Role of hepatitis C virus in chronic liver disease occurring after orthotopic liver transplantation

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Abstract
Paediatric orthotopic liver transplant recipients may develop chronic hepatitis after surgery. To investigate the role of hepatitis C virus in this pathology a cohort of 249 paediatric orthotopic liver transplant recipients was studied. Sixteen children (6.4%) were found to have chronic hepatitis C virus hepatitis after orthotopic liver transplantation. All but one of them had serum transaminase values which were persistently raised two to eight times the upper limit of normal. Thirteen were positive for both serology and serum hepatitis C virus RNA. Serum hepatitis C virus RNA detection occurred five to 33 months before hepatitis C virus antibodies. Liver tissue hepatitis C virus RNA and hepatitis C virus core antigen were detected in five. In one patient, tissue hepatitis C virus core antigen was detected when other tests for hepatitis C were negative. Two patients had positive human cytomegalovirus serum antibodies and RNA before transplantation. Although serum hepatitis C virus RNA was not detected after transplantation, serum enzyme immunosorbent assay and tissue core antigen were still detectable in both patients. In another child, serum hepatitis C virus RNA was positive and hepatitis C virus core antigen was found on a liver biopsy specimen but antihepatitis C virus antibodies were negative as well as liver hepatitis C virus RNA. No patient developed severe liver disease or cirrhosis during a follow up of up to 72 months. It is concluded that hepatitis C virus is a significant cause of morbidity after paediatric orthotopic liver transplantation. Diagnosis cannot rely on serological testing only. The patients remained stable on follow up, but longer prospective histological studies remain necessary to establish prognosis.

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Viral infections are a major concern for paediatricians involved in the care of paediatric liver transplant recipients.1 2 Of particular concern is hepatitis C which is characterised by a high propensity to chronicity and may lead to severe damage in the liver graft. In contrast to adult recipients who are commonly infected by hepatitis C virus before transplantation,3–6 children are more likely to contract hepatitis C virus during or after the transplant procedure itself.7 The pathogenicity and natural history of hepatitis C acquired on or after transplantation in childhood may not necessarily mimic recurrence of hepatitis C virus infection as observed in adult liver transplant recipients.

In the present series, we analyse the prevalence of hepatitis C virus infection in a cohort of 249 paediatric liver transplant recipients, and report our experience of hepatitis C virus antibody and hepatitis C virus genome detection and the natural history of this infection in liver transplantation recipients in childhood.

Patients and methods
PATIENTS
Between March 1984 and September 1993, orthotopic liver transplantation was performed in 314 children at our centre. The present study includes 249 of these 314 (79.3%) children who have been regularly screened for the presence of hepatitis C virus antibodies using second and third generation antihepatitis C virus (EIAs). The screening took place at each major check up performed on the patients, that is three, six, and 12 months after transplant and yearly thereafter, starting with the availability of tests for hepatitis C virus in January 1990. A minimum of three hepatitis C virus tests were performed in each patient, and the minimal interval between the first and last investigation was six months. A total of 151 patients had transplant after January 1990 and 98 before. Blood and organ donors were also screened for antihepatitis C virus antibodies from that date. Sixty five (20.7%) children were not included in this study because they had missed one or more major follow up visits and/or hepatitis C virus serum marker determination.

Liver biopsy and biochemical data were obtained in all patients at six and 12 months after transplant and thereafter every year. In addition, liver biopsies were performed if abnormal liver function tests were present. In patients with abnormal liver histology or biochemical tests, additional serum samples were tested for antihepatitis C virus by third generation recombinant immunoblot assay (RIBA3.0) and for hepatitis C virus RNA by polymerase chain reaction (PCR) (see laboratory techniques). In seven out of 16 children, a well preserved frozen biopsy sample obtained at the time of the liver enzyme abnormalities...
was available for hepatitis C virus RNA and hepatitis C virus antigen detection, using a new immunohistochemical assay, as described elsewhere. Liver tissue hepatitis C virus RNA was also detected by the PCR technique.

**LABORATORY TECHNIQUES**

Serum samples were stored at −70°C until used for testing. Antihepatitis C virus antibodies were detected with second or third generation enzyme immunoassorbent assay (EIA) (screening and confirmatory assay (Ortho Diagnostic Systems). These tests were performed according to the manufacturer’s recommendations. In the third generation EIA (Ortho), an additional recombinant antigen NS5 has been added to the other recombinant antigens (core, C33c, C110–3) used in the second generation assay.

In the third generation RIBA test (RIBA3.0), the hepatitis C virus antigens, immobilised on nitrocellulose strips, are either recombinant proteins (C33c and NS5) or synthetic peptides (c100p and c22p). The results, expressed in accordance with the rating system (from 1+ to 4+) proposed by the manufacturer, were defined as positive when reactive for at least two antigens and indeterminate when positive for only one antigen.

Hepatitis C virus RNA was extracted from 50 µl of serum sample using the acid guanidinium thiocyanate-phenol-chloroform method. Each serum sample was coextracted with one H20 sample used as negative control. The reverse transcription and nested PCR steps were done using primer pairs corresponding to the highly conserved 5’ non-coding region of the viral genome. The specificity of the PCR products was confirmed by Southern blot hybridisation under stringent conditions with an internal oligonucleotide probe (position from −185 to −148) enzymatically labelled with digoxigenin using DIG-System (oligonucleotide 3’-end labelling and detection kits; Boehringer Mannheim). Results were recorded only if the samples and the negative and positive controls were reproducible in at least two independent experiments.

Serum aspartate and alanine aminotransferases activities were determined at 37°C according to the Deutsche Gesellschaft für Klinische Chemie method on the Hitachi 737 analyser.

Hepatitis C virus RNA was detected in liver tissue using a previously described reverse transcription PCR assay with primers derived from the highly conserved 5’ non-coding region (UTR) of the viral genome. Enhanced detection of hepatitis C virus RNA was found in some samples using other 5’ sense primers (outer sense primers: GTATCTCGAGGC-GACACTCCACCATAGT=position −333 to −305 of 5’ UTR and inner sense primers: CGACCATTGATCTTTCCTCCTGTC=position −315 to −294 of 5’ non-coding region according to Han et al and Carson et al.

Hepatitis C virus antigens in liver biopsy samples of transplanted patients was detected immunohistochemically using a specific monoclonal antibody (1F-2) directed against a synthetic core peptide, as described elsewhere.

**LIVER HISTOLOGY**

The histological diagnosis of hepatitis was based on the presence of portal and/or parenchyma mononuclear infiltration and isolated hepatocyte necrosis as evidenced by acidophilic bodies. Hepatitis was classified as chronic if the changes were present for at least six months as determined clinically or histologically. Chronic liver damage was further classified as chronic active hepatitis, chronic persistent hepatitis, and chronic lobular hepatitis. For a diagnosis of chronic active hepatitis, a portal mononuclear infiltrate with piecemeal necrosis and disruption of the limiting plate by inflammatory cells was necessary. Chronic persistent hepatitis was characterised by expansion of the portal zone by mononuclear cells, with some fibrosis. Chronic lobular hepatitis was diagnosed when predominant intralobular inflammation with individual cell necrosis was found.

**CRITERIA FOR DIAGNOSIS OF HEPATITIS C VIRUS INFECTION**

Hepatitis C virus infection was established by using at least two of the following criteria: (1) positive second or third generation EIA test with positive RIBA3.0; (2) positive serum hepatitis C virus RNA; or (3) positive liver hepatitis C virus RNA with or without positive hepatitis C virus antigen in liver tissue.

**RESULTS**

**LABORATORY DATA**

Among 249 patients, 21 (8.4%) children had at least one positive test for hepatitis C virus infection. None of them had positive serological and/or liver tissue markers for hepatitis B virus infection.

In five patients, a positive second or third generation antihepatitis C virus EIA, was not confirmed by RIBA3.0 and hepatitis C virus RNA was not found in their serum. A slight increase of serum transaminases and minimal histological signs of non-specific hepatitis were demonstrated in four out of these five patients, while one patient had normal liver function tests and normal liver histology. Liver tissue was not available for hepatitis C virus RNA and hepatitis C virus antigen detection in these patients. None of these five patients displayed hypergammaglobulinaemia, a factor which is known to give false positive results in antihepatitis C virus EIAs. Because of uncertain diagnosis, these patients were excluded from the hepatitis C virus series. In the remaining 16 children (6.4%) a diagnosis of hepatitis C virus infection was firmly established. Their pretransplant diagnoses were as follows: extrahepatic biliary atresia (8), Wilson’s disease (3), congenital hepatic fibrosis (1) Byler’s disease (1), Alagille’s syndrome (1), non-A, non-B
fulminant hepatitis (1), and cryogenic cirrhosis (1). The mean follow up period after transplant in this group was 57 months (range: 18–87 months).

Eleven patients belonged to the subgroup of 98 who underwent orthotopic liver transplantation before screening of blood products and donor organs for hepatitis C virus infection (11/98 = 11.2%). Their pretransplant hepatitis C virus status was unknown. The five remaining patients had orthotopic liver transplantation performed after the donor and recipient screening tests for hepatitis C virus became available (5/151 = 3.3%). All five donors had no hepatitis C virus antibodies. The pretransplant hepatitis C virus status of these five patients was: three patients were antihepatitis C virus negative (second and third generation. EIA and RIBA3.0 test) with undetectable serum hepatitis C virus RNA levels and two were positive for all these serological assay (EIAs, RIBA3.0, and PCR). After transplant 13 of 16 patients were found to be positive for antihepatitis C virus antibodies detected by EIA and RIBA tests and for serum hepatitis C virus RNA. In addition, in five of these 13 patients, liver biopsy specimens were tested for hepatitis C virus RNA and hepatitis C virus antigen and were found to be positive for both of them. The two patients who were known to be hepatitis C virus antibody and serum hepatitis C virus RNA positive before transplantation had no more detectable hepatitis C virus RNA in serum and liver after transplantation, but they remained antihepatitis C virus positive (second and third generation EIA) with an indeterminate RIBA3.0 test. Hepatitis C virus antigen was still found in their liver biopsy specimens (table). In the patient with fulminant hepatitis, serum hepatitis C virus RNA was positive on two subsequent determinations but thereafter it became undetectable. Hepatitis C virus antibodies were always negative using EIAs and RIBA3.0 test and no hepatitis C virus RNA was found in liver tissue. However, hepatitis C virus antigen was positive in liver tissue (table).

In five children, serum hepatitis C virus RNA preceded the appearance of hepatitis C virus antibodies, the mean delay of seroconversion was about 20 months (range 5–33 months). All but one hepatitis C virus infected child had persistently raised transaminases fluctuating from twice to eight times the upper normal limit.

**Discussion**

The 6.4% prevalence of hepatitis C virus infection reported in our paediatric liver transplant series is similar to that reported from the Cochin Bicétre paediatric series, while prevalence as high as 35% has been reported after orthotopic liver transplantation in previously hepatitis C virus negative adult recipients.

All our patients showed to variable degrees histological features compatible with the diagnosis of chronic hepatitis C. The histological aspect of chronic hepatitis C was not specific in our experience and this is generally explained by immunosuppression itself, which may alter the histological appearance of viral hepatitis after transplantation. In addition associated problems may interfere, such as rejection, cytomegalovirus related hepatitis, ischaemic injury, drug toxicity, and vascular and biliary problems. Although none of the patients had fibrosis nor cirrhosis, a longer follow up is needed to establish long term prognosis of this slowly progressive disease, whose natural history in paediatric patients who have not been transplanted also remains poorly documented. It has been shown that only 10% of children infected with hepatitis C virus but not transplanted achieve sustained biochemical remission and that severe active hepatitis or even cirrhosis may develop within six years of infection. No case of acute graft failure or rapidly progressive hepatitis was observed in our series, in agreement with other paediatric and adult series. Development of cirrhosis is described in adult patients, and 50% of adult patients may have chronic active hepatitis two years after the initial acute hepatitis C virus hepatitis. Newly acquired hepatitis C seems to have a slower progress after transplantation than recurrent hepatitis C in adult patients. Even if some children may have been infected by hepatitis C virus before transplantation, hepatitis C was usually not the original cause of liver disease, and patients may be infected at the time of transplantation or shortly after. Pretransplant hepatitis C virus infection was diagnosed in two of our patients, one of them having cryogenic cirrhosis compatible with hepatitis C virus infection as the primary cause of liver disease.

Systematic screening of blood donors for hepatitis C virus has reduced but not eliminated the risk of hepatitis C virus infection acquired after transfusion. For solid organ
transplantation, however, hepatitis C virus may also be transmitted by the graft. This mode of transmission may have contributed to de novo hepatitis C virus infections in our patients, although we cannot exclude reinfection of their liver by pre-existing hepatitis C virus infection. They may also become infected by blood transfusions administered during and after orthotopic liver transplantation. This stresses the importance of testing organ donors for the presence of hepatitis C virus RNA both in serum and graft, a time consuming test which is unfortunately not available routinely in laboratories.

A complete concordance of tests for hepatitis C virus (EIA, RIBA, and serum hepatitis C virus RNA) was observed in 13 out of 16 (81%) transplant children. Two patients were already positive for hepatitis C virus markers before transplant; antihepatitis C virus antibodies remained positive after orthotopic liver transplantation, but serum and liver tissue hepatitis C virus RNA were undetectable. Positive hepatitis C virus antigens in these grafts suggest, however, that infection recurred after transplant unless the donors were themselves infected with hepatitis C virus. One of these two patients had normal liver function tests and normal histology during follow up (18 months), and hepatitis C virus antigen was the only evidence of hepatitis C virus infection after transplant.

The high sensitivity of the hepatitis C virus antigen detection assay was further demonstrated in the patient with fulminant hepatitis, who had serum hepatitis C virus RNA detected on two occasions with negative serology. Positive liver hepatitis C virus antigen remained the only positive test in this patient whose liver histology after transplant and biochemical profile was consistent with chronic hepatitis C.

An important finding in our series is the frequent negative hepatitis C virus serology in our patient despite proved hepatitis C virus infection (positive serum or tissue hepatitis C virus RNA), with seroconversion delay, as long as 33 months after infection. The possibility of false negative results in antihepatitis C virus EIA and RIBA tests has been demonstrated in transplant patients with positive serum hepatitis C virus RNA suggesting that immunosuppressive treatment might delay the appearance of serum antibodies against hepatitis C virus.

This and the intermittent viremia of hepatitis C virus may hamper the positive diagnosis of hepatitis C virus infection in many patients. Non-detection of serum hepatitis C virus RNA can be due to low concentration of circulating virions or intermittent viremia.

In five of 20 patients, hepatitis C virus antibodies were found using a second or third generation EIA, but no hepatitis C virus RNA was found in serum and RIBA3.0 was negative. These false positive results in antihepatitis C virus assays have been previously reported and may lead to overestimation of the prevalence of hepatitis C virus infection. RIBA negativity does exclude hepatitis C virus infection.

Conclusion

Systematic screening of patients and donors for hepatitis C virus is mandatory, before and after transplantation. Negative antihepatitis C virus assays do not exclude hepatitis C virus infection, and positive tests must be confirmed by RIBA test and detection of serum hepatitis C virus RNA. Whenever possible, hepatitis C virus RNA must be looked for in liver tissue. The detection of hepatitis C virus antigen seems very promising as a tool for diagnosis of hepatitis C virus infection in children who have had orthotopic liver transplantation. Donor serum and liver hepatitis C virus RNA testing, and perhaps donor liver hepatitis C virus antigen testing, may decrease the prevalence of de novo hepatitis C virus infection in transplanted children. Short term prognosis of the transplant child is not affected by hepatitis C virus infection. We did not observe rapid progressive hepatitis, nor severe liver damage on follow up in this series, but long term prospective histological studies are needed to determine the natural course of hepatitis C virus infection in these patients.

The psychology of smallness

One reason given for offering growth hormone treatment to short children who are not demonstrably growth hormone deficient is that it might save them psychological trauma. Workers in Buffalo, New York State (David E Sandberg and colleagues, *Pediatrics* 1994; 94: 832–40) have measured psychosocial functioning in their short patients. The study involved 258 children and adolescents (180 boys, 78 girls) aged 4 to 18 years referred to a paediatric endocrinology clinic because of short stature (height less than 5th centile). Their height Z (SD) scores were: mean -2.3, SD 0.5, range -4.0 to -1.6. Fifty four per cent had 'normal variant' short stature, 33% had various diagnoses including growth hormone deficiency (14%), undernutrition, skeletal dysplasia and chromosomal abnormality, and the remaining 13% had short stature of undetermined cause. Psychosocial assessments were made using three questionnaires; the child behaviour checklist (CBCL) completed by parents, and the youth self report and self perception profile, both completed by the patients. Boys in the study were described by their parents as being less socially competent and showing more behavioural and emotional problems than the CBCL norms and the boys described themselves as being less socially active but had normal estimates of their own abilities and self worth. Nevertheless the psychosocial morbidity in boys was not great and the findings in girls were normal. Neither the degree of short stature nor its cause was significantly related to psychosocial functioning. The authors conclude that psychosocial malfunction is not a reason for growth hormone treatment of most children with normal variant short stature.
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