Disseminated herpes simplex virus infection with hepatitis in an adult renal transplant recipient

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SUMMARY A rare occurrence of disseminated herpes simplex virus infection with hepatitis in an adult renal transplant recipient is described.

Disseminated herpes simplex virus (HSV) infection is well recognised in neonates and produces a high mortality.1, 2 In contrast it is very rare in adults; less than a score of cases have been documented, and only nine cases appear to have been recorded in the United Kingdom in the last 32 years.3–7 This report describes disseminated HSV infection with hepatitis in a 39-year-old female renal transplant recipient.

Case report

JP, aged 39, developed nephrotic syndrome in 1962 and went on to regular home dialysis in 1968. In 1975 thyrotoxicosis was diagnosed and treated medically. In 1978 she developed severe dyspnoea on exertion and mixed aortic valve disease was found. Ventricular extra-systoles were noted and atrial fibrillation occurred intermittently during dialysis. Cadaver renal transplantation was carried out on 2 February 1980. Dialysis was required in the initial post-transplant period but the graft then functioned well. On the fourth day after transplantation, she developed ventricular fibrillation and required resuscitation and cardioversion. Her further course was complicated by persisting multifocal ventricular ectopic beats resistant to drug treatment and congestive heart failure supervened.

From the 6th day, liver enzyme activities were raised. Alanine-aminotransferase activity was initially 113 IU/l but over 1000 IU/l by the 23rd day. The serum bilirubin remained normal throughout. There was no clinical suggestion of liver failure. She collapsed suddenly on the 23rd day and could not be resuscitated.

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proximal ascending aorta and mild calcific aortic stenosis.

The liver (2010 g) had a very mottled cut surface with multiple irregular, rather ill-defined yellowish areas 1-4 mm in maximum diameter; histologically, these were foci of necrosis distributed randomly throughout the lobules, and there was only a mild associated infiltrate by polymorphs, lymphocytes and macrophages. Many hepatocytes at the periphery of the necrotic areas contained eosinophilic intranuclear inclusions (Cowdry Type A).

The spleen was enlarged (460 g) and unduly soft on section. Histologically very occasional small foci of necrosis were present, and a few cells in those areas contained intranuclear inclusions.

The adrenals appeared normal microscopically, but multiple small foci of necrosis with very little associated inflammatory cell infiltration and intranuclear inclusions were seen microscopically.

The patient's kidneys showed "end-stage" renal disease. Other organs including brain, pituitary, thyroid, salivary glands, gastrointestinal tract, internal genitalia and the transplanted kidney, were normal.

**VIROLOGICAL LABORATORY FINDINGS**

The patient was reviewed weekly for evidence of virus infection following transplantation. Blood, saliva, throat swabs, and urine were collected and liver was obtained at necropsy. Specimens for isolation were inoculated into monkey kidney, HEp-2 and human embryo fibroblast cell cultures. Specimens taken two days after transplantation did not yield virus, but urine and saliva taken at 10, 15, and 18 days yielded HSV. Virus isolated from liver was identified as HSV type 1 using methods described by Pauls and Dowdle.\(^8\)

To demonstrate HSV antigen in all organs examined histologically, two techniques were used. One involved an indirect immunofluorescence method, after digestion of the tissues with trypsin,\(^9\) using rabbit antiserum to HSV type 1 and fluorescein-conjugated sheep anti-rabbit antibody (Wellcome Laboratories); the other was an immunoperoxidase method using antisera to HSV type 1 and 2 (Dako-immunoglobulins, supplied by Mercia Brocades Ltd, Surrey). With both techniques, viral antigens were demonstrated in liver and adrenal; these were maximal around the periphery of necrotic areas (Figure). The pattern of immunoperoxidase staining was similar with antisera to both HSV types 1 and 2, illustrating the considerable antigenic relation between the two strains.

To confirm the histological and immunofluorescence results, small pieces of formalin-fixed liver tissue were processed for electron microscopy and embedded in Spurr resin.\(^10\) Thin sections of this material were viewed in the electron microscope, and although tissue preservation was poor, numerous herpes virus-like particles could be recognised in the cytoplasm and nuclei of many hepatocytes.

The diagnosis of HSV infection was supported by serological studies. During the year preceding transplantation, persistently low concentrations of complement-fixing (CF) antibody to HSV (CF titre 1/10) had been found. HSV CF antibody titres remained at 1/10 on the 5th and 10th days after transplantation but had risen to 1/160 on the 18th and 22nd days. IgM antibodies to HSV were detected in the last two sera but not in serum taken 26 days before transplantation. Low unchanging concentrations of CF antibody to cytomegalovirus (titre 1/10) were present in all sera examined both before and after transplantation.
Discussion

Disseminated HSV infection in adults is usually accompanied by stomatitis and as in infants, the outcome is usually, but not invariably, fatal. One of the few patients to survive was a 24-year-old pregnant woman who recovered from severe illness but delivered a macerated fetus. Disseminated HSV infection in both children and adults may be caused by HSV type 1 or type 2. In adults, it is often associated with pre-existing clinical conditions where cellular immunity has been compromised—for example, severe burns, thymic dysplasia, Hodgkin's disease and after steroid treatment for asthma and renal transplantation. The observation of HSV-specific IgM complement-fixing antibodies in the patients (heat-treated) sera may have been spurious, resulting from the banding of heat-induced aggregates of HSV-specific IgG antibody with the IgM immunoglobulin fraction in the sucrose density gradients. Rheumatoid factor was not detected in any of the sera. Patients with recurrent HSV disease have been found to have titres of HSV-specific IgM antibodies which remained constant before and after a recurrence. In the case described here, quantitative data on HSV-specific IgM antibody concentrations were not available for comparison.

The clinical, histological and virological investigations on the patient were compatible with a diagnosis of disseminated HSV infection although the cause of death was myocardial infarction. Serological evidence indicated a recrudescence of latent HSV type 1. Previous observations in cases of HSV hepatitis in adults were confirmed and in addition, immunofluorescence and immunoperoxidase labelling techniques demonstrated HSV antigen within areas of necrosis. In retrospect, it is clear that insufficient attention was paid to the original HSV isolate obtained from urine. Although it was appreciated that the isolation of HSV from urine was most unusual and the possibility of a viraemia was considered, the full significance of the finding was not grasped. Indeed, a week later the patient developed clinical herpes genitalis, and this merely strengthened an erroneous belief in specimen contamination. It is now clear that the original isolate was a finding of utmost importance, and had the patient been treated at this stage—that is, 14 days after transplant, with vidarabine or acyclovir she might have been spared cataclysmic dissemination of HSV.

This case further demonstrates the association between immunosuppression and disseminated HSV with hepatitis in adults. It is probable that steroid treatment is instrumental in exacerbating the infection in renal transplant recipients.

References


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