigations on the specific causes of childhood mortality for birth cohorts for a total population.

Previous work in Western Australia has highlighted the importance of considering the Indigenous (see editors’ note) status of the child when analysing mortality data in infancy because Indigenous infants have been shown to have much higher postneonatal and infant mortality than non-Indigenous infants.5–7 The leading causes of death in infancy also differ in the two populations, with sudden infant death syndrome (SIDS) being the major cause of death for Indigenous infants and birth defects for non-Indigenous infants.8

The aim of our retrospective, longitudinal study was to investigate childhood mortality for all children aged 1–6 years inclusive born from 1980 to 1989 and dying from 1981 to 1996 inclusive in Western Australia. The available data for these birth cohorts allowed us to compare trends and causes of childhood mortality for the Indigenous and non-Indigenous populations, and to investigate the data according to selected maternal and infant variables, to obtain information that would be useful in assisting with the planning of preventive strategies for childhood deaths.

Methods

All children born in Western Australia from 1980 to 1989 inclusive were ascertained from the Western Australia maternal and child health research database (MCHRDB).10 This database includes death information linked to birth data for all births in Western Australia from 1980 onwards. Infant survivors from the 1980 to 1989 birth cohorts were followed to their 7th birthdays to ascertain which children had died between the ages of 1 and 6 years inclusive. Cause of death data for the MCHRDB are obtained from the Registrar General of Western Australia via the Australian Bureau of Statistics (ABS). The Registrar General’s Office collects descriptive data pertaining to the cause of death from death certificates; these data are then forwarded to the ABS, where they are coded according to the International classification of diseases 9th revision (ICD–9).11

Table 1  Childhood deaths for children aged 1–6 years born in Western Australia 1980–89

<table>
<thead>
<tr>
<th>Year of birth</th>
<th>Rate/1000 infant survivors (n)</th>
<th>Indigenous</th>
<th>Non-Indigenous</th>
<th>Total</th>
</tr>
</thead>
<tbody>
<tr>
<td>1980</td>
<td>7.0 (7)</td>
<td>2.7 (52)</td>
<td>3.0 (61)</td>
<td></td>
</tr>
<tr>
<td>1981</td>
<td>11.2 (12)</td>
<td>2.5 (51)</td>
<td>2.9 (64)</td>
<td></td>
</tr>
<tr>
<td>1982</td>
<td>3.7 (4)</td>
<td>2.0 (41)</td>
<td>2.1 (46)</td>
<td></td>
</tr>
<tr>
<td>1983</td>
<td>10.8 (12)</td>
<td>1.8 (39)</td>
<td>2.3 (51)</td>
<td></td>
</tr>
<tr>
<td>1984</td>
<td>9.6 (11)</td>
<td>2.1 (44)</td>
<td>2.4 (55)</td>
<td></td>
</tr>
<tr>
<td>1985</td>
<td>6.7 (8)</td>
<td>1.6 (35)</td>
<td>1.9 (44)</td>
<td></td>
</tr>
<tr>
<td>1986</td>
<td>6.7 (8)</td>
<td>1.3 (28)</td>
<td>1.5 (36)</td>
<td></td>
</tr>
<tr>
<td>1987</td>
<td>6.1 (8)</td>
<td>1.8 (41)</td>
<td>2.1 (49)</td>
<td></td>
</tr>
<tr>
<td>1988</td>
<td>5.8 (8)</td>
<td>1.5 (36)</td>
<td>1.8 (45)</td>
<td></td>
</tr>
<tr>
<td>1989</td>
<td>7.1 (10)</td>
<td>1.7 (41)</td>
<td>2.0 (51)</td>
<td></td>
</tr>
<tr>
<td>1980–1989</td>
<td>7.2 (88)</td>
<td>2.1 (408)</td>
<td>2.2 (502)*</td>
<td></td>
</tr>
</tbody>
</table>

p value 0.370 0.001 0.001

*There were six childhood deaths where the race was unknown.
Table 2  Cause of childhood deaths by Indigenous status for children aged 1–6 years born in Western Australia from 1980–89

<table>
<thead>
<tr>
<th>Cause of death*</th>
<th>Indigenous (n=88)</th>
<th>Non-Indigenous (n=407)</th>
<th>Relative risk (95% CI)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Birth defects</td>
<td>9.3 (11)</td>
<td>3.4 (74)</td>
<td>2.7 (1.5 to 5.1)</td>
</tr>
<tr>
<td>Infections</td>
<td>16.0 (19)</td>
<td>1.7 (36)</td>
<td>9.7 (5.6 to 16.9)</td>
</tr>
<tr>
<td>Sudden infant death syndrome</td>
<td>1.7 (2)</td>
<td>0.7 (15)</td>
<td>2.5 (0.6 to 0.7)</td>
</tr>
<tr>
<td>Accidents</td>
<td>37.9 (45)</td>
<td>8.3 (181)</td>
<td>4.6 (3.3 to 6.3)</td>
</tr>
<tr>
<td>Cancer and leukaemias</td>
<td>2.5 (3)</td>
<td>2.8 (60)</td>
<td>0.9 (0.3 to 2.9)</td>
</tr>
<tr>
<td>Neurological conditions</td>
<td>5.1 (6)</td>
<td>0.7 (15)</td>
<td>7.3 (2.9 to 18.9)</td>
</tr>
<tr>
<td>Other</td>
<td>1.7 (2)</td>
<td>0.8 (18)</td>
<td>2.0 (0.5 to 8.8)</td>
</tr>
<tr>
<td>Unascertainable</td>
<td>-</td>
<td>0.4 (8)</td>
<td>-</td>
</tr>
</tbody>
</table>

There was one non-Indigenous childhood death caused by perinatal conditions.

Excludes six deaths where race was unknown.

Table 3  Deaths by age at death and major cause for non-Indigenous children aged 1–6 years born in Western Australia 1980–89

<table>
<thead>
<tr>
<th>Age</th>
<th>Birth</th>
<th>Accidents</th>
<th>Cancer and leukaemias</th>
<th>All causes</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Incidents</td>
<td>Infections</td>
<td></td>
<td></td>
</tr>
<tr>
<td>1</td>
<td>2.8 (61)</td>
<td>1.6 (34)</td>
<td>0.4 (9)</td>
<td>0.6 (13)</td>
</tr>
<tr>
<td>2</td>
<td>1.8 (40)</td>
<td>0.9 (19)</td>
<td>0.5 (11)</td>
<td>0.3 (6)</td>
</tr>
<tr>
<td>3</td>
<td>1.6 (34)</td>
<td>0.5 (10)</td>
<td>0.3 (7)</td>
<td>0.7 (15)</td>
</tr>
<tr>
<td>4</td>
<td>0.9 (19)</td>
<td>0.2 (4)</td>
<td>0.1 (2)</td>
<td>0.6 (12)</td>
</tr>
<tr>
<td>5</td>
<td>0.7 (16)</td>
<td>0.1 (2)</td>
<td>0.1 (3)</td>
<td>0.4 (9)</td>
</tr>
<tr>
<td>6</td>
<td>0.6 (12)</td>
<td>0.2 (5)</td>
<td>0.2 (4)</td>
<td>0.2 (5)</td>
</tr>
</tbody>
</table>

To validate the cause of death data from the MCHRDB, postmortem records were obtained for as many childhood deaths as possible. Postmortem data were collected from three major centres, the forensic pathology section of the Western Australia State Health Laboratory Services in Perth, Princess Margaret Hospital for Children (PMH) (the only tertiary paediatric hospital in Western Australia), and the Perth Coroner’s Office. Since 1990, all postmortem examinations are conducted centrally at the forensic pathology section. Although some of the accidental deaths that occur in the country areas have postmortem examinations performed locally, postmortem records from 1986 onwards have been held at the Perth Coroner’s Office. For country deaths from 1980 to 1985, the records are held at the Coroner’s Court in the appropriate country region. There are eight such country regions and all were contacted by mail to obtain postmortem information.

Since 1990, all postmortem examinations carried out under the State Coroner on sudden and unexpected deaths up to the age of 2 years, other than those related to accidents, have been attended by and the material reviewed by specialised paediatric pathologists based at PMH. Postmortem examinations on children who were admitted and died at PMH were conducted by the same pathologists at the hospital. The birth defects registry of Western Australia,9 which linked to the MCHRDB, was also used as a data source to ascertain postmortem information for those deaths where records could not be found at the above centres. After all information was gathered, these data were compared with the cause of death information recorded on the MCHRDB to determine if there were any discrepancies.

A simple and broad classification system for cause of childhood death was designed with the aim of identifying major changes in causation over time, and such that it was useful in identifying preventable causes of death. Under this system, all deaths were classified into major groupings according to the condition that was thought to be the underlying cause. The major categories “other” included uncommon but specific causes of death (such as cardiomyopathy and mechanical bowel obstruction). Accidents were classified further into road traffic accidents, drowning, poisoning, deaths attributable to fire, other accidents, and homicide. The “other accident” group included farm accidents, accidental mechanical suffocation, choking, and blows from falling objects. All the childhood deaths were classified independently by three of the authors (LA, AR, and HC). Any disagreement was resolved by discussion. If the cause of death data from the MCHRDB differed from that derived from the postmortem data, the latter took precedence.

Information on maternal race, age, parity, and marital status (as recorded at the birth of the child, de facto partnerships being included in the married category), and on sex, birth weight, and gestational age of the child at birth were obtained from the MCHRDB. Children with Aboriginal or Torres Strait Islander mothers were classified as Indigenous children. Other data such as age at death were collected from death certificates. SAS computer software14 was used for data entry and manipulation and for obtaining frequencies. Overall rates of childhood death for birth cohorts 1980 to 1989 were calculated for each 1000 infant survivors by Indigenous status and for each 10 000 infant survivors by cause of death. Factor specific childhood mortality rates were calculated for all children for total mortality and, for non-Indigenous children for major cause of death categories. This latter analysis was not performed for Indigenous children because of small numbers in some categories. Unknown values were few and were excluded from all calculations. Chi squared tests for trend were calculated to investigate trends in mortality over time, and relative risks with Taylor series 95% confidence intervals (CI) were calculated to compare rates using the EpiInfo software package.14 The denominators for all rates were obtained from the MCHRDB.

Results

There were 502 deaths of children aged 1–6 years ascertained for our study. Death information from the ABS could not be linked to birth information on the MCHRDB for four of these deaths; two of these four children had been adopted. Postmortem records were found for 67.3% (n=338) of the total deaths. Most of these records (66.9%; n=226) were located...
Table 4  Deaths by Indigenous status and selected maternal and infant variables for children aged 1–6 years born in Western Australia 1980–89

<table>
<thead>
<tr>
<th>Marital status†</th>
<th>Sex</th>
<th>Birth weight</th>
<th>Gestational age</th>
<th>Maternal age*</th>
<th>Parity†</th>
<th>Birth defects</th>
<th>Infections</th>
<th>Accidents</th>
<th>Cancer and leukaemias</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Female</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td>Rate/1000 infant survivors (n)</td>
<td>Relative risk (95% CI)</td>
<td>Rate/1000 infant survivors (n)</td>
<td>Relative risk (95% CI)</td>
</tr>
<tr>
<td></td>
<td>Single/other</td>
<td>≥ 2500 g</td>
<td>≥ 37 weeks</td>
<td>≥ 20 years</td>
<td>≥ 1</td>
<td>3.2 (6)</td>
<td>0.9 (0.4 to 2.1)</td>
<td>1.3 (25)</td>
<td>7.7 (152)</td>
</tr>
<tr>
<td></td>
<td>Married/de facto</td>
<td>&lt; 2500 g</td>
<td>&gt; 37 weeks</td>
<td>&lt; 20 years</td>
<td>&lt; 1</td>
<td>3.4 (44)</td>
<td>1.0 (0.6 to 1.5)</td>
<td>1.5 (25)</td>
<td>6.5 (55)</td>
</tr>
<tr>
<td></td>
<td>Boy</td>
<td>&lt; 2500 g</td>
<td>&lt; 37 weeks</td>
<td>&lt; 20 years</td>
<td>&lt; 1</td>
<td>3.4 (48)</td>
<td>0.9 (0.4 to 1.5)</td>
<td>1.8 (375)</td>
<td>8.3 (137)</td>
</tr>
<tr>
<td></td>
<td>Girl</td>
<td>≥ 2500 g</td>
<td>≤ 37 weeks</td>
<td>≥ 20 years</td>
<td>≥ 1</td>
<td>3.4 (62)</td>
<td>1.0 (0.7 to 1.5)</td>
<td>1.9 (21)</td>
<td>8.3 (108)</td>
</tr>
<tr>
<td></td>
<td>Single/other</td>
<td>≥ 37 weeks</td>
<td>≥ 20 years</td>
<td>≥ 1</td>
<td>3.4 (68)</td>
<td>0.9 (0.4 to 2.1)</td>
<td>1.3 (25)</td>
<td>7.7 (152)</td>
<td>2.7 (54)</td>
</tr>
<tr>
<td></td>
<td>Married/de facto</td>
<td>&lt; 37 weeks</td>
<td>&lt; 20 years</td>
<td>&lt; 1</td>
<td>3.4 (21)</td>
<td>1.0 (0.6 to 1.5)</td>
<td>1.5 (20)</td>
<td>6.5 (55)</td>
<td>3.4 (29)</td>
</tr>
<tr>
<td></td>
<td>Boy</td>
<td>≥ 2500 g</td>
<td>&gt; 37 weeks</td>
<td>&gt; 20 years</td>
<td>&gt; 1</td>
<td>3.4 (48)</td>
<td>0.9 (0.4 to 1.5)</td>
<td>1.8 (375)</td>
<td>8.3 (137)</td>
</tr>
<tr>
<td></td>
<td>Girl</td>
<td>&lt; 2500 g</td>
<td>≤ 37 weeks</td>
<td>&lt; 20 years</td>
<td>&lt; 1</td>
<td>3.4 (62)</td>
<td>1.0 (0.7 to 1.5)</td>
<td>1.9 (21)</td>
<td>8.3 (108)</td>
</tr>
</tbody>
</table>

*At birth of child.
†Separated, divorced, or widowed.
‡At birth of child.
§Married/de facto.

at the forensic pathology section of the State Health Laboratories, with 17.2% (n = 58) being ascertained from PMH, 13.6% (n = 46) from the Perth Coroner’s Office, 1.8% (n = 6) from Country Coroners’ Courts, and 0.6% (n = 2) from the Birth Defects Registry. Postmortem records could not be found for five deaths that were identified as Coroner’s cases on the death certificate. All of the eight Country Coroners’ Courts replied to our request for data but only three could provide information, the other regions being unable to locate the records required. The proportion of Indigenous childhood deaths (71.6%) for which postmortem information was ascertained was slightly higher than that for non-Indigenous children (66.4%). There was discrepancy between postmortem information and the MCHRDB in only nine (1.8%) of the total childhood deaths. Of the 17 deaths (3.4%) where there was initial disagreement between the authors as to the classified cause of death, all were subsequently reclassified with complete agreement.

The cumulative incidence of death for children aged from 1 to 6 years born from 1980 to 1989 in Western Australia was 2.2/1000 infant survivors. The rate decreased significantly over time for the total population and for non-Indigenous children but not for Indigenous children (table 1). The relative risk for Indigenous compared with non-Indigenous children was 4.0 (95% CI, 3.1 to 5.0).

In both Indigenous and non-Indigenous populations, the cumulative mortality for the ages studied was highest for accidents. The leading causes of death for Indigenous children were accidents, followed by infections, and birth defects, whereas for non-Indigenous children they were accidents, followed by birth defects, and cancer and leukaemias. The Indigenous population had higher rates of childhood deaths for all causes except cancer and leukaemias, where the rates were very similar. The greatest disparity between the Indigenous and non-Indigenous populations was for childhood deaths caused by infections, the relative risk being 9.7 (table 2).

Within the accident category, differences were seen between Indigenous and non-Indigenous children, with the former being more likely to die from road traffic accidents (rate, 12.6/10 000 infant survivors) and the latter from drowning (rate, 3.2/10 000 infant survivors). The mortality rate for non-Indigenous children for road traffic accidents was 2.3/10 000 and the rate for Indigenous children for drowning was 8.4/10 000. Death as a result of fire was grossly excessive in the Indigenous population with a rate of 7.6/10 000 infant survivors with a rate of 0.1/10 000 for non-Indigenous children (data not shown).

As age increased, the incidence of death decreased but accidents were the major cause of death in each age group. Drownings accounted for the major proportion of accidental deaths in the younger age groups, whereas road traffic accidents were responsible for most of the accidental deaths in the older age groups (data not shown). Of the other causes, birth

Table 5  Deaths by selected maternal and infant variables and major cause for non-Indigenous children aged 1–6 years born in Western Australia 1980–89

<table>
<thead>
<tr>
<th>Birth defects</th>
<th>Infections</th>
<th>Accidents</th>
<th>Cancer and leukaemias</th>
</tr>
</thead>
<tbody>
<tr>
<td>Rate* (n)</td>
<td>RR (95% CI)</td>
<td>Rate* (n)</td>
<td>RR (95% CI)</td>
</tr>
<tr>
<td>Sex</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Female</td>
<td>3.3 (35)</td>
<td>1.4 (15)</td>
<td>6.9 (73)</td>
</tr>
<tr>
<td>Male</td>
<td>3.5 (39)</td>
<td>1.9 (21)</td>
<td>9.7 (108)</td>
</tr>
<tr>
<td>Birth weight</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>&lt; 2500 g</td>
<td>3.0 (62)</td>
<td>1.5 (31)</td>
<td>8.2 (169)</td>
</tr>
<tr>
<td>≥ 2500 g</td>
<td>10.9 (12)</td>
<td>4.5 (5)</td>
<td>10.9 (12)</td>
</tr>
<tr>
<td>Gestational age</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>≥ 37 weeks</td>
<td>3.1 (63)</td>
<td>1.5 (31)</td>
<td>8.3 (168)</td>
</tr>
<tr>
<td>&lt; 37 weeks</td>
<td>7.5 (11)</td>
<td>2.2 (134)</td>
<td>8.9 (13)</td>
</tr>
<tr>
<td>Maternal age‡</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>≥ 20 years</td>
<td>3.4 (69)</td>
<td>1.6 (33)</td>
<td>7.9 (162)</td>
</tr>
<tr>
<td>&lt; 20 years</td>
<td>4.3 (5)</td>
<td>2.6 (3)</td>
<td>16.3 (19)</td>
</tr>
<tr>
<td>Parity†</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>0</td>
<td>3.3 (35)</td>
<td>1.9 (16)</td>
<td>6.5 (55)</td>
</tr>
<tr>
<td>≥ 1</td>
<td>3.4 (44)</td>
<td>1.5 (20)</td>
<td>9.6 (126)</td>
</tr>
<tr>
<td>Marital status‡</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Married/de facto</td>
<td>3.4 (68)</td>
<td>1.3 (25)</td>
<td>7.7 (152)</td>
</tr>
<tr>
<td>Single/other</td>
<td>3.2 (6)</td>
<td>0.9 (0.4 to 2.1)</td>
<td>5.8 (11)</td>
</tr>
</tbody>
</table>

*Rate/10 000 infant survivors.
†At birth of child.
‡Married/de facto.
defects and infections were more important in the younger age groups, whereas cancer and leukaemias were more important causes of death for older children (table 3).

In the Indigenous population, low birth weight, preterm birth, and young maternal age were significant risk factors for all causes of childhood mortality. In addition to these factors, for non-Indigenous children, single marital status was significant (table 4). When analysing the variables of interest by the major cause of death groups for non-Indigenous children only (table 5), it was found that the variables analysed were not necessarily risk factors for each cause of death. Boys were at significantly increased risk of dying from accidents but not from the other major causes of childhood death. Low birth weight was an important risk factor for childhood deaths attributable to birth defects and infections. Children of teenage mothers were at significantly increased risk of dying from accidents, whereas those of single mothers had a significant increase in risk from both infections and accidents. None of the factors investigated were associated with an increased risk of childhood death from cancer and leukaemias.

Discussion

In Western Australia, the 1–6 years mortality rate for non-Indigenous children born during the 1980s decreased significantly over time. No such decreases were seen for Indigenous children, who remained at much higher risk. Children of young mothers and unsupported mothers were also at increased risk of childhood death, as were children who were preterm or low birth weight. The major cause of death for all children was accidents.

Longitudinal research of this type is rare in the literature and we have been unable to find any similar studies using the cumulative incidence of mortality with which to compare our results. Data to document and analyse total mortality for individual children during the entire course of childhood are rarely available. Thus, most researchers use the population prevalence when investigating childhood mortality. Cumulative incidence allows more precise estimation of rates and more accurate investigation of time trends and risk factors.

Substantial declines in mortality for children (aged 1–4 and 1–14 years) have been reported recently from both industrialised and other countries, and the excess in mortality for Indigenous children has been described in other Australian studies. Our study highlights the ongoing disadvantage for these children. Indigenous populations such as Indians in the USA and Canada also have higher rates of childhood mortality than the corresponding white population or the nation as a whole.

Indigenous families in Australia are known to live in poorer socioeconomic and environmental conditions than non-Indigenous families, and the relation between low socioeconomic class and high childhood mortality has been well documented.

Key messages

- In Western Australia, all cause mortality in early childhood has significantly decreased for later born cohorts
- There was no decrease for Indigenous children
- Much of the disparity between the two populations was attributable to potentially preventable causes, such as accidents and infections
- There is scope to reduce this excess mortality but any intervention must be implemented in close collaboration with the Indigenous community
- Accidents and infections are also excessive for the infants of young and unsupported mothers in the overall population

The overall ranking of causes of death is similar to that seen in Sweden and the USA, with the potentially preventable cause of accidents being of major importance. The most striking difference between Indigenous and non-Indigenous was in the rate of deaths attributable to infection, another potentially preventable cause, and one for which Indigenous children are about 10 times more likely to be admitted to hospital.

The association of factors such as preterm birth, low birth weight, young mothers, and single mothers with high rates of perinatal and infant mortality is well established, and it appears from this study and others that the disadvantage conferred by their impact persists throughout childhood. In the UK, it has been suggested that the increased injury rates for the children of single mothers can be explained by the poverty, poor housing conditions, and social isolation of these mothers. The importance of low social class in terms of childhood mortality has been highlighted further by another recent study in England and Wales where the authors found that the decline in death rates from injury for children in social classes I and II had been greater than for children in social classes IV and V. In the USA, Singh and Yu noted that the association of socioeconomic status with accidents was stronger than for overall mortality and Mare came to similar conclusions. For accidents only, we found that boys had significantly higher rates of death than girls. Studies in the UK and Norway have also shown an excess of childhood accidents in boys, and Singh and Yu have reported a higher overall childhood mortality rate for boys than for girls in the USA.

To develop effective programmes aimed at reducing childhood mortality it is essential to have high quality data systems that allow good epidemiological studies to be conducted. Monitoring of the incidence, prevalence, and trends of childhood deaths, and a clear understanding of their epidemiology, is vital for the development, implementation, and evaluation of appropriate policies and programmes aimed...
Mortality in 1–6 year olds in Western Australia

Statistics are acknowledged for providing access to birth and birth defects registry of Western Australia for their cooperation. Services, Princess Margaret Hospital for Children, and the for her support and access to data. We would also like to thank Mrs V Gee (Western Australian midwives’ the National Health and Medical Research Council of public health research and development grants committee of recognised by a number of authors.1 39–41

The most important groups in terms of the prevention of childhood mortality were Indigenous children, young mothers, single mothers, and children who were preterm or of low birth weight. In close collaboration with the Indigenous community, attempts must be made to ascertain and implement the most appropriate strategies to prevent deaths in this population. Measures must be taken to curb road traffic accidents and living conditions must be improved to decrease the risk of infections. Extra information and advice about safety guidelines and current preventive strategies, such as the importance of adequate immunisation, adult supervision of children while swimming, child restraints in motor vehicles, and fire detectors should be given to all at risk groups. In the UK, the importance of the role of the health visitor in preventing childhood accidents has long been recognised,39 and a similar scheme might be an appropriate way to deliver such information and services to these groups. For the Indigenous population, such programmes would have to be closely linked with community development processes and need to be conducted and staffed from within the community as far as possible.40 There appear to be social class differences in compliance with safety practices and it has been suggested that this reflects class differences in cultural and material resources. Thus, the groups at higher risk require not only the knowledge of safety laws and guidelines but also the resources to implement them.

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