GUMMA OF THE LIVER IN CONGENITAL SYPHILIS

by

D. Whitehouse and W. V. MacFarlane

From the Departments of Child Health and Venereology, Newcastle General Hospital, Newcastle upon Tyne

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Modern methods of serological testing for syphilis in pregnancy combined with highly efficient antisyphilitic therapy should now be making congenital syphilis a relatively rare disease. In addition, past reports have frequently emphasized the ease with which gummata of the liver, and, indeed, elsewhere, may be misdiagnosed. In these circumstances unusual manifestations of this condition are more than ever likely to be missed, and for this reason we feel our recent experience with gumba of the liver should be described.

Case Report

W.G., a boy of 6 years, the only son of parents both aged 40 years living in a small market town, was sent to the Out-patient Department with the suggested diagnosis of leukaemia. His history was of vague ill-health for one year, beginning with an episode called by his parents 'the blues', consisting of a period of extreme sensitivity to cold, with shivering and a blue-black appearance of the face and legs during the winter weather. This was followed by episodes of loss of appetite, pallor and pain in the left side of the abdomen lasting three or four weeks, but apparently without fever and without vomiting.

On examination he was pale, lethargic and febrile. The main physical sign consisted of a striking enlargement of the liver, especially the left lobe. The edge was well defined, reaching half-way to the umbilicus, and its surface was nodular. There was no generalized lymphadenopathy and the spleen was not palpable. The provisional diagnosis was secondary neoplasm, involving the liver.

During his first few days in hospital he showed a sustained fever averaging 100° F. and had a persistently high E.S.R. to 100 mm. in one hour. The blood picture was as follows: Hb 73%, P.C.V. 35%, M.C.H.C. 31%, W.B.C. 7,000/c.mm. (polymorphs 47%, eosinophils 2%, lymphocytes 42%, abnormally large lymphocytes 5% and monocytes 7%).

Liver function was tested by zinc sulphate, thymol turbidity, thymol flocculation and cephalin cholesterol tests, which were all normal. The alkaline phosphatase was 27.5 units (Jenner and Kay) and the serum bilirubin was 0.3 mg./100 ml.

Mantoux testing (10 international units) was negative and x-ray examination of the chest, abdomen and long bones showed no abnormality.

At laparotomy the liver was found to be studded throughout with round greyish nodules with a large mass coalescing in the left lobe to cause the enlargement which constituted the presenting physical sign. Careful search of the abdomen did not reveal any primary neoplasm, but the findings were thought to bear out the original diagnosis.

Histology. The report, by Dr. J. Hart-Mercer, was as follows:

'There is no sign of neoplasm. The nodular lesion is essentially a necrotizing chronic granuloma with almost complete obliteration of the local parenchyma. Included in the specimen are surviving areas of liver tissue but these are also being invaded by the granulomatous process which is spreading here mainly by the lymphoreticular tissue of the larger portal tracts. With a picture of necrotic foci rather resembling caseation associated with epithelioid reaction and Langhans giant cells, tubercle has to be considered.'

Serological Tests. As the Mantoux test was again negative a Wassermann reaction was performed and found to be positive. Blood Wassermann tests were repeatedly strongly positive, but examination of the cerebrospinal fluid revealed no evidence of syphilis. The parents were then investigated and the mother was found to have latent syphilis. She had a strongly positive blood Wassermann reaction but normal cerebrospinal fluid with a negative Wassermann reaction. There was no clinical evidence of syphilis. The father was entirely negative. Detailed review of the child revealed no other signs of congenital syphilis, and, in particular, re-examination of radiographs of the long bones showed no evidence of osteoperiostitic changes.

Treatment. Treatment was initially with potassium iodide, starting with 1 grain daily and increasing rapidly up to 10 grains three times daily. To this was added bismuth metal suspension ('chlorostab' brand) beginning with 50 mg. weekly and increasing gradually until the patient was receiving 150 mg. per injection, which was given deeply intramuscularly in the upper outer quadrant of alternate buttocks.
An aqueous crystalline penicillin preparation was started on the tenth day after commencing treatment, beginning with 5,000 units six hourly and increasing to 50,000 units per injection in a further 10 days. After two weeks a delayed absorption product ('prolopen', Messrs. Glaxo Laboratories) was given once daily until a total of 63 mega units of penicillin had been administered.

After nearly three months, out-patient treatment was begun, consisting of 0.2 ml. bismuth weekly for 10-week periods alternating with 900,000 units penicillin per week for seven-week periods. A total of six courses of bismuth and 39 mega units of penicillin has been given to date.

Even the short pre-penicillin course produced a dramatic decrease in the size of the liver, and by the time penicillin was begun it had receded to only half an inch below the costal margin.

Liver function tests on discharge from hospital and repeated one year later were normal and the boy has now attained a negative serology for syphilis.

Comment

Nabarro (1954), with his very extensive experience of congenital syphilis, states that hepatic gummatas are of great rarity in this variety of the disease. He says that the usual histological picture in the liver consists of a diffuse small cell infiltration occurring in utero and in the first months of life, succeeded later in some cases by fibrosis or cirrhosis varying from a microscopic lesion to complete destruction of the liver architecture. He quotes only two cases of gumma of the liver, described by Carpenter (1901) and Hutinel (1926), the former in an infant aged 8 months and the latter in a newborn infant, and says that he personally has not seen one.

McCrae and Caven (1926) reviewed 100 cases of tertiary syphilis of the liver, including five patients diagnosed as having congenital syphilis, aged 3, 13, 16, 18 and 19 years respectively. In the latter the liver was diffusely enlarged but the exact histology was unrecorded. Hahn (1943) examined 66 tertiary syphilitic livers at necropsy in adults, and pointed out that in only nine was the true diagnosis made in life. Shapiro and Weiner (1951) described a further 79 necropsies, all in adults, and made the observation that 'the visceral manifestations of late syphilis through their rarity have been neglected'.
Fiegel (1951) gave very detailed accounts of the
livers in 22 children examined at necropsy at ages
ranging from a stillborn infant to one aged 2\frac{1}{2}
years of age. All had syphilitic changes, but in very few
had the liver architecture been replaced by granulation
tissue, and in none was there a striking picture of
naked-eye gummatà, the liver being always
enlarged but usually smooth. His account of the
histology accords closely with that of Nabarro.

Jeans (1957) states that 'gummatà may occur in
almost any part of the body but they are rare in the
liver, spleen and lungs'. He refers to one case he
had seen which probably had gummatà of the liver,
but without any details.

Only one other reference to gumma of the liver,
in an infant, was found in an article by Gerlóczy
(1939), but this article could not be obtained.
Isolated gummatà of the pancreas (Raeburn, 1951),
and the stomach (Willeford, Childers and Hepner,
1952) have also been recorded.

A recent report by Macfarlane, Johns and Schofield
(1955) has emphasized these points. Their survey
covered 200 patients with congenital syphilis of whom
172 (86\%) were found to have late syphilis. Of these, 105
had ocular lesions, including interstitial keratitis,
optic atrophy choroiditis and iridocyclitis. Twenty-five
had evidence of neurosyphilis, including juvenile and
general paresis, tabes dorsalis, eighth nerve deafness,
epilepsy and mental changes. Bone changes
were found in 11 patients and joint changes in 16.

Only two patients had gummatà, a girl of 9 years
with a gumma of the larynx, and a man aged 28 years
with a gumma of the testis, which had been removed
as a malignant tumour.

These reports serve to emphasize the rarity of
gumma in the viscera, and particularly in the liver,
in childhood, and also the difficulty which may arise
in diagnosis at any age. The striking features in
this child are that such a condition can develop so
early and with no other signs of the disease, presenting,
as in many adults with tertiary syphilis, signs
imitating malignant disease.

Summary

The case is reported of a child aged 6 years
presenting with an isolated picture of gummatà of the
liver.

Allowing for the rarity of gummatous hepatitis in
children, the remarkable ability of this disease to
masquerade as other conditions serves as a reminder
that a serological test for syphilis before operation
in similar cases is worthy of consideration.

Bibliography

Carpenter, G. (1901). The Syphilis of Children in Every-day


Ch. 26, pp. 9 and 33. Hagerstown, Maryland.


Background of Congenital Syphilis. (Dept. of Venereology,
Newcastle upon Tyne General Hospital.)


10, 162.
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