Erratum to "Averting the legacy of kidney disease-Focus on childhood" [Nephrol. Ther. 12 (1) (2016) 1-5].

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Erratum


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Table 2

Etiology of chronic kidney disease in children.

<table>
<thead>
<tr>
<th>CKD Etiology</th>
<th>Percentage (range)</th>
<th>ESRD Etiology</th>
<th>Percentage (range)</th>
</tr>
</thead>
<tbody>
<tr>
<td>CAKUT</td>
<td>48–59</td>
<td>CAKUT</td>
<td>34–43</td>
</tr>
<tr>
<td>GN</td>
<td>5–14</td>
<td>GN</td>
<td>15–29</td>
</tr>
<tr>
<td>HN</td>
<td>10–19</td>
<td>HN</td>
<td>12–22</td>
</tr>
<tr>
<td>HUS</td>
<td>2–6</td>
<td>HUS</td>
<td>2–6</td>
</tr>
<tr>
<td>Cystic</td>
<td>5–9</td>
<td>Cystic</td>
<td>6–12</td>
</tr>
<tr>
<td>Ischemic</td>
<td>2–4</td>
<td>Ischemic</td>
<td>2</td>
</tr>
</tbody>
</table>

Rare causes include congenital NS, metabolic diseases, cystinosis. Miscellaneous causes depend on how such entities are classified. CAKUT: congenital anomalies of the kidney and urinary tract; GN: glomerulonephritis; HN: hypertension; HUS: hemolytic uremic syndrome. From Harambat et al. CKD data are from NAPRTCS, the Italian Registry and the Belgian Registry. ESRD data are from ANZDATA, ESPN/ERA-EDTA, UK Renal Registry and the Japanese Registry.

Correction notice:

In proofing, the abbreviation HN was changed to hypertension. HN stands for hereditary nephropathy and should have been defined as such, when a legend was added in proofing. However, due to the change in the abbreviation to hypertension, hypertension was erroneously inserted in the legend. We apologize for this error.

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