Congenital syphilis and glomerulonephritis with evidence for immune pathogenesis

J. WIGGELINKHUIZEN, R. O. C. KASCHULA, C. J. UYS, R. H. KUIJTEN,* and J. DALE
From the Red Cross War Memorial Children's Hospital, Rondebosch, Cape, and the Departments of Paediatrics and Pathology, University of Cape Town, South Africa

Wiggelinkhuizen, J., Kaschula, R. O. C., Uys, C. J., Kuijten, R. H., and Dale, J. (1973). Archives of Disease in Childhood, 48, 375. Congenital syphilis and glomerulonephritis with evidence for immune pathogenesis. In 3 infants with congenital syphilis the dominant clinical manifestation of syphilitic kidney disease was the nephrotic syndrome. Mesangioendothelial proliferation was present in 2 cases and mixed proliferative glomerulonephritis with crescent formation in the third. The severity of the clinical and histopathological abnormalities could be related to the apparent duration of the illness.

In all 3 cases immune complex deposition could be shown within and along the epithelial aspect of the glomerular basement membrane on light, electron, and immunofluorescent microscopy. These features, together with a reduced total serum haemolytic complement, suggest an immune pathogenesis of the glomerulonephritis associated with early congenital syphilis.

Congenital syphilis, though uncommonly associated with kidney disease, is nevertheless a well-documented cause of the nephrotic syndrome in young infants. We present the clinical, laboratory, histopathological, electron microscopy, and immunofluorescent findings of the nephropathy before, during, and after treatment of early congenital syphilis. Evidence suggestive of an immune pathogenesis for the nephropathy of acquired syphilis was recently published (Braunstein et al., 1970; Bhorade et al., 1971). Our findings suggest a similar immunopathogenetic mechanism for the renal lesions of congenital syphilis.

Case reports

Case 1. A coloured girl (mixed race) aged 4 months was admitted for oedema of 2 months' duration. Pregnancy and delivery had been uneventful but the mother had received no antenatal care. No immunizations had been given. The infant had gross generalized oedema with ascites, a healing erythematous circinate rash on the trunk, thin sparse hair, anaemia, and 3 cm hepatosplenomegaly.

Pertinent laboratory findings are given in Table I. X-rays of the long bones revealed syphilitic periostitis and osteochondritis. An intravenous pyelogram was normal.

Received 14 August 1972.

*Present address: Wilhelmina Children's Hospital, Utrecht, The Netherlands.
<table>
<thead>
<tr>
<th></th>
<th>Case 1</th>
<th>Case 2</th>
<th>Case 3</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Laboratory findings</strong></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td><strong>in 3 cases of congenital syphilis with glomerulonephritis</strong></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td><strong>Case 1</strong></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td><strong>Urine microscopy</strong></td>
<td>+ + + +</td>
<td>+ + +</td>
<td>+ + +</td>
</tr>
<tr>
<td><strong>WBC</strong></td>
<td>+ + + +</td>
<td>+ + +</td>
<td>+ + +</td>
</tr>
<tr>
<td><strong>Casts</strong></td>
<td>390–1000</td>
<td>350–420</td>
<td>260–990</td>
</tr>
<tr>
<td><strong>Urine protein (mg/100 ml)</strong></td>
<td>390–1000</td>
<td>350–420</td>
<td>260–990</td>
</tr>
<tr>
<td><strong>Serum proteins (g/100 ml)</strong></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td><strong>Albumin</strong></td>
<td>1.7</td>
<td>1.3</td>
<td>1.0</td>
</tr>
<tr>
<td><strong>α1-globulin</strong></td>
<td>0.3</td>
<td>0.3</td>
<td>0.3</td>
</tr>
<tr>
<td><strong>β2-globulin</strong></td>
<td>1.4</td>
<td>0.5</td>
<td>1.2</td>
</tr>
<tr>
<td><strong>γ-globulin</strong></td>
<td>0.6</td>
<td>0.6</td>
<td>0.6</td>
</tr>
<tr>
<td><strong>IgG immunoelectrophoresis</strong></td>
<td>755</td>
<td>969*</td>
<td></td>
</tr>
<tr>
<td><strong>IgM</strong></td>
<td>650</td>
<td>256*</td>
<td></td>
</tr>
<tr>
<td><strong>ASO titre (IU)</strong></td>
<td>200</td>
<td>125</td>
<td>182</td>
</tr>
<tr>
<td><strong>Total serum haemolytic complement (normal 160–210 C′H50 units/ml)</strong></td>
<td>106</td>
<td>Very low</td>
<td>182</td>
</tr>
<tr>
<td><strong>WR, VDRL, FTA-ABS† tests</strong></td>
<td>Positive</td>
<td>Positive</td>
<td>Positive</td>
</tr>
</tbody>
</table>

*In remission after treatment.
†Fluorescent treponema antibody absorption.

Glomeruli still showed mild mesangioendothelial proliferation with capsular adhesions (Fig. 1).

Electron microscopy: the glomerular basement membrane showed irregular thickening with numerous electron-dense hump-like deposits mostly along the subepithelial aspect. The epithelial cell foot processes were fused and showed dense granularity in the neighbourhood of the deposits (Fig. 5). Villous hyperplasia of the epithelial cells, crescents, and mesangial proliferation were prominent.

The second biopsy showed advanced hyalinization. Identifiable glomerular basement membrane showed gross thickening with incorporation of the electron-dense deposits.

Immunofluorescence of the second biopsy: in non-hyalinized glomeruli finely granular immunofluorescence...
Congenital syphilis and glomerulonephritis with evidence for immune pathogenesis

### TABLE II

**Immunofluorescence studies* in renal biopsy specimens**

<table>
<thead>
<tr>
<th>Case</th>
<th>Intensity</th>
<th>Distribution</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>1</td>
<td>2</td>
</tr>
<tr>
<td>Anti-IgG</td>
<td>++</td>
<td>++</td>
</tr>
<tr>
<td>Anti-IgA</td>
<td>±</td>
<td>+</td>
</tr>
<tr>
<td>Anti-IgM</td>
<td>+</td>
<td>+</td>
</tr>
<tr>
<td>Anti-C'</td>
<td>+</td>
<td>+</td>
</tr>
<tr>
<td>Antifibrinogen</td>
<td>±</td>
<td>±</td>
</tr>
</tbody>
</table>

*Brethens et al., 1969.

Note: Intensity is graded from 0 to +++. Distribution is graded from an occasional deposit (trace) to diffusely along the glomerular basement membrane (+++).

of moderate intensity was present along the outer aspect of and within the glomerular basement membrane. This was most marked with anti-IgG, anti-IgM, and anticomplement (Table II). In the hyalinized glomeruli some immunofluorescence could occasionally be seen with anti-IgM and anticomplement sera, but not with anti-IgG. Fluorescence in these glomeruli could not clearly be located along the basement membrane, but seemed to involve the mesangium and was of a patchy character.

**Case 2.** A 2-week-old African boy was admitted with abdominal distension and bilateral inguinal herniae. Pregnancy and delivery had been uneventful. Progressive generalized oedema and ascites developed in hospital. The infant was anaemic, had a 3 cm enlargement of liver and spleen, and palpably enlarged kidneys.

The infant was given intramuscular procaine penicillin 300,000 units daily and a blood transfusion. The oedema and ascites resolved within 2 weeks, but severe proteinuria persisted. Open renal biopsy was performed 10 days after penicillin therapy. He was discharged at the age of 2 months, oedema-free, but with persisting moderate proteinuria. Clinical examination and laboratory investigations were normal when seen at 5 months.

**Biopsy findings.** Light microscopy: there was diffuse enlargement of glomeruli with mesangioendothelial proliferation and mild polymorph exudation. Capsular adhesions but no crescents were seen (Fig. 2). Methenamine silver stains showed irregular thickening with 'spiking' on the epithelial surface of the glomerular basement membrane. The interstitium showed a moderate X-rays of the long bones revealed the bone lesions of congenital syphilis. An intravenous pyelogram was normal.

**Fig. 2.—**Case 2. Glomeruli show mesangioendothelial proliferation with capsular adhesions. The interstitium is infiltrated with mononuclear inflammatory cells. (H. and E. × 133.)
infiltrate of plasma cells, lymphocytes, and polymorphonuclear leucocytes.

Electron microscopy: numerous electron-dense deposits were seen within and particularly along the epithelial aspect of the glomerular basement membrane. Mesangial cells and matrix were increased.

Immunofluorescent microscopy: all glomeruli showed finely granular immunofluorescence of marked intensity diffusely along the outer aspect of and within the basement membrane (Fig. 3). It was most marked with anti-IgG and anticomplement, less so with anti-IgA and anti-IgM, and minimal with antifibrinogen (Table II).

Case 3. A coloured boy aged 2 months was admitted with gastroenteritis. Gross generalized oedema with ascites developed in hospital. The clinical, serological, and x-ray findings resembled the features of Cases 1 and 2. Pertinent laboratory findings are given in Table I.

The infant was treated with diuretics and a blood transfusion. Open renal biopsy was performed before the administration of penicillin. Within 4 weeks of the start of penicillin therapy (6·3 megaunits) the infant had improved, with loss of oedema and minimal residual proteinuria.

Renal biopsy was repeated 3 months after the first.

Biopsy findings. First biopsy (pretreatment): under light microscopy the ultrastructure (Fig. 4) and immunofluorescence showed features similar to those found in Case 2.

Second biopsy: under light microscopy the glomeruli showed mild mesangioendothelial hypercellularity while methenamine silver stains still revealed 'spikes' on the epithelial aspect of the glomerular basement membrane.

Electron microscopy: there was obvious diminution in the number of subepithelial and intramembranous deposits, many of which now had an unusual coarse particulate appearance under very high magnification.
Congenital syphilis and glomerulonephritis with evidence for immune pathogenesis

Fig. 5.—Case 1, first biopsy. Subepithelial electron-dense deposits in glomerular basement membrane, similar to those seen in Fig. 4, but shown here in higher magnification. (×10312.)

Discussion

Hepatosplenomegaly, snuffles, sparse hair, skin lesions, anaemia, and thrombocytopenia suggested the diagnosis of congenital syphilis. This was confirmed by the positive serological tests on the infants and their mothers, and the x-ray findings of the bone lesions of early congenital syphilis.

The infants had features of severe renal disease with generalized oedema, ascites, massive proteinuria, hypoproteinaemia, and raised blood urea. One infant had macrohaematuria and another significant microhaematuria (Table I). Causes of renal disease other than syphilis were not apparent.

Penicillin therapy resulted in slow resolution of the clinical manifestations of congenital syphilis and healing of the histopathological lesions in the kidney. Herxheimer reactions were not encountered.

The clinical manifestations in our cases were similar to previous descriptions (Herrmann and Marr, 1935; Yampolsky and Mullins, 1945; Scully and Yamazaki, 1949; Mitchell, 1951; Taitz, Isaachsen, and Stein, 1961; Papaiaonnou, Asrow, and Schucknell, 1961; Pollner, 1966; McDonald, Wiggelinkhuizen, and Kaschula, 1971). Oedema is the most prominent clinical feature, though a nephritic element with haematuria and uraemia is common.

The serum protein electrophoretic pattern is characteristic of the nephrotic syndrome, though y-globulin may be normal or even raised due to markedly raised IgM levels. The latter finding in the early weeks of life denotes intrauterine infection.

Urinary protein electrophoresis in 2 of the cases revealed a nonselective proteinuria.

The published histopathological features have varied from 'minimal changes' (Yuceoglu et al., 1966), membranoproliferative glomerulonephritis (McDonald et al., 1971); membranous glomerulonephritis (Pollner, 1966), to mixed endothelial and epithelial proliferation with crescent formation (Yampolsky and Mullins, 1945; Taitz et al., 1961).

In the first infant, who had macrohaematuria and moderate uraemia with the nephrotic syndrome, biopsy showed severe mixed proliferative glomerulonephritis with crescent formation and glomerular basement membrane thickening. The duration of symptoms and thus presumed exposure to the injurious agent was prolonged in this infant, and we suggest that this had a bearing on the severity of the inflammatory reaction in the glomeruli. Repeat biopsy 6 months later revealed scarring with glomerulosclerosis of more than half of the glomeruli (Fig. 1).

The other two infants had been ill for less than 2 weeks; both showed reversible mesangioendothelial proliferation with glomerular basement membrane thickening (Fig. 2), a lesion similar to that seen in the nephropathy of acquired syphilis (Braunstein et al., 1970; Bhorade et al., 1971; Brophy et al., 1964).

The tubules showed no gross abnormalities and interstitial infiltration with inflammatory cells was not prominent.

It is evident that resolution of the microscopical abnormalities, even if reversible, is slower than the clinical improvement.

Pathogenesis. Direct injury during the spirochaetic phase, antisypilillic therapy with heavy metals and penicillin, increased susceptibility to poststreptococcal glomerulonephritis, hypersensitivity, and antigen-antibody reactions have been suggested as possible pathogenetic mechanisms.

On the basis of electron microscopy findings, Falls et al. (1965) first suggested an immune pathogenesis for the nephropathy of acquired syphilis. This was recently corroborated by the studies of Braunstein et al. (1970) and Bhorade et al. (1971).

The evidence presented suggests that the nephropathy of congenital syphilis similarly is an immune complex glomerulonephritis.

Light microscopy revealed glomerular injury with mesangioendothelial proliferation, epithelial crescents, and thickening of the glomerular basement membrane due to nonargyrophilic subepithelial deposits. On electron microscopy discrete electron-dense deposits suggestive of immune complexes...
were seen in subepithelial positions along the glomerular basement membrane. On immunofluorescent staining, immunoglobulins and complement were shown in similar sites.

Braunstein et al. (1970), in their case of acquired syphilitic nephropathy, were able to show selective deposition of $\gamma-G_1$ heavy chains and $\gamma$-light chains, but complement was not found, and neither were complement activity of the deposited $\gamma$-globulin fractions. Bhorade et al. (1971) showed complement in the deposits on immunofluorescence, though serum complement was normal. We found a reduced total serum haemolytic complement in 2 infants and complement was shown in the immune complex deposits in the glomeruli in all 3 cases. The serum complement levels returned to normal after treatment with penicillin.

Damage to, and interference with, the function of the glomerulus occurs as a result of the inflammatory reaction induced by soluble antigen-antibody complexes trapped in the glomerular basement membrane. This is mediated by the complement cascade, polymorphonuclear leucocytes, the coagulation system, and probably other mechanisms (Dixon, 1968; Dixon and Cochrane, 1970; McCluskey, 1970; Lewis and Couser, 1971).

The clinical and morphological resemblance between human immune complex-induced glomerulonephritis and experimental serum sickness glomerulonephritis in rabbits suggests similar pathogenic mechanisms. Experimentally, the earliest histopathological lesion is membranous glomerulonephritis, but as the experiment is continued, proliferative and progressive glomerulonephritis with scarring occurs (Germuth, Senterfit, and Pollack, 1967).

Congenital syphilis acquired during intrauterine life may be present for weeks or months before diagnosis. Development of immune complex glomerulonephritis may be dependent, as under experimental conditions, on the degree of antigenic exposure and antibody response. The morphological lesions in our 3 cases correlated with the apparent duration of antigenic exposure.

Autologous IgG production in the fetus is limited, and it is interesting to speculate that, in the neonate, maternal IgG transmitted across the placental barrier participates in the formation of immune complexes.

Eradication of the antigen may be expected to be followed by clinical and histological cure, provided that the glomeruli have not sustained irreversible damage. Reversibility is probably related to the type, severity, and duration of the initial insult and to the presence or absence of a mechanism for perpetuating the disease process. Prolonged exposure to the injurious agent appears to result in severe and irreversible glomerular damage.

**Treatment and prognosis.** Congenital syphilis is preventable by good antenatal care. Serological tests for syphilis on umbilical cord blood will assure an early diagnosis postnatally. The IgM fluorescent treponemal antibody absorption test (IgM-FTA-ABS) appears reliable in the differentiation of passively transferred maternal antibody from that produced by the fetus in response to syphilitic infection.

Recommended treatment is either an intramuscular injection of benzathine penicillin G 100,000 units/kg which should be repeated 1 to 2 weeks later, or procaine penicillin G 100,000 units/kg per day for 10 days (Kempe, Silver, and O’Brien, 1970).

In the past, susceptibility to intercurrent infections frequently led to a fatal outcome. Modern supportive therapy and antibiotics have markedly improved the prognosis. Early and adequate penicillin therapy is curative, but, if diagnosis and treatment are delayed, irreversible renal damage may result.

**Conclusion**

On the basis of the findings in these 3 infants we suggest that the *Treponema pallidum* in congenital syphilis be added to the list of known or suspected antigens in human immune complex-induced glomerulonephritis. This list so far includes DNA antigen in disseminated lupus erythematosus nephritis (Koffier, Schur, and Kunkel, 1967), nephritogenic types of the Lancefield group A $\beta$-haemolytic-streptococcus in acute poststreptococcal glomerulonephritis (Michael et al., 1966), *Plasmodium malariae* in quartan malaria nephrotic syndrome (Ward and Kibukamuseke, 1969), coagulase negative and positive staphylococci in glomerulonephritis associated with infected ventriculotrial shunts (Stickler et al., 1968), and penicillamine (Lachmann, 1968).

We thank Professor R. McDonald for constant encouragement and advice; Dr. H. Meyer, Acting Medical Superintendent, for permission to publish; M. v.d. Giessen from the Central Laboratory of the Red Cross Blood Transfusion Service, Amsterdam, for preparing the antisera; and Mrs. V. Eyles for preparation of the manuscript.

**References**

Congenital syphilis and glomerulonephritis with evidence for immune pathogenesis


Correspondence to Dr. Jan Wiggelinkhuizen, Red Cross War Memorial Children's Hospital, Rondebosch, Cape, South Africa.

Addendum

Since preparation of this paper two case reports have been published.


Congenital syphilis and glomerulonephritis with evidence for immune pathogenesis

J. Wiggelinkhuizen, R. O. C. Kaschula, C. J. Uys, R. H. Kuijten and J. Dale

Arch Dis Child 1973 48: 375-381
doi: 10.1136/adc.48.5.375

Updated information and services can be found at:
http://adc.bmj.com/content/48/5/375

These include:

Email alerting service
Receive free email alerts when new articles cite this article. Sign up in the box at the top right corner of the online article.

Notes

To request permissions go to:
http://group.bmj.com/group/rights-licensing/permissions

To order reprints go to:
http://journals.bmj.com/cgi/reprintform

To subscribe to BMJ go to:
http://group.bmj.com/subscribe/