Association of Nephrotic Syndrome with Intestinal Lymphangiectasia

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The occurrence of exudative enteropathy in children with the nephrotic syndrome was first reported by Nüssle et al. in 1961, but no intestinal pathology has previously been described in nephrotic children, whether with or without protein-losing enteropathy. The finding in small bowel biopsy specimens of intestinal lymphangiectasia, in 4 of 7 cases of nephrotic syndrome, suggests that this association might not be uncommon.

Material and Methods

Seven children aged from 10 months to 10 years, admitted to the Clínica Pediátrica Universitária de Lisboa with nephrotic syndrome, were submitted to intestinal biopsy, using the paediatric version of the Crosby-Kugler capsule (Crosby and Kugler, 1957), and following a technique previously described (Salazar de Sousa and Cunha, 1967). The specimens were fixed in buffered formalin and stained with haematoxylin-eosin.

In 2 patients the intestinal loss of plasma protein was studied with 131I albumin (RISA, Abbott Laboratories) according to the method of Jeejeebhoy and Coghill (1961).

Results

Clinical and biochemical details of Cases 1–4 are summarized in Tables I and II. The results of renal biopsy and intestinal biopsy are given in Table I, and illustrated in Fig. 1–4.

Table III gives the results of RISA studies in Cases 2 and 4. Protein loss from the gut was 13% and 8% per day, compared with 2% per day in a control case.

Cases 5–7 need be mentioned only briefly.

Case 5. A 6-year-old boy with a pure nephrosis that responded well to steroid therapy.

Case 6. A 4-year-old girl with a pure nephrosis that responded well to steroid therapy.

Case 7. A 10-month-old boy with a familial form of nephrosis. A (surgical) renal biopsy showed a lobular glomerulonephritis. He proved non-responsive to steroids.

Jejunal biopsy in Cases 5, 6, and 7 showed a marked oedema of the lamina propria as the sole abnormality.

Discussion

Though intestinal protein loss during the acute phase of the nephrotic syndrome is an accepted fact (Nüssle et al., 1961; Royer et al., 1963), there is no previous reference to the association of nephrosis and intestinal lymphangiectasia. Nor, in two recent review articles of protein-losing enteropathy (Dawson, 1965; Waldmann, 1966), is the nephrotic syndrome referred to in the list of diseases in which intestinal lymphangiectasia may be found.

Of 7 children with the nephrotic syndrome submitted to intestinal biopsy, the histological picture was typical of intestinal lymphangiectasia in 2 (Cases 2 and 4), and was consistent with this diagnosis in two others (Cases 1 and 3). In the authors’ experience of over 20 intestinal biopsies in children, lymphatic dilatation of such a magnitude as that seen in Cases 1 and 3 has never been seen.

In the 2 cases (Cases 2 and 4) in which the metabolism of albumin was studied with RISA, the faecal loss of labelled albumin was found to be much increased.

We conclude that since protein-losing enteropathy is a frequent occurrence during the acute phase of the nephrotic syndrome, in at least some of these cases, intestinal lymphangiectasia is the responsible factor.

The finding of intestinal lymphangiectasia in nephrotic children has some interesting therapeutic implications. Long-chain fatty acids are largely converted to chylomicrons which are transported by the intestinal lymph, whereas short-chain fatty acids of 10 carbon atoms or less are mainly found in the portal blood (Peterson, 1963). With a low fat
### TABLE I

#### Clinical and Pathological Findings in 4 Cases of Nephrotic Syndrome

<table>
<thead>
<tr>
<th>Case No.</th>
<th>Age (yr.) and Sex</th>
<th>Duration of Nephrosis</th>
<th>Oedema</th>
<th>BP (mm. Hg)</th>
<th>Renal Biopsy (percutaneous)</th>
<th>Intestinal Biopsy (peroral)</th>
<th>Outcome</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>10M</td>
<td>1 yr.</td>
<td>++</td>
<td>115/80</td>
<td>Thickened capillary basement membrane</td>
<td>Several ectatic lymphatics in villi and among crypts; ratio of normal villi to villi with abnormal lymphatics 5:1 (Fig. 1)</td>
<td>Complete clinical and biochemical remission after steroid therapy</td>
</tr>
<tr>
<td>2</td>
<td>8M</td>
<td>8 mth.</td>
<td>++</td>
<td>120/80</td>
<td>Thickened capillary basement membrane; marked endothelial cell proliferation, some hyalinized glomeruli</td>
<td>Typical lymphangiectasia (Fig. 2)</td>
<td>Steroids and 6-mercaptopurine ineffective; oedema cleared after hydrochlorothiazide and spironolactone, but blood chemistry unchanged</td>
</tr>
<tr>
<td>3</td>
<td>4M</td>
<td>3 wk.</td>
<td>++</td>
<td>105/75</td>
<td>—</td>
<td>Lymphatic dilatation in several villi; ratio of normal villi to villi with abnormal lymphatics 7:1 (Fig. 3)</td>
<td>Complete clinical and biochemical remission after steroid therapy</td>
</tr>
<tr>
<td>4</td>
<td>7M</td>
<td>3 mth.</td>
<td>++</td>
<td>110/80</td>
<td>—</td>
<td>Typical lymphangiectasia</td>
<td>Steroids ineffective; oedema fluctuated; biochemically unchanged</td>
</tr>
</tbody>
</table>

**FIG. 1.—Case 1: dilated lymphatics in the villi and among the crypts.**

### Urine and Blood Findings

<table>
<thead>
<tr>
<th>Case No.</th>
<th>Protein Output (g. day)</th>
<th>Deposit</th>
<th>Hb (g. 100 ml.)</th>
<th>BSR (mm. 1 hr.)</th>
<th>BUN (mg. 100 ml.)</th>
<th>Creatinine Clearance (ml./min.)</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>1</td>
<td>Nil</td>
<td>12·3</td>
<td>61</td>
<td>29</td>
<td>23·5</td>
</tr>
<tr>
<td>2</td>
<td>1</td>
<td>WBC+</td>
<td>8·2</td>
<td>79</td>
<td>143</td>
<td></td>
</tr>
<tr>
<td>3</td>
<td>3</td>
<td>Nil</td>
<td>10·9</td>
<td>56</td>
<td>64</td>
<td>95·7</td>
</tr>
<tr>
<td>4</td>
<td>7</td>
<td>Few RBC; few casts</td>
<td>11·5</td>
<td>69</td>
<td>17</td>
<td>88·6</td>
</tr>
</tbody>
</table>
**Fig. 2.**—Case 2: dilated lymphatics give the villi a ballooned appearance.

**Fig. 3.**—Case 3: (a) dilated lymphatic at the tip of the villous; (b) dilated lymphatic following the villous axis.

**Fig. 4.**—Case 4: tip of villi occupied by dilated lymphatics.

### Cases of Nephrotic Syndrome

<table>
<thead>
<tr>
<th>Total Serum Protein (g./100 ml.)</th>
<th>Albumin (g./100 ml.)</th>
<th>( \alpha_1 ) (% of total)</th>
<th>( \alpha_2 ) (% of total)</th>
<th>( \beta ) (% of total)</th>
<th>( \gamma ) (% of total)</th>
<th>Total Serum Cholesterol (mg./100 ml.)</th>
<th>Esterified Serum Cholesterol (mg./100 ml.)</th>
</tr>
</thead>
<tbody>
<tr>
<td>4.25</td>
<td>16</td>
<td>5</td>
<td>56</td>
<td>11</td>
<td>11</td>
<td>666</td>
<td>483</td>
</tr>
<tr>
<td>4.5</td>
<td>14</td>
<td>5</td>
<td>60</td>
<td>9</td>
<td>11</td>
<td>691</td>
<td>391</td>
</tr>
<tr>
<td>4.0</td>
<td>38</td>
<td>3</td>
<td>35</td>
<td>18</td>
<td>7</td>
<td>456</td>
<td>308</td>
</tr>
<tr>
<td>5.0</td>
<td>12</td>
<td>6</td>
<td>55</td>
<td>11</td>
<td>15</td>
<td>756</td>
<td>516</td>
</tr>
</tbody>
</table>
diet or with a diet in which the fat predominantly contains medium-chain fatty acids, the chyle flow may be reduced and the leakage of proteins into the gut minimized (Jeffries, Chapman, and Sleisenger, 1964).

Summary

Seven children with nephrotic syndrome were submitted to intestinal biopsy. In 2, the histological picture was typical of intestinal lymphangiectasia and in a further 2 it was consistent with this diagnosis. In 2 of the patients with lymphatic dilatation in the intestinal mucosa, the metabolism of albumin was studied with RISA and the faecal loss of labelled albumin was found to be markedly increased.

It is concluded that in at least some nephrotic children, intestinal lymphangiectasia is responsible for an associated protein-losing enteropathy.

### References