International incidence of childhood cancer, 2001–10: a population-based registry study

Eva Steliarova-Foucher, Murielle Colombet, Lynn A G Ries, Florencia Moreno, Anastasia Dolya, Freddie Bray, Peter Hesseling, Hee Young Shin, Charles A Stiller, and the IICC-3 contributors *

Summary
Background Cancer is a major cause of death in children worldwide, and the recorded incidence tends to increase with time. Internationally comparable data on childhood cancer incidence in the past two decades are scarce. This study aimed to provide internationally comparable local data on the incidence of childhood cancer to promote research of causes and implementation of childhood cancer control.

Methods This population-based registry study, devised by the International Agency for Research on Cancer in collaboration with the International Association of Cancer Registries, collected data on all malignancies and non-malignant neoplasms of the CNS diagnosed before age 20 years in populations covered by high-quality cancer registries with complete data for 2001–10. Incidence rates per million person-years for the 0–14 years and 0–19 years age groups were age-adjusted using the world standard population to provide age-standardised incidence rates (WSRs), using the age-specific incidence rates (ASRs) for individual age groups (0–4 years, 5–9 years, 10–14 years, and 15–19 years). All rates were reported for 19 geographical areas or ethnicities by sex, age group, and cancer type. The regional WSRs for children aged 0–14 years were compared with comparable data obtained in the 1980s.

Findings Of 532 invited cancer registries, 153 registries from 62 countries, departments, and territories met quality standards, and contributed data for the entire decade of 2001–10. 385 509 incident cases in children aged 0–19 years occurring in 2.64 billion person-years were included. The overall WSR was 140·6 per million person-years in children aged 0–14 years (based on 284 649 cases), and the most common cancers were leukaemia (WSR 46·4), followed by CNS tumours (WSR 28·2), and lymphomas (WSR 15·2). In children aged 15–19 years (based on 100 860 cases), the ASR was 185·3 per million person-years, the most common being lymphomas (ASR 41·8) and the group of epithelial tumours and melanoma (ASR 39·5). Incidence varied considerably between and within the described regions, and by cancer type, sex, age, and racial and ethnic group. Since the 1980s, the global WSR of registered cancers in children aged 0–14 years has increased from 124·0 (95% CI 123·3–124·7) to 140·6 (140·1–141·1) per million person-years.

Interpretation This unique global source of childhood cancer incidence will be used for aetiological research and to inform public health policy, potentially contributing towards attaining several targets of the Sustainable Development Goals. The observed geographical, racial and ethnic, age, sex, and temporal variations require constant monitoring and research.

Funding International Agency for Research on Cancer and the Union for International Cancer Control.

Copyright © 2017 World Health Organization; licensee Elsevier. This is an Open Access article published under the CC BY-NC-ND 3.0 IGO license which permits users to download and share the article for non-commercial purposes, so long as the article is reproduced in the whole without changes, and provided the original source is properly cited. This article shall not be used or reproduced in association with the promotion of commercial products, services or any entity. There should be no suggestion that WHO endorses any specific organisation, products, or services. The use of the WHO logo is not permitted. This notice should be preserved along with the article’s original URL.

Introduction
Cancers rarely occur before age 20 years, and when they do, they raise a range of medical, psychological, ethical, and societal concerns. These distinct types of neoplasms require specific treatment paths. Furthermore, the extent of the cancer burden in this young population is unknown in many low-income and middle-income countries (LMICs), where data on cancer incidence are not collected. Even in the presence of population-based cancer registries, collection of information about childhood cancers is often neglected because they represent a small proportion of all cancers, additional data sources might be required, and the resulting statistics must be subjected to meticulous quality control because they are more sensitive to imprecision or missing information.

Since the publication of the International Incidence of Childhood Cancer, volume 1 (IICC-1) in 1988 and IICC-2 in 1998, no internationally comparable data on incidence patterns of childhood cancer have been published. To address this problem, the International Agency for Research on Cancer (IARC), in collaboration with the
Research in context

Evidence before this study

The worldwide incidence of cancer in children aged 0–14 years was reported in 1988 and 1998 in two volumes of the International Incidence of Childhood Cancer (IICC), which described incidence patterns in the 1970s and 1980s, respectively. We searched MEDLINE and participating registries for studies on the incidence of childhood cancer worldwide, with the search terms “childhood cancer”, “registry”, “incidence”, and “population”, in February, 2016, without language or publication date restrictions. We found that no globally comparable data on cancer types affecting children have been published since IICC-2, and no comparison of incidence patterns in children aged 15–19 years has been attempted on a global scale.

Added value of this study

This study provides an overview of the incidence of cancer in 2001–10 for children aged 0–19 years, based on data collected in 153 population-based cancer registries in 62 countries, departments, and territories on five continents. In addition to the sex, age, and tumour-specific incidence rates for the entire decade of 2001–10, which eliminated some high-quality registries that covered non-overlapping time periods. Analyses for the 0–14 years age group were based on a larger database (the paediatric dataset) than those for the 0–19 years age group (general dataset), because the paediatric dataset included data from the paediatric cancer registries not collecting data in children older than 15 years and these registries tended to have a wider coverage than the cancer registries used in the general dataset (appendix pp 2–4). All malignancies and non-malignant tumours of the CNS diagnosed before age 20 years (before age 15 years in most of the paediatric cancer registries) in the covered populations in 2001–10 were eligible for inclusion. The USA, non-malignant tumours were only registered from 2004 onwards, so they were excluded. The submission from each registry contained cancer and population data, information about registration coverage, practices, and data sources, and a short narrative describing registry operations. Cancer data included individual records of cases with codes for the following: sex; age; date of birth; date of incidence; and tumour sequence (ie, the numerical order of occurrence of the neoplasm), site, morphology, behaviour, laterality, and most valid basis of diagnosis. The 3rd edition of the International Classification of Diseases for Oncology was required for coding of the tumour site, morphology, behaviour, multiple primary tumours, and basis of diagnosis. Non-conforming coding systems were converted and tumours were classified centrally according to the 3rd edition of the International Classification of Childhood Cancer (ICCC-3). We requested each contributing cancer registry to provide total population

Methods

Study design and data sources

In this epidemiological, population-based registry study, we invited all population-based cancer registries that we identified among the membership of the IACR and from published literature in MEDLINE or other published sources, such as annual reports, in any language. The search terms used for the literature search were “population”, “childhood cancer”, “incidence”, and “registry”. Of the invited registries, we selected only those that had submitted their data and met standard data quality criteria. We included only registries covering 19 world regions or populations for 2001–10, we report an increase in the incidence of neoplasms since the 1980s in children aged 0–14 years. This study updates information on cancer incidence in children published almost 20 years ago and adds the first global overview of cancer incidence in young people aged 15–19 years. It comprehensively summarises the most recent and globally comparable data and presents information per sex, age group, geographical region, and tumour type.

Implications of all the available evidence

The wealth of information provided by this study constitutes a solid baseline to assess needs and define priorities in the area of paediatric oncology, supporting Sustainable Development Goal 3 to ensure healthy lives and promote wellbeing for all at all ages. The identified variations in cancer incidence according to region, sex, age group, and tumour type will provide a springboard for specific aetiological research. The shortage of high-quality local information should stimulate formation of new and more accurate data sources.
adjusted via direct standardisation. We calculated for the 0–14 years and 0–19 years age groups were regions and time periods, so all reported incidence rates purpose of this study was to make comparisons among registries, based on an assumption of regular linear interpolation (six registries) or extrapolation we estimated the covered population counts by ethnicity. In the absence of all required details, 2010, categorised by sex and age and, if relevant, race or area surveyed by a registry in each year from 2001 to 1980s, the time periods of the contributing registries differed both in length and starting years, because they the 1980s, the time periods of the contributing registries differed both in length and starting years, because they the 1980s, the time periods of the contributing registries differed both in length and starting years, because they the 1980s, the time periods of the contributing registries differed both in length and starting years, because they the 1980s, the time periods of the contributing registries differed both in length and starting years, because they the 1980s, the time periods of the contributing registries differed both in length and starting years, because they the 1980s, the time periods of the contributing registries differed both in length and starting years, because they the 1980s, the time periods of the contributing registries differed both in length and starting years, because they the 1980s, the time periods of the contributing registries differed both in length and starting years, because they the 1980s, the time periods of the contributing registries differed both in length and starting years, because they the 1980s, the time periods of the contributing registries differed both in length and starting years, because they the 1980s, the time periods of the contributing registries differed both in length and starting years, because they the 1980s, the time periods of the contributing registries differed both in length and starting years, because they the 1980s, the time periods of the contributing registries differed both in length and starting years, because they the 1980s, the time periods of the contributing registries differed both in length and starting years, because they the 1980s, the time periods of the contributing registries differed both in length and starting years, because they the 1980s, the time periods of the contributing registries differed both in length and starting years, because they the 1980s, the time periods of the contributing registries differed both in length and starting years, because they the 1980s, the time periods of the contributing registries differed both in length and starting years, because they the 1980s, the time periods of the contributing registries differed both in length and starting years, because they
The ASRs were higher in children aged 0–4 years and 15–19 years than in those aged 5–9 years and 10–14 years (table 2). ASRs also differed between regions, especially for the age groups with high ASRs (appendix pp 6, 9). In young people aged 15–19 years, the lowest ASR was observed in girls (incidence sex ratio was 1·17 in the 0–14 years age group and 1·14 in the 0–19 years age-group) and varied with age, region, and diagnostic group (table 2; appendix p 8). Across all regions, incidence was higher in boys than in girls (incidence sex ratio ranged from 1·1 to 1·4 in the 0–19 years age group; appendix p 8), except for the 15–19 years age group of Native Americans in the USA (incidence sex ratio of 0·9) and in east Asia (1·0), where girls had a higher incidence (ratios rounded to one decimal place; appendix p 8). The highest incidence sex ratio was observed in the age group of 5–9 years in south Asia (India; 1·7; appendix p 8). Sex-specific incidence varied by diagnostic group, with renal tumours and epithelial tumours more common in girls, with variations by age group (appendix p 8). Germ cell and gonadal tumours were also more common in girls than in boys in the 0–14 years age group (appendix p 8).

ASRs were higher in children aged 0–4 years and 15–19 years than in those aged 5–9 years and 10–14 years (table 2). ASRs also differed between regions, especially for the age groups with high ASRs (appendix pp 6, 9). In the 0–14 years group, overall WSRs varied from less than 100 per million person-years in sub-Saharan Africa, for Native American children in the USA, and in south Asia (India), to more than 150 per million person-years in young people aged 15–19 years (table 2). The ASR in young people aged 15–19 years was 185·3 per million person-years.

Incidence rates were slightly higher in boys than in girls (incidence sex ratio was 1·17 in the 0–14 years age group and 1·14 in the 0–19 years age-group) and varied with age, region, and diagnostic group (table 2; appendix p 8). Across all regions, incidence was higher in boys than in girls (incidence sex ratio ranged from 1·1 to 1·4 in the 0–19 years age group; appendix p 8), except for the 15–19 years age group of Native Americans in the USA (incidence sex ratio of 0·9) and in east Asia (1·0), where girls had a higher incidence (ratios rounded to one decimal place; appendix p 8). The highest incidence sex ratio was observed in the age group of 5–9 years in south Asia (India; 1·7; appendix p 8). Sex-specific incidence varied by diagnostic group, with renal tumours and epithelial tumours more common in girls, with variations by age group (appendix p 8). Germ cell and gonadal tumours were also more common in girls than in boys in the 0–14 years age group (appendix p 8).

ASRs were higher in children aged 0–4 years and 15–19 years than in those aged 5–9 years and 10–14 years (table 2). ASRs also differed between regions, especially for the age groups with high ASRs (appendix pp 6, 9). In the 0–14 years group, overall WSRs varied from less than 100 per million person-years in sub-Saharan Africa, for Native American children in the USA, and in south Asia (India), to more than 150 per million person-years in some subpopulations of North America and Europe, and in Oceania (figure 1A; appendix p 8). In young people aged 15–19 years, the lowest ASR was observed in 6 105 345 342 person-years, with the lowest being 0·8% in sub-Saharan Africa. Overall, 385 509 registered cases occurring in 2·64 billion person-years contributed to the analyses (tables 1, 2). Registration of non-malignant CNS tumours differed by registry, and was higher in the paediatric dataset than in the general dataset (table 1, 2). The overall WSRs were 140·6 per million person-years in children aged 0–14 years and 155·8 per million person-years in those aged 0–19 years (table 2). The ASR in young people aged 15–19 years was 185·3 per million person-years.

Incidence rates were slightly higher in boys than in girls (incidence sex ratio was 1·17 in the 0–14 years age group and 1·14 in the 0–19 years age-group) and varied with age, region, and diagnostic group (table 2; appendix p 8). Across all regions, incidence was higher in boys than in girls (incidence sex ratio ranged from 1·1 to 1·4 in the 0–19 years age group; appendix p 8), except for the 15–19 years age group of Native Americans in the USA (incidence sex ratio of 0·9) and in east Asia (1·0), where girls had a higher incidence (ratios rounded to one decimal place; appendix p 8). The highest incidence sex ratio was observed in the age group of 5–9 years in south Asia (India; 1·7; appendix p 8). Sex-specific incidence varied by diagnostic group, with renal tumours and epithelial tumours more common in girls, with variations by age group (appendix p 8). Germ cell and gonadal tumours were also more common in girls than in boys in the 0–14 years age group (appendix p 8).

ASRs were higher in children aged 0–4 years and 15–19 years than in those aged 5–9 years and 10–14 years (table 2). ASRs also differed between regions, especially for the age groups with high ASRs (appendix pp 6, 9). In the 0–14 years group, overall WSRs varied from less than 100 per million person-years in sub-Saharan Africa, for Native American children in the USA, and in south Asia (India), to more than 150 per million person-years in some subpopulations of North America and Europe, and in Oceania (figure 1A; appendix p 8). In young people aged 15–19 years, the lowest ASR was observed in 6 105 345 342 person-years, with the lowest being 0·8% in sub-Saharan Africa. Overall, 385 509 registered cases occurring in 2·64 billion person-years contributed to the analyses (tables 1, 2). Registration of non-malignant CNS tumours differed by registry, and was higher in the paediatric dataset than in the general dataset (table 1, 2). The overall WSRs were 140·6 per million person-years in children aged 0–14 years and 155·8 per million
south Asia (India), whereas the highest ASRs were seen in some predominantly white populations of North America, Europe, and Oceania (figure 1B; appendix p 6).

The range of tumour types varied markedly with age group (figure 2). In children aged 0–4 years, leukaemia represented 36.1% (45 849 of 127 096) of all cases; however, the proportion of leukaemia cases was 15.4% (15 520 of 100 860) in young people aged 15–19 years. Conversely, lymphomas represented 5.3% (6766 of 127 096) of cases in children aged 0–4 years, and 22.5% (22 740 of 100 860) of cases in those aged 15–19 years. CNS tumours were the second most frequent tumour type after leukaemia in children aged 0–4 years (21 804 [17.2%] of 127 096), 5–9 years (19 471 [26.3%] of 74 175), and 10–14 years (16 680 [20.0%] of 83 378; figure 2). Epithelial tumours and melanoma represented 0.9% of all cases in children aged 0–4 years (1197 of 127 096), but were the second most common tumour group in young people aged 15–19 years (21 480 [21.3%] of 100 860).

The most common sympathetic nervous system tumour across all age groups was neuroblastoma. It was most frequent in children aged 0–4 years (21 804 [17.2%] of 127 096), 5–9 years (19 471 [26.3%] of 74 175), and 10–14 years (16 680 [20.0%] of 83 378; figure 2). Epithelial tumours and melanoma represented 0.9% of all cases in children aged 0–4 years (1197 of 127 096), but were the second most common tumour group in young people aged 15–19 years (21 480 [21.3%] of 100 860).

Renal tumours were common in children aged 0–4 years (11 297 [8.9%] of 127 096), and their relative frequency decreased in older age groups, to 0.7% (756 of 100 860) in young people aged 15–19 years (figure 2). Bone tumours constituted 4.7% (13 320...
<table>
<thead>
<tr>
<th>Region</th>
<th>Leukaemia</th>
<th>Lymphomas</th>
<th>CNS tumours</th>
<th>Sympathetic nervous system</th>
<th>Retino-blastoma</th>
<th>Renal tumours</th>
<th>Hepatic tumours</th>
<th>Bone tumours</th>
<th>Soft tissue sarcomas</th>
<th>Germ cell and gonadal tumours</th>
<th>Epithelial tumours and melanoma</th>
<th>Other and unspecified</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>n</td>
<td>WSR</td>
<td>n</td>
<td>WSR</td>
<td>n</td>
<td>WSR</td>
<td>n</td>
<td>WSR</td>
<td>n</td>
<td>WSR</td>
<td>n</td>
<td>n</td>
</tr>
<tr>
<td>North Africa</td>
<td>5067</td>
<td>28.2</td>
<td>473</td>
<td>25.6</td>
<td>304</td>
<td>16.5</td>
<td>142</td>
<td>8.8</td>
<td>33</td>
<td>2.2</td>
<td>104</td>
<td>6.4</td>
</tr>
<tr>
<td>Sub-Saharan Africa</td>
<td>2060</td>
<td>12.5</td>
<td>1664</td>
<td>9.8</td>
<td>1041</td>
<td>63</td>
<td>411</td>
<td>27</td>
<td>563</td>
<td>37</td>
<td>1031</td>
<td>6.7</td>
</tr>
<tr>
<td>Central America and Caribbean</td>
<td>2145</td>
<td>45.5</td>
<td>920</td>
<td>18.0</td>
<td>1023</td>
<td>21.1</td>
<td>277</td>
<td>6.6</td>
<td>155</td>
<td>3.8</td>
<td>292</td>
<td>6.7</td>
</tr>
<tr>
<td>South America</td>
<td>6734</td>
<td>49.8</td>
<td>2319</td>
<td>16.2</td>
<td>3272</td>
<td>23.7</td>
<td>969</td>
<td>7.8</td>
<td>655</td>
<td>5.4</td>
<td>845</td>
<td>6.7</td>
</tr>
<tr>
<td>Canada</td>
<td>2780</td>
<td>55.5</td>
<td>949</td>
<td>16.2</td>
<td>1683</td>
<td>31.8</td>
<td>674</td>
<td>14.9</td>
<td>215</td>
<td>4.9</td>
<td>435</td>
<td>9.3</td>
</tr>
<tr>
<td>USA</td>
<td>297</td>
<td>31.5</td>
<td>76</td>
<td>75</td>
<td>140</td>
<td>144</td>
<td>49</td>
<td>5.7</td>
<td>15</td>
<td>1.8</td>
<td>49</td>
<td>5.7</td>
</tr>
<tr>
<td>Asian and Pacific Islanders</td>
<td>1294</td>
<td>43.9</td>
<td>378</td>
<td>12.1</td>
<td>596</td>
<td>19.9</td>
<td>214</td>
<td>7.7</td>
<td>107</td>
<td>3.9</td>
<td>117</td>
<td>4.2</td>
</tr>
<tr>
<td>Black</td>
<td>2994</td>
<td>31.3</td>
<td>1442</td>
<td>13.4</td>
<td>2541</td>
<td>26.2</td>
<td>881</td>
<td>10.2</td>
<td>444</td>
<td>5.3</td>
<td>968</td>
<td>10.9</td>
</tr>
<tr>
<td>White Hispanic</td>
<td>7366</td>
<td>65.4</td>
<td>1885</td>
<td>16.2</td>
<td>3249</td>
<td>28.6</td>
<td>933</td>
<td>8.6</td>
<td>619</td>
<td>5.7</td>
<td>811</td>
<td>7.4</td>
</tr>
<tr>
<td>White non-Hispanic</td>
<td>17709</td>
<td>54.5</td>
<td>5905</td>
<td>15.8</td>
<td>13009</td>
<td>38.2</td>
<td>4452</td>
<td>15.0</td>
<td>1219</td>
<td>4.2</td>
<td>3029</td>
<td>10.0</td>
</tr>
<tr>
<td>East Asia</td>
<td>5855</td>
<td>47.4</td>
<td>1730</td>
<td>12.7</td>
<td>2674</td>
<td>20.6</td>
<td>1236</td>
<td>12.0</td>
<td>506</td>
<td>5.1</td>
<td>542</td>
<td>5.1</td>
</tr>
<tr>
<td>South Asia*</td>
<td>2176</td>
<td>34.1</td>
<td>825</td>
<td>12.0</td>
<td>825</td>
<td>12.4</td>
<td>180</td>
<td>12.0</td>
<td>165</td>
<td>31.1</td>
<td>229</td>
<td>4.1</td>
</tr>
<tr>
<td>Southeast Asia</td>
<td>3925</td>
<td>52.7</td>
<td>745</td>
<td>9.5</td>
<td>997</td>
<td>12.9</td>
<td>312</td>
<td>4.4</td>
<td>404</td>
<td>6.0</td>
<td>378</td>
<td>5.4</td>
</tr>
<tr>
<td>West Asia</td>
<td>3459</td>
<td>45.2</td>
<td>1807</td>
<td>22.6</td>
<td>1918</td>
<td>24.5</td>
<td>727</td>
<td>10.1</td>
<td>326</td>
<td>4.6</td>
<td>534</td>
<td>7.3</td>
</tr>
<tr>
<td>Eastern Europe</td>
<td>526</td>
<td>44.3</td>
<td>2164</td>
<td>15.6</td>
<td>3793</td>
<td>30.0</td>
<td>1323</td>
<td>12.5</td>
<td>424</td>
<td>4.1</td>
<td>1062</td>
<td>9.8</td>
</tr>
<tr>
<td>Northern Europe</td>
<td>6987</td>
<td>49.1</td>
<td>2258</td>
<td>13.9</td>
<td>5821</td>
<td>38.9</td>
<td>1327</td>
<td>10.1</td>
<td>576</td>
<td>4.5</td>
<td>1235</td>
<td>9.3</td>
</tr>
<tr>
<td>Southern Europe</td>
<td>4200</td>
<td>51.1</td>
<td>1924</td>
<td>21.6</td>
<td>3156</td>
<td>37.6</td>
<td>1111</td>
<td>14.5</td>
<td>344</td>
<td>4.6</td>
<td>700</td>
<td>9.1</td>
</tr>
<tr>
<td>Western Europe</td>
<td>13630</td>
<td>50.7</td>
<td>4932</td>
<td>16.3</td>
<td>10628</td>
<td>38.0</td>
<td>3162</td>
<td>13.0</td>
<td>1082</td>
<td>4.5</td>
<td>2447</td>
<td>9.8</td>
</tr>
<tr>
<td>Oceania</td>
<td>2703</td>
<td>56.4</td>
<td>817</td>
<td>15.3</td>
<td>1285</td>
<td>26.2</td>
<td>500</td>
<td>11.4</td>
<td>187</td>
<td>4.3</td>
<td>384</td>
<td>8.5</td>
</tr>
<tr>
<td>Total</td>
<td>92085</td>
<td>46.4</td>
<td>3273</td>
<td>15.2</td>
<td>57955</td>
<td>28.2</td>
<td>4234</td>
<td>10.4</td>
<td>1832</td>
<td>4.5</td>
<td>15192</td>
<td>8.2</td>
</tr>
</tbody>
</table>

Regions are represented by the contributing cancer registries (appendix pp 3–4). Tumours classified by International Classification of Childhood Cancer, volume 3. Data are based on the paediatric dataset. WSR = age-standardised rate (world standard) per million person-years. *Comprising data from India only.

Table 3: Numbers of cases and age-standardised incidence of tumours in children aged 0–14 years, 2001–10, by main diagnostic group
<table>
<thead>
<tr>
<th>Region</th>
<th>Leukaemia</th>
<th>Lymphomas</th>
<th>CNS Tumours</th>
<th>Sympathetic Nervous System</th>
<th>Retino-Blastoma</th>
<th>Renal Tumours</th>
<th>Hepatic Tumours</th>
<th>Bone Tumours</th>
<th>Soft Tissue Sarcomas</th>
<th>Germ Cell and Gonadal Tumours</th>
<th>Epithelial Tumours and Melanoma</th>
<th>Other and Unspecified</th>
</tr>
</thead>
<tbody>
<tr>
<td>North Africa</td>
<td>184</td>
<td>26·4</td>
<td>258</td>
<td>37·0</td>
<td>97</td>
<td>13·9</td>
<td>5</td>
<td>100</td>
<td>143</td>
<td>85</td>
<td>12·2</td>
<td>40</td>
</tr>
<tr>
<td>Sub-Saharan Africa</td>
<td>129</td>
<td>19·0</td>
<td>183</td>
<td>27·0</td>
<td>41</td>
<td>6·0</td>
<td>2</td>
<td>0</td>
<td>0·3</td>
<td>16</td>
<td>24</td>
<td>3</td>
</tr>
<tr>
<td>Central America and Caribbean</td>
<td>366</td>
<td>22·8</td>
<td>479</td>
<td>29·9</td>
<td>226</td>
<td>14·1</td>
<td>11</td>
<td>0</td>
<td>1·6</td>
<td>22</td>
<td>14</td>
<td>3</td>
</tr>
<tr>
<td>North America</td>
<td>464</td>
<td>33·8</td>
<td>424</td>
<td>30·9</td>
<td>196</td>
<td>14·3</td>
<td>6</td>
<td>3</td>
<td>0·4</td>
<td>10</td>
<td>7</td>
<td>0</td>
</tr>
<tr>
<td>Canada</td>
<td>570</td>
<td>27·2</td>
<td>1210</td>
<td>57·8</td>
<td>519</td>
<td>24·8</td>
<td>15</td>
<td>1</td>
<td>0·7</td>
<td>34</td>
<td>16</td>
<td>20</td>
</tr>
<tr>
<td>USA</td>
<td>252</td>
<td>24·1</td>
<td>51</td>
<td>14·4</td>
<td>48</td>
<td>13·6</td>
<td>2</td>
<td>0</td>
<td>0·6</td>
<td>1</td>
<td>6</td>
<td>17</td>
</tr>
<tr>
<td>Asian and Pacific Islanders</td>
<td>1508</td>
<td>43·0</td>
<td>1306</td>
<td>37·3</td>
<td>558</td>
<td>15·9</td>
<td>19</td>
<td>0</td>
<td>0·5</td>
<td>38</td>
<td>11</td>
<td>50</td>
</tr>
<tr>
<td>Black</td>
<td>776</td>
<td>22·1</td>
<td>1385</td>
<td>39·4</td>
<td>542</td>
<td>15·4</td>
<td>29</td>
<td>0</td>
<td>0·8</td>
<td>86</td>
<td>24</td>
<td>44</td>
</tr>
<tr>
<td>White Hispanic</td>
<td>3845</td>
<td>29·5</td>
<td>7178</td>
<td>55·1</td>
<td>3176</td>
<td>24·4</td>
<td>101</td>
<td>0</td>
<td>0·8</td>
<td>199</td>
<td>15</td>
<td>154</td>
</tr>
<tr>
<td>White Non-Hispanic</td>
<td>1683</td>
<td>31·5</td>
<td>1037</td>
<td>19·4</td>
<td>811</td>
<td>15·2</td>
<td>36</td>
<td>0</td>
<td>0·7</td>
<td>53</td>
<td>10</td>
<td>90</td>
</tr>
<tr>
<td>East Asia</td>
<td>565</td>
<td>21·8</td>
<td>346</td>
<td>13·3</td>
<td>263</td>
<td>10·2</td>
<td>7</td>
<td>0</td>
<td>0·3</td>
<td>18</td>
<td>0</td>
<td>7</td>
</tr>
<tr>
<td>Southeast Asia</td>
<td>767</td>
<td>26·5</td>
<td>418</td>
<td>14·4</td>
<td>215</td>
<td>7·4</td>
<td>10</td>
<td>0</td>
<td>0·3</td>
<td>28</td>
<td>10</td>
<td>63</td>
</tr>
<tr>
<td>West Asia</td>
<td>786</td>
<td>33·7</td>
<td>1222</td>
<td>52·4</td>
<td>513</td>
<td>22·0</td>
<td>23</td>
<td>0</td>
<td>0·0</td>
<td>26</td>
<td>11</td>
<td>25</td>
</tr>
<tr>
<td>Eastern Europe</td>
<td>1048</td>
<td>21·8</td>
<td>2047</td>
<td>42·6</td>
<td>1001</td>
<td>20·8</td>
<td>41</td>
<td>0</td>
<td>0·9</td>
<td>80</td>
<td>17</td>
<td>51</td>
</tr>
<tr>
<td>Northern Europe</td>
<td>512</td>
<td>25·5</td>
<td>954</td>
<td>47·5</td>
<td>726</td>
<td>36·2</td>
<td>17</td>
<td>0</td>
<td>0·8</td>
<td>26</td>
<td>13</td>
<td>18</td>
</tr>
<tr>
<td>Southern Europe</td>
<td>650</td>
<td>30·0</td>
<td>1562</td>
<td>72·1</td>
<td>655</td>
<td>30·2</td>
<td>27</td>
<td>1</td>
<td>1·2</td>
<td>36</td>
<td>17</td>
<td>23</td>
</tr>
<tr>
<td>Western Europe</td>
<td>753</td>
<td>28·4</td>
<td>1466</td>
<td>55·4</td>
<td>814</td>
<td>30·7</td>
<td>19</td>
<td>0</td>
<td>0·7</td>
<td>38</td>
<td>14</td>
<td>30</td>
</tr>
<tr>
<td>Oceania</td>
<td>577</td>
<td>33·0</td>
<td>918</td>
<td>52·5</td>
<td>294</td>
<td>16·8</td>
<td>12</td>
<td>0</td>
<td>0·7</td>
<td>23</td>
<td>13</td>
<td>20</td>
</tr>
<tr>
<td>Total</td>
<td>15520</td>
<td>28·5</td>
<td>22400</td>
<td>41·8</td>
<td>10822</td>
<td>19·9</td>
<td>391</td>
<td>0</td>
<td>0·7</td>
<td>756</td>
<td>1·4</td>
<td>675</td>
</tr>
</tbody>
</table>

Regions are represented by the contributing cancer registries (appendix pp 3–4). Tumours classified by International Classification of Childhood Cancer, volume 3. Data are based on the general dataset. ASR=age-specific rate per million person-years.

*Comprising data from India only.

Table 4: Numbers of cases and age-specific incidence of tumours in young people aged 15-19 years, 2001-10, by main diagnostic group.
of 284649) of all cancers in children aged 0–14 years, and 7.8% (7851 of 100860) in those aged 15–19 years (figure 2; appendix p 10). Soft tissue sarcomas were present in a similar proportion of cases in children aged 0–14 years and those aged 15–19 years in most regions (table 3, 4; figure 2; appendix p 10). In sub-Saharan Africa, 46% (579 of 1262) of all soft tissue sarcomas in children aged 0–14 years were Kaposi’s sarcoma compared with 57% (119 of 208) in those aged 0–19 years, whereas in all the other regions Kaposi’s sarcoma represented less than 1% of all soft tissue sarcomas in children younger than 15 years and a maximum of 2% (14 of 592) in black children in the USA (data for all regions not shown). Germ cell and gonadal tumours were rare in children younger than 15 years (10 200 [3.6%] of 284649; table 3), and more common in those aged 15–19 years (12 105 [12.0%] of 100860; table 4). The group of epithelial neoplasms and melanoma comprises several specific types of carcinomas (adrenocortical, thyroid, nasopharyngeal, skin), all other types of carcinoma (except those occurring in kidney, liver, and gonads), and melanoma. This group constituted 3.8% (10 679 of 284649) of all tumours in children aged 0–14 years and 21.3% (21 480 of 100860) in those aged 15–19 years (appendix p 10). The group of other and unspecified tumours comprised 0.9% (2469 of 284649) of all cases in children aged 0–14 years and 1.5% (1500 of 100860) of all cases in those aged 15–19 years. In 22 registries no tumours were classified in this category; other and unspecified tumours comprised 5% or more of all tumours in ten registries, including one registry for which 12% (68 of 561) of all tumours were classified as other and unspecified.

In children aged 0–14 years, the leading cancers in all regions combined were leukaemia, followed by CNS tumours, and then lymphomas, and this ranking was
observed in most world regions (table 3). The exception to this pattern was seen in the two African regions (appendix p 10). Incidence rates of all tumour types varied considerably between the regions (figure 3). The ratios of highest to lowest regional WSR were higher than 5·0 for CNS tumours (6·2), germ cell and gonadal tumours (5·9), neuroblastoma (5·6), epithelial tumours and melanoma (5·5), and leukaemia (5·2).

In young people aged 15–19 years, lymphomas were the most common cancers in all regions combined, followed by epithelial tumours and melanoma (table 4; figure 4). However, leukaemia was the most common neoplasm in this age group in South America, Native American and white Hispanic children in the USA, south Asia (India), and southeast Asia (table 4; figure 4). In Oceania, white non-Hispanic children in the USA, in east Asia, and in Central America and the Caribbean the most common cancer type was the group of epithelial cancers and melanoma (table 4; figure 4). Geographical variations in incidence, expressed as the ratio of the highest to the lowest ASR across regions, were 5·0 or higher for hepatic tumours (7·8), germ cell tumours (6·4), CNS tumours (6·0), lymphomas (5·4), epithelial tumours and melanoma (5·2), and soft tissue sarcomas (5·0).

From the 1980s to 2001–10, the overall WSR for all tumours in children aged 0–14 years increased from 124·0 per million person-years (123·3–124·7) in the 1980s to 140·6 per million person-years (140·1–141·1) in 2001–10. The increase was seen in all regions except sub-Saharan Africa, and was smallest in Central America and the Caribbean (WSR 125·3 [95% CI 125·5–132·5] in 2001–10) and highest in southeast Asia (92·0 [95% CI 89·2–94·8]; 119·8 [95% CI 117·3–122·3]). In sub-Saharan Africa, a decrease was noted between the 1980s (119·8 [95% CI 117·3–122·3]) and 2001–10 (56·3 [95% CI 55·1–57·5]; figure 5; appendix p 7).

**Discussion**

Using data provided by 153 high-quality cancer registries, we report internationally comparable incidence rates of cancer in children aged 0–19 years during 2001–10. The incidence of cancer in children aged 0–14 years was 140·6 per million person-years, and in those aged 0–19 years was 155·8 per million person-years. We report considerable variations in incidence by diagnostic group, sex, age group, and world region. We have also shown that the overall WSR in children aged 0–14 years has increased between the preceding comparable publication covering the 1980s, and 2001–10. To our knowledge, this report presents the best available information on cancer incidence for the given period and age group.

We chose WSR as a relative estimate of cancer burden, but it should not be used for estimating numbers of cases in a population, because the value of WSR depends on the choice of the standard population. The world standard population has been used in previous international publications, choosing the same standard thus ensures comparability between populations and periods.

Although 20% of the cancers occurring in young people aged 15–19 years were epithelial tumours (the most prevalent histology types in adults) or melanoma, this age group also had high proportions of lymphomas, leukaemias, germ cell tumours, and sarcomas. Therefore, ICCC-3 seems to be well adapted to reporting incidence in this age group, because the named tumour groups represent the main diagnostic categories in ICCC-3. We found that the incidence of cancer in young people aged 15–19 years was 185·3 per million person-years, based on 100860 cases.

Because not all the contributing paediatric cancer registries could provide data for young people aged 15–19 years, the analyses for children aged 0–14 years were based on a different dataset (paediatric dataset) than were the analyses for children aged 0–19 years or those aged 15–19 years (general dataset). The two resulting datasets
gave different estimates of incidence for the (common) age group of 0–14 years, and this incidence was 5% lower in the paediatric than in the general dataset. This difference can be explained by variations in population coverage between the registries, and possibly distinctive data sources used by the two types of registries (ie, those that included data for young people aged 15–19 years and those that did not), as was shown previously in an analysis of data on childhood cancer in Europe.4 Although the results presented from both datasets are based on the best data available for each age group, they also emphasise the importance of quality assurance, particularly when dealing with a rare disease such as childhood cancer. We selected the contributing registries because they provided quality-assured data for the entire decade of 2001–10. Inevitably, the reported rates were influenced by this selection; however, they provide the best and unique comparable global incidence estimates for the given period because they are not affected by intermittent contributions for parts of the target period.

A particular strength of our study comes from separate presentation of the incidence patterns for the five racial and ethnic groups distinguished in the US data, despite the difficulties in classifying the US population into racial and ethnic groups due to population mixing, migration, and self-declaration of ethnicity.11 Unfortunately, the ethnic differences within multi-ethnic populations of Europe, Canada, and some other regions could not be readily studied. In other countries (South Africa, Israel, Australia) it was possible to gather incidence data for specific ethnicities within the population. For the purpose of simplicity these data were not included in this paper; however, they are available online on the IICC-3 website.

Our study improved the overall data quality in all participating cancer registries, particularly in 72 of 153 whose datasets were not acceptable at the beginning of the study. The study raised the registries’ awareness of additional data sources (haematology, ophthalmology, orthopaedics, dermatological, neurology, endocrinology, and paediatric clinics) and of specific quality control. We required a high level of quality and completeness for our study, because small errors or omissions would have had a large impact on the resulting incidence rates. Approximately 30 cancer registries dropped out of the study, irrespective of the quality of their data. Such loss is especially regrettable in LMICs, where data are sparse and health policy has to be based on information from other countries. Data collected on more than 40 million person-years in 12 cancer registries (Canada [New Brunswick, Prince Edward Island], Denmark, Finland, Germany [Berlin, Brandenburg, Bremen, Mecklenburg-Vorpommern, Sachsen-Anhalt, and the Free States Saxony and Thuringia], and Singapore) could not be used because these registries were bound by requirements of a specific administrative procedure to approve provision of their data.

Although the cancer registries follow international guidelines, we observed wide variation in registration techniques, access to data sources, case eligibility criteria, application and interpretation of registration standards, and coding systems. Non-malignant tumours represented 0–10% of the total cases and affected international comparability. Non-malignant tumours were eligible for registration more commonly in Europe than in other parts of the world. Other practices reduced data quality (eg, restricted or no access to pathology reports), completeness (eg, restricted or no access to identifiable death certificates), or timeliness (eg, third-party data encryption). In some registries cases were coded as
microscopically verified only if the pathology report could be reviewed by registry staff. In others, precise morphology codes were not available for a large proportion of microscopically verified diagnoses. Registries are often under-resourced, and large investments for a small proportion of cases in general cancer registries might not be seen as a priority; however, these variations highlight the need for support to enable production of high-quality and comparable data.

A high proportion of specified diagnoses cannot be ensured without available specialised diagnostic facilities. The category of unspecified morphology might have included misclassified specific cancers, but in the absence of precise documentation it might have also included cases that were not malignant or cancers that did not occur in children.

Although the differences in sex ratios by diagnostic group and age might have to a large extent reflected true differences in disease occurrence, the differences by geographical region could have more readily reflected sociocultural customs whereby boys are favoured over girls to seek medical attention when sick. The potential for such selective treatment in India and in some African populations was supported by our registry contacts in those regions when we inquired about a possibility of inequality in seeking health care.

The registries with no access to a national database of all causes of death might not have registered the cancers that were only discovered at the time of death, even though the proportion of missed cases might be low in childhood populations. In some countries death certification is non-existent whereas in others access to these records is hindered by data protection laws.

Factors external to the registry operations should also be considered. Higher socioeconomic level of development, better treatment facilities, and unobstructed access to data sources are linked to more advanced registration systems and better data quality. The registries operating in some LMICs were particularly affected by missing or infrequent official population estimates, and these data needed to be estimated by interpolation or extrapolation, influencing the reported rates. The low incidence rates observed in low-resource regions might in part have resulted from underdiagnosis, but our analysis could not quantify its extent. Treatment abandonment, common in LMICs, might also have led to under-registration if identified patients who refused treatment were not sufficiently well described (eg, missing dates of place of residence). Under-ascertainment of diagnosed cases might have resulted from administrative restrictions of access to medical files, political or social instability, competing needs, and inadequate political will, causing a dearth of resources, shortage or volatility of registry personnel, loss of personnel expertise, and missing or broken links with relevant data sources. Overestimates of incidence might have occurred in areas with superior treatment facilities if the place of permanent residence could not be correctly determined for registered patients, and national coverage would neutralise such artefactual regional differences within a country. Accurate incidence rates are also difficult to obtain in ethnic minorities, such as the Native American population in the USA, possibly because of imprecise classification of the at-risk population, as well as patients with cancer. Cancer statistics in high-income countries might be influenced by overdiagnosis of some cancers detected by non-invasive imaging and screening tests, including neuroblastoma, thyroid cancer, melanoma, and kidney cancer.

Although a complete assessment of time trends in incidence falls outside the scope of this report, our data showed that the overall incidence of registered neoplasms in childhood increased between the 1980s and the 2000s. Our comparison is certainly influenced by the composition of the two data pools (different registries have contributed to the two compared periods); however, the increase in the overall rates is clear. The reasons might include improved diagnosis (eg, imaging of brain tumours in high-income countries and less under-diagnosis in some parts of world). Cancer registration is also developing constantly, and more effective ascertainment techniques might have been used since the 1980s. Future detailed analyses of time trends by cancer type and subpopulations will bring more clarity to the interpretation of these secular changes.

The reduction of cancer incidence in sub-Saharan Africa probably had two main causes. First, this IICC-3 study included data from the national childhood cancer registry of South Africa, which contributed 70% of all cases for this region and reported very low incidence rates. Of the five datasets classified as sub-Saharan African in IICC-2, only one (from Namibia) reported a lower overall incidence rate than the South African registry in IICC-3. The possible reasons for the low rates in South Africa are discussed elsewhere. Second, the implementation of antiretroviral therapy in some areas affected by HIV has contributed to a decrease in incidence of Kaposi’s sarcoma. This cancer type represented more than a third of the total cancer incidence in Kampala, Uganda, in the IICC-2 (WSR 182.7 per million person-years). The incidence of Kaposi’s sarcoma in Kampala halved between the IICC-2 and this IICC-3 study, and a decreasing trend between the 1990s and the 2000s was also noted in a Kampala registry study.

Leukaemia, the most common cancer in children worldwide, had the largest impact on total cancer incidence. The strikingly low incidence of leukaemia in the African regions, notably in sub-Saharan Africa, might give some clues as to the cause of this cancer. Even though strong underdiagnosis of leukaemia is suspected in sub-Saharan Africa, leukaemia was also less commonly diagnosed in black children in the USA than in the other US ethnic groups; however, whether
leukaemia might be underdiagnosed as a result of reduced access to health care associated with lower economic status in this group is unclear. In children aged 0–14 years, the highest leukaemia rates were in Hispanic white children in the USA (40% of total incidence). The Native American component of Hispanic ancestry was a presumed risk factor, based on the observations that known risk alleles at loci identified in genome-wide association studies of European-ancestry populations in CDKN2A, PIP4K2A, CEBPE, and ARID5B were all significantly associated with Native American ancestry. Comparatively high incidence rates and the largest proportion of leukaemia among all cancers were reported in southeast Asia. A link to the massive use of pesticides in this world region to protect crops and increase yields should be examined in specific studies, since exposure to pesticides has been associated with leukaemia risk.

The male predominance among patients with lymphoma is probably a result of innate sex differences in susceptibility, but with increasing age other factors might play a role, such as the increased risk of HIV infection in boys compared with girls. The highest incidence of lymphomas worldwide has been reported in the Mediterranean region (as shown in our study and others), and in HIV-infected populations, with B-cell non-Hodgkin lymphoma an AIDS-defining malignancy. Burkitt’s lymphoma is endemic and common in some sub-Saharan African childhood populations, in which HIV infection can further accentuate its incidence. Hodgkin’s lymphoma is rarer in populations of south and east Asia than in other parts of the world; the undergoing socioeconomic changes might gradually result in a similarly high incidence to that observed in Western populations.

The highest incidence of all CNS tumours was noted in high-income countries, which is clearly related to the wide availability of diagnostic facilities. The lower incidence rates in LMICs most probably (and proportionately) reflect poor access to neuroimaging facilities (eg, low number of CT or MRI scans, long waiting lists, and prohibitive costs of diagnostics tests). This poor access causes delay in diagnosis and possibly underdiagnosis of brain tumours. Comparability of incidence of CNS tumours across the continents would be greatly improved if all registries attempted to collect information on non-malignant CNS tumours, at least in children. This study describes the global cancer incidence patterns in children younger than 20 years for 2001–10, providing an update to comparable information that is now almost 20 years old. Despite possible artefacts influencing data availability, quality, and comparability, the size of the studied populations and the observed differences in our study suggest that our data are sufficiently robust for international comparisons of childhood cancer occurrence, and provide useful pointers for further studies. The collected data can be used for further research, including continued development of childhood-specific registration standards and guidelines, detailed studies in subpopulations and by tumour subtype, and global estimates of cancer indicators. This report constitutes a springboard for attaining several targets of the Sustainable Development Goal aiming at ensuring healthy lives and promoting wellbeing of all at all ages. High-quality data over time is vital to devising cancer control mechanisms in childhood populations worldwide. Local data that are internationally comparable are therefore indispensable for aetiological research and for effective health policy actions. Sharing of data internationally should not impose an unaffordable burden on data acquisition and validation in coordinated studies. To secure data availability and quality, constant support of cancer registration is required at local, national, and international levels.

Contributors
ES-F, LAGR, FM, PH, HYS, and CAS designed the study. ES-F, AD, and MC, in collaboration with LAGR, FM, PH, HYS, and CAS acquired the data for the study. MC did the analyses. ES-F and MC produced the tables and figures. AD and ES-F did the literature search. ES-F drafted the manuscript, which was reviewed, modified, and approved by ES-F, LAGR, FM, AD, FB, PH, HYS, and CAS.

Declaration of interests
We declare no competing interests.

Acknowledgments
International Incidence of Childhood Cancer, volume 3 is supported by the International Agency for Research on Cancer and the Union for International Cancer Control. The cooperation of all cancer registries and members of the International Association of Cancer Registries is gratefully acknowledged.

References


