Rates of radiologically confirmed pneumonia as defined by the World Health Organization in Northern Territory Indigenous children

Kerry-Ann F O’Grady, Debbie M Taylor-Thomson, Anne B Chang, Paul J Torzillo, Peter S Morris, Grant A Mackenzie, Gavin R Wheaton, Paul A Bauert, Margaret P De Campo, John F De Campo and Alan R Ruben

ABSTRACT

Objective: To determine the burden of hospitalised, radiologically confirmed pneumonia (World Health Organization protocol) in Northern Territory Indigenous children.

Design, setting and participants: Historical, observational study of all hospital admissions for any diagnosis of NT resident Indigenous children, aged between ≥ 29 days and < 5 years, 1 April 1997 to 31 March 2005.

Intervention: All chest radiographs taken during these admissions, regardless of diagnosis, were assessed for pneumonia in accordance with the WHO protocol.

Main outcome measure: The primary outcome was endpoint consolidation (dense fluffy consolidation [alveolar infiltrate] of a portion of a lobe or the entire lung) present on a chest radiograph within 3 days of hospitalisation.

Results: We analysed data on 24 115 hospitalised episodes of care for 9492 children and 13 683 chest radiographs. The average annual cumulative incidence of endpoint consolidation was 26.6 per 1000 population per year (95% CI, 25.3–27.9); 57.5 per 1000 per year in infants aged 1–11 months, 38.3 per 1000 per year in those aged 12–23 months, and 13.3 per 1000 per year in those aged 24–59 months. In all age groups, rates of endpoint consolidation in children in the arid southern region of NT were about twice that of children in the tropical northern region.

Conclusion: The rates of severe pneumonia in hospitalised NT Indigenous children are among the highest reported in the world. Reducing this unacceptable burden of disease should be a national health priority.

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METHODS

Design

We conducted a historical observational study between 1 April 1997 and 31 March 2005 of all NT hospital admissions, irrespective of diagnosis, of Indigenous children aged ≥ 29 days and < 5 years at the time of admission. Exclusions were admitted if the child did not reside in the NT, or the location of residence was unknown or not specified. All chest x-rays taken during all admissions were reviewed, but those taken more than 3 days from the date of admission were excluded as being more likely to represent nosocomial infections. The primary outcome was endpoint consolidation seen on a chest x-ray taken ≤ 3 days after admission.

Setting

In 2001, the NT had an estimated resident population of about 197 800 people4 dispersed across 1 346 200 km²; 29% of the population identify as Aboriginals or Torres Strait Islanders. Two climate zones exist in the NT: tropical in the north (the “Top End”) and arid in the south (the “Centre”). The average annual population of NT resident Indigenous children aged ≥ 29 days and < 5 years over the study period was 7214 (range, 7047–7382).

All Indigenous children requiring hospitalisation are admitted to one of the five public hospitals in the NT, including those who need subsequent transfer to larger institutions in other states. The nearest public hospitals in other states are hundreds of kilometres from the NT borders. While not formally documented, expert opinion indicates that out-of-hospital deaths in the NT are rare.

X-ray facilities are not available in remote communities. However, given the high incidence of multiple morbidities in Indigenous children, clinical algorithms specify that all Indigenous children admitted to hospital with respiratory illnesses, gastroenteritis, malnutrition, failure to thrive and/or anaemia have a chest x-ray on admission.

Hospitalisation data

Up to 30 June 1998, the International classification of diseases, 9th revision, clinical modification (ICD-9-CM), was used for morbidity coding of records; for the remaining years, the 10th revision, Australian modification (ICD-10-AM) was used. Box 1 shows the hierarchy of diagnosis codes selected. Unique health record numbers permit linking of data and tracking of individuals in the NT public hospital system.

Radiological data

All chest x-rays corresponding to inpatient episodes of care were obtained, de-identified, and a randomly generated, unique study number was assigned to each film. Inpatient episodes of care for which no x-ray was taken, or the x-ray taken could not be found, were recorded as “not done”, or “missing”, respectively.

A panel of seven paediatric and respiratory physicians assigned the presence of endpoint consolidation (yes or no) for each
1 Hierarchical selection of ICD-9-CM and ICD-10-AM codes for analysis

<table>
<thead>
<tr>
<th>ICD-9-CM</th>
<th>ICD-10-AM</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>460–519</td>
<td>J00–J99</td>
<td>Major diagnostic category of diseases of the respiratory system</td>
</tr>
<tr>
<td>466–487</td>
<td>J10–J22</td>
<td>Acute lower respiratory infections</td>
</tr>
<tr>
<td>481–486</td>
<td>J13–J16</td>
<td>Non-viral pneumonias (includes pneumonia unspecified)</td>
</tr>
<tr>
<td>481–482</td>
<td>J13–J15</td>
<td>Specified bacterial pneumonias</td>
</tr>
<tr>
<td>481–482</td>
<td>J13–J15</td>
<td>Pneumococcal pneumonia</td>
</tr>
<tr>
<td>460–466</td>
<td>J21</td>
<td>Bronchitis</td>
</tr>
</tbody>
</table>


2 Demographic characteristics of children hospitalised and the episodes of care

<table>
<thead>
<tr>
<th>Characteristic</th>
<th>No. (%) children</th>
<th>No. (%) episodes</th>
</tr>
</thead>
<tbody>
<tr>
<td>Sex</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Male</td>
<td>5115 (53.9%)</td>
<td>13 281 (55.1%)</td>
</tr>
<tr>
<td>Female</td>
<td>4376 (46.1%)</td>
<td>10 828 (44.9%)</td>
</tr>
<tr>
<td>Not specified</td>
<td>1 (0.0)</td>
<td>6 (0.02%)</td>
</tr>
<tr>
<td>Total</td>
<td>9492 (100%)</td>
<td>24 115 (100%)</td>
</tr>
<tr>
<td>Age group (months)</td>
<td></td>
<td></td>
</tr>
<tr>
<td>at the time of admission</td>
<td></td>
<td></td>
</tr>
<tr>
<td>1–11</td>
<td>5426 (57.2%)</td>
<td>9997 (41.5%)</td>
</tr>
<tr>
<td>12–23</td>
<td>4432 (46.7%)</td>
<td>7 225 (30.0%)</td>
</tr>
<tr>
<td>24–59</td>
<td>4291 (45.2%)</td>
<td>6 893 (28.6%)</td>
</tr>
<tr>
<td>Northern Territory region</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Top End</td>
<td>5887 (62.0%)</td>
<td>13 402 (55.6%)</td>
</tr>
<tr>
<td>Centre</td>
<td>3543 (37.3%)</td>
<td>10 612 (44.0%)</td>
</tr>
<tr>
<td>Key comorbidities diagnosed at least once†</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Gastrointestinal disorders</td>
<td>4891 (51.5%)</td>
<td>8 439 (34.9%)</td>
</tr>
<tr>
<td>Metabolic disturbance‡</td>
<td>1468 (15.5%)</td>
<td>2 319 (9.6%)</td>
</tr>
<tr>
<td>Anaemia</td>
<td>3288 (34.6%)</td>
<td>4 977 (20.6%)</td>
</tr>
</tbody>
</table>

* Individual children may be represented more than once (eg, admitted when aged 1–11 months and readmitted when aged 12–23 months). † Number of episodes of care in which a comorbidity was diagnosed. ‡ Includes malnutrition, failure to thrive.

Statistical analyses

The primary analysis was the cumulative incidence of endpoint consolidation, stratified by age group of the child and NT health region, with calculation of incidence rate ratios. All instances of endpoint consolidation, irrespective of the discharge diagnosis, were included in the analysis to maximise the capture of cases and to ensure that the case definition for the primary analysis was systematically applied over time. Incidence rate ratios were computed to compare rates by NT region and sex. Denominators — the estimated resident population for each year of the study — were obtained from the Australian Bureau of Statistics and the NT Department of Health and Families. Data were analysed using Stata SE, version 9.1 (StataCorp, College Station, Tex, USA).

Ethics approval

The study was approved by the joint institutional Human Research Ethics Committee of the NT Department of Health and Community Services and the Menzies School of Health Research (HREC ID: 02/63), and by the Human Research Ethics Committee of Central Australia. Both committees have Indigenous Health Research Ethics Subcommittees.

RESULTS

Final dataset

Between 1 April 1997 and 31 March 2005, there were 24 820 admitted episodes of care for 9787 Indigenous children (regardless of NT residence) aged between ≥29 days and <5 years (median age, 2; range, 1–39 episodes per child). In 10 911 episodes (43.9%) no chest x-ray was taken, and in 64 episodes (0.3%) the film was missing; 18 224 x-rays were located and assessed for endpoint consolidation (median, 1; range, 0–142 per episode). The proportion of admissions in which x-rays were taken remained stable over the study period (an average of 56% of all hospitalisations).

Of the 24 820 admitted episodes of care, 705 episodes for 295 children and 4541 chest x-rays were excluded from the analysis as they did not meet eligibility criteria (child not an NT resident, x-ray taken ≥3 days after admission). The final dataset comprised 24 115 admitted episodes of care for 9492 children and 13 683 x-rays. Box 2 gives the demographic characteristics of the children and the episodes of care.
Radiologically confirmed pneumonia

Overall, in 13,205 episodes of care (54.7%), irrespective of diagnosis, chest x-rays were taken within 3 days; end-point consolidation was found in 11.6% (1535) of these; film quality was inadequate in 14.0% (1854).

The 1535 episodes of end-point consolidation occurred in 1211 children with a range of 1–16 episodes per child (one episode, 937 children; two or more episodes, 274 children). Median age at the time of admission was 15 months (range, 1–59 months); 40.8% of episodes were for children aged <12 months and 23% were for children aged <6 months. Rates were 1.22 times higher for boys (95% CI, 1.19–1.26; P<0.001), and 2.1 times higher in children aged less than 12 months in the Central Australian region of the NT (Box 3). The average annual cumulative incidence of end-point consolidation was 26.6 per 1000 population per year (95% CI, 25.3–27.9); 57.5 per 1000 per year in infants aged 1–11 months, 38.3 per 1000 per year in those aged 12–23 months, and 13.3 per 1000 per year in those aged 24–59 months. Annual rates for each year of the study by age group are shown in Box 4.

Chest x-rays were taken within 3 days of admission in 6852 of the 8518 episodes (80.4%) with any diagnosis in the category, acute lower respiratory infection (ALRI). Film quality was inadequate for determining end-point consolidation in 1305 (19.0%) episodes with ALRI. End-point consolidation was diagnosed in 20.4% of episodes with ICD-defined ALRI (1401/6852), 33.2% with non-viral pneumonia (1178/3551); 40.4% with pneumococcal pneumonia (21/52), 13.8% with influenza (32/232) (of which 3% had a concomitant diagnosis of non-viral pneumonia), and 9.2% with bronchiolitis (226/2455) (of which 43% had a concomitant diagnosis of non-viral pneumonia).

Sixty-four episodes with end-point consolidation had no respiratory diagnosis recorded (4.2% of end-point consolidation episodes). There were 258 diagnoses attached to these episodes; 59 (22.9%) were conditions listed under the major diagnostic category of infectious and parasitic diseases (predominantly gastrointestinal infections); 55 (21.3%) were haematological and metabolic disorders; 36 (14.0%) were related to external causes and injury (eg, trauma or near drowning) and 10 (3.9%) were ear diseases. The remainder were scattered across other diagnostic categories. As coding errors cannot be excluded, these episodes were included in the analysis.

There were differing seasonal patterns of end-point consolidation and other respiratory diagnoses between the NT regions. Episodes of end-point consolidation were more frequent in the winter and spring months in the Central Australian region of NT; in the Top End region, episodes were more evenly distributed across the year.

**DISCUSSION**

We found that the annual incidence of end-point consolidation in NT children under 5 years of age approximates 3%, and in infants aged under 12 months it is as high as 7% Differences in study design, case ascertainment, and populations complicate direct comparisons, however, rates of WHO-defined end-point consolidation in NT children are between three and 25 times higher than found elsewhere. The rates in children aged less than 12 months in the Central Australian region of the NT are the highest reported in the world.

Case ascertainment differences may partially explain the high rates of end-point consolidation we found. First, the entire NT hospitalised population was included, and hospital access has been improving over the past two decades; and, second, we included remote-living children. Comparable data on children in other disadvantaged populations are from studies in predominantly urban or peri-urban populations; children from rural and remote areas, with substantially less access to health services and with differing risk-factor profiles, would have been missed. These studies are from The Gambia (53.0/1000; aged 1–11 months), Philippines (13.5/1000; aged 6 weeks to 23 months), Indonesia (8.9/1000; aged 1.5–23 months), Chile (5.0/1000; aged 4–23 months), Fiji (4.3/1000; aged 1–59 months), Uruguay (up to 16.9/1000; aged 0–59 months) and South Africa (4.9/1000; aged 1.5–30 months). Furthermore, we included all chest x-rays taken within the first 3 days of admission. Studies including only x-rays taken on the day of admission may miss some cases, as clinical pneumonia may precede radiologically confirmed end-point consolidation.

Indigenous children in the NT may have a different risk factor and health care access profile, predisposing them to infection and hospitalisation. Underlying medical conditions or comorbidities (eg, malnutrition, anaemia and gastrointestinal infections) are common, and the high prevalence of low-birthweight infants is well documented. overcrowding, excessive pneumococci and non-typeable Haemophilus influenzae carriage rates in early infancy, and repeated infections leading to bron-
chietasis and chronic lung disease\textsuperscript{19,20} may play an important role in disease burden.

The major limitations of our study were its retrospective nature, the inclusion of hospitalised children only, and the use of a specific case definition as the primary outcome. The results should be viewed as an underestimate of the true incidence of disease. Most children with ALRI are treated in the community without a chest x-ray and antibiotic use is high. Measurement error leading to an overestimation of disease incidence is unlikely. All readers might have been influenced by knowing that they were examining x-rays from a high-risk population, the incidence of endpoint consolidation found was lower than that anticipated from a priori estimates based on data from Central Australia.\textsuperscript{21} Furthermore, the proportion of admitted episodes of care deemed positive for endpoint consolidation (20\% of all episodes of ALRI in which a chest x-ray was taken) was similar to that reported in other studies.\textsuperscript{5,11,22}

The rates of pneumonia in Indigenous children hospitalised in the NT are among the highest reported in the world. This is unacceptable in a wealthy country like Australia, and reducing this disease burden should be a national priority. Ongoing surveillance programs incorporating aetiological studies and innovative interventions are urgently required.

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COMPETING INTERESTS

Wyeth Vaccines provided funding for the study, but had no role in the design, data collection, analysis and interpretation, writing, or publication of the article. Kerry-Ann O’Grady has been a senior research officer on sponsored vaccine trials (GlaxoSmithKline, Wyeth, Merck Sharp & Dohme, MedImmune, and CSL) and a recipient of funds for epidemiological research (GlaxoSmithKline).

AUTHOR DETAILS

Kerry-Ann F O’Grady, GDipPH, Mappepid, PhD, NHMRC Post-Doctoral Training Fellow, Child Health Division\textsuperscript{1,2,3}

Debbie M Taylor-Thomson, BPharm, Senior Project Officer, Menzies School of Health Research\textsuperscript{1}

Anne B Chang, FRACP, MPH, PhD, Professor, and Head, Child Health Division\textsuperscript{1}

Paul J Torzillo, AM, MB BS, FRACP, FIFJCIM, Associate Professor, Department of Respiratory Medicine,\textsuperscript{4} and Director, Nganampa Health Service

Peter S Morris, MBBS, FRACP, PhD, Paediatrician and Deputy Leader, Child Health Division\textsuperscript{1,5}

Grant A Mackenzie, MB BS, PhD, Clinical Epidemiologist\textsuperscript{6}

Gavin R Wheaton, MBBS, FRACP, FCSANZ, Cardiologist, Women’s and Children’s Hospital Adelaide\textsuperscript{7}

Paul A Bauert, MB BS, FRACP, Head, Department of Paediatrics\textsuperscript{8}

Margaret P De Campo, FRANZCR, MPH, GDipEpiBiostats, Associate Professor, Department of Medicine\textsuperscript{9}

John F De Campo, FRANZCR, MHA, FRACMA, Professor, Department of Medicine\textsuperscript{9}

Alan R Ruben, MB BS, MAppEpipid, Paediatrician\textsuperscript{5}

1 Menzies School of Health Research, Charles Darwin University, Darwin, NT.

2 School of Population Health and Department of Paediatrics, University of Melbourne, Melbourne, VIC.

3 Vaccine and Immunisation Research Group, Murdoch Childrens Research Institute, Melbourne, VIC.

4 Royal Prince Alfred Hospital, University of Sydney, Sydney, NSW.

5 Northern Territory Clinical School, Flinders University, Darwin, NT.

6 Bacterial Diseases Program, Medical Research Council (UK) Laboratories, Fajara, The Gambia.

7 Department of Clinical Effectiveness, School of Medicine, Faculty of Health Sciences, Flinders University, Adelaide, SA.

8 Royal Darwin Hospital, Darwin, NT.

9 Bond University, Gold Coast, QLD.

Correspondence: k.o'grady@uq.edu.au

REFERENCES


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