Mucopolysaccharidosis type VII associated with hydrops fetalis: histopathological and ultrastructural features with genetic implications

A J Molyneux, E Blair, N Coleman, P Daish

Abstract
A case of mucopolysaccharidosis type VII (MPS VII, β glucuronidase deficiency) causing fatal hydrops fetalis in the third trimester is presented. The diagnosis was suspected on histopathological examination by the presence of foam cells in many of the viscera and foamy change in the placental Hofbauer cells. Electron microscopy showed empty cytoplasmic inclusion bodies within macrophages and in the Hofbauer cells. Enzyme assay of cultured fibroblasts showed markedly deficient β glucuronidase activity, thus confirming the diagnosis. A detailed and thorough histopathological examination of hydrops fetalis cases is important to detect subtle features of inherited metabolic disorders. Use of a structured necropsy protocol is recommended for cases of non-immune hydrops. Electron microscopy is a useful adjunct to light microscopy in cases where an inherited metabolic disorder is suspected. Precise necropsy diagnosis is important as there are implications for genetic counselling and possible prenatal diagnosis in subsequent pregnancies.

(J Clin Pathol 1997;50:252–254)

Keywords: mucopolysaccharidosis type VII; hydrops fetalis; electron microscopy.

PATHOLOGICAL FINDINGS
External examination confirmed a grossly hydropic male baby. Assessment of the facies was complicated by oedema but it was not obviously abnormal. Internal examination revealed a dilated heart and pulmonary hypoplasia (combined weight 10.6 g, expected 34±11 g; lung/body weight ratio 0.003). The liver was pale and slightly enlarged (82.4 g, expected 65±22 g) and there was splenomegaly (19.6 g, expected 4.1±2.1 g). There were blood stained pleural and peritoneal effusions and there was fresh haemorrhage into soft tissues in the neck and mesentry. Other gross features were unremarkable and there were no malformations.

The placenta was large (fixed weight 765 g) but otherwise appeared grossly normal.

Microscopic examination showed finely vacuolated interstitial foamy cells present in many organs, but most noticeably in the spleen, lung, myocardium, bowel mucosa, and bone marrow (fig 1). Similar vacuolation was seen in hepatocytes, proximal convoluted tubules, and cerebral cortex, probably in microglial cells. The placenta also showed a subtle vacuolation of villous Hofbauer cells and a normal appearance of the cytotrophoblast, features suggestive of MPS VII. Electron microscopy of lung tissue showed numerous mainly empty cytoplasmic vacuoles (fig 2). Although not specific, the ultrastructural findings also suggested the possibility of a metabolic storage disorder.

The diagnosis of MPS VII was confirmed by quantitation of enzyme activity in cultured fibroblasts. β Glucuronidase enzyme activity was negligible (0.27 nmol/h/mg protein, normal range 48–360 nmol/h/mg protein). Control enzymes (β galactosidase and u fucosidase) showed activity within their normal ranges.

Discussion
A wide variety of conditions may give rise to non-immune hydrops fetalis, and a structured approach to the perinatal necropsy is therefore
especially important. Perinatal necropsy protocols for non-immune hydrops cases emphasise the importance of a complete examination including whole body x ray, thorough gross dissection, placental examination, microscopic examination, microbiological cultures, and tissue (fibroblast) culture. The latter may be unsuccessful, particularly in cases of stillbirth with postmortem retention in excess of 24 hours. Although placental amnion is a good backup tissue for culture, we also advocate inclusion of sample collection for electron microscopy in the protocol for cases where the initial gross examination fails to show a cause for the hydrops. Electron microscopy may significantly reduce the differential diagnosis where an inherited metabolic disorder is suspected on light microscopy. In some cases (for example, GM, gangliosidosis and Gaucher’s disease) the inclusions seen on electron microscopy may be diagnostic even in the macerated fetus.

Mucopolysaccharidosis type VII is a rare autosomal recessive condition which results from a deficient function of the enzyme β-glucuronidase. This glycoprotein is required for the hydrolysis of glucuronic acid residues from the non-reducing termini of glycos-amuraglycans of lysosomes. Deficiency of glucuronidase results in the accumulation of glycosaminglycans in many organs and increased urinary excretion of glycosaminglecans.

The β-glucuronidase gene has been mapped to 7q and has been isolated and characterised by molecular genetic studies. Twelve separate disease causing mutations have been identified and this molecular heterogeneity may in part explain the variable onset from the perinatal period to early childhood. Those affected may present with hydrops fetalis or later with Hurler features.

The pathological features of MPS VII have not been studied extensively. The degree and distribution of cellular vacuolation within organs appears to be variable, but the liver and spleen have been reported as showing the changes consistently. Vacuolation of chorionic villous Hofbauer cells in the presence of a normal cytrophoblast layer is a microscopic feature which should alert the pathologist dealing with a case of hydrops to a probable diagnosis of an inherited metabolic disorder. In this context, this pattern of vacuolation within the chorionic villus is characteristic of MPS VII, but other metabolic disorders (notably GM, gangliosidosis, sialidosis, and mucolipidosis II) may show vacuolation of the cytrophoblast with less conspicuous villous Hofbauer cells.

A diagnosis of MPS VII following necropsy examination has important implications for future pregnancies. Prenatal diagnosis is now possible following chorionic villus sampling or amniocentesis by β-glucuronidase assay or direct gene analysis.

In our case the diagnosis of MPS VII was suspected on the basis of the histopathological examination. The ultrastructural appearances aided the diagnosis by excluding many other types of inherited metabolic disorder with characteristic inclusion bodies, for example GM, gangliosidosis. The electron microscopic appearances reflected the empty appearance of cytoplasmic vacuoles on light microscopy and the failure to stain with the periodic-acid-Schiff reaction.

We thank Dr A H Fensom and C P Ward of the Supra-regional Laboratory for Genetic Enzyme Defects, SE Thames Regional Genetics Centre, Guy’s Hospital, London, for carrying out the enzyme tests. Thanks also to Dr S M Hsuon, Department of Medical Genetics, Churchill Hospital, Oxford, for her helpful comments.

Hashimoto’s thyroiditis associated with urticaria and angio-oedema: disappearance of cutaneous and mucosal manifestations after thyroidectomy

Antonio Amoroso, Pier Luigi Garzia, Cynthia Pasquarelli, Giuseppe Sportelli, Antonella Afeltra

Abstract
A 60 year old woman affected by Hashimoto’s thyroiditis presented with a history of recurring episodes of urticaria and angio-oedema. Clinical and laboratory evaluation of the patient excluded allergy to external agents, hereditary angio-oedema, and occult infections. A pathogenic relation between Hashimoto’s thyroiditis and chronic urticaria/angio-oedema was suspected. However, treatment with L-thyroxine had no influence on the frequency and severity of the cutaneous and mucosal manifestations, which occurred almost daily and required repeated administration of steroids. The patient therefore underwent total thyroidectomy. Cytometric analysis of intrathyroidal lymphocyte subsets showed unusual abnormalities. Urticaria and angio-oedema completely remitted after surgery; 18 months postoperatively the patient was still asymptomatic.

Keywords: Hashimoto’s thyroiditis; urticaria-angio-oedema; intrathyroidal lymphocyte subsets.

Hashimoto’s thyroiditis is an organ specific autoimmune disease characterised by an intense thyroid infiltrate of mononuclear cells. It predominantly affects women. Clinical presentation is thyroid enlargement associated with hypothyroidism. Antithyroid antibodies are found in most cases.

Urticaria is a well demarcated skin reaction characterised by oedema involving the superficial portion of the dermis. Lesions are raised and erythematous, and usually pruritic. Angio-oedema differs from urticaria in that the oedematous process is located in the deep dermis and subcutaneous or submucosal tissues. Furthermore, lesions are more painful than pruritic. The involvement of the upper respiratory tract may result in severe and sometimes fatal complications.

Recently an association between idiopathic chronic urticaria and autoimmune thyroiditis has been reported, but its pathogenic mechanisms are still unknown.” In this paper we describe the case of a 60 year old woman affected by Hashimoto’s thyroiditis associated with cutaneous and mucosal manifestations of urticaria and angio-oedema. The patient underwent total thyroidectomy, and a cytometric analysis of intrathyroidal and peripheral blood lymphocyte subsets was performed. Surgery was followed by a complete remission of urticaria and angio-oedema; 18 months postoperatively the patient was still asymptomatic.

Case report
A 60 year old woman was admitted to our department in January 1995. Her family history was unremarkable. She had been healthy until July 1993 when thyroid enlargement appeared and raised serum titres of thyroid microsomal and thyroglobulin antibodies were documented; at the same time thyroid function tests showed FT3 and FT4 values within the normal range and high TSH levels. Because of these results, L-thyroxine treatment (0.1 mg/day) was started.

In March 1994 the patient presented with episodes characterised by urticarial lesions involving the face and trunk, accompanied by swelling of the lips and tongue. Allergy tests were performed, but total IgE (PRIST) and C1-INH values were within the normal range. Routine laboratory analysis revealed only an increase of erythrocyte sedimentation rate (ESR) of 40 mm/h. During the following months attacks of urticaria and angio-oedema recurred, with progressively increasing frequency and severity; on several occasions, airway involvement caused respiratory obstruction. Attacks were treated with steroids (betamethasone, 4 mg intravenously). Continuous
Mucopolysaccharidosis type VII associated with hydrops fetalis: histopathological and ultrastructural features with genetic implications.
A J Molyneux, E Blair, N Coleman and P Daish

J Clin Pathol 1997 50: 252-254
doi: 10.1136/jcp.50.3.252

Updated information and services can be found at:
http://jcp.bmj.com/content/50/3/252

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/