Ultrafiltration for intractable ascites after liver transplantation

G Noble-Jamieson, N Jamieson, N D Barnes

Abstract
A 5-7 year old boy with α1-antitrypsin deficiency developed intractable ascites after liver transplantation. Conservative treatment was unsuccessful and after 18 days concentrated ascitic fluid was infused intravenously using a Gambro haemofilter. The ascitic loss resolved rapidly. This new method of ultrafiltration proved simple and effective.

Case report
Severe intractable ascites was the main indication for liver transplantation in a 5-7 year old boy with α1-antitrypsin deficiency. After the operation the ascites returned and massive amounts were lost through the abdominal drains from the fourth postoperative day. The next day the serum bilirubin concentration and alanine transaminase activity rose and ultrasound Doppler demonstrated poor hepatic compliance. These are all indications of possible acute rejection.

A liver biopsy specimen confirmed moderately severe rejection and treatment with high dose intravenous methylprednisolone was given for three days. There was a rapid response with the liver function tests returning to normal within seven days. The ascitic fluid losses, however, increased to a maximum of 10 l/day. The fluid lost was a sterile exudate which contained 39 g/l of protein. Repeated examinations with ultrasound Doppler excluded obstruction of the inferior vena cava and the hepatic and portal venous systems. Despite massive replacement with blood products the protein loss caused a severe coagulopathy (fig 1). From the 11th day all ascitic fluid loss was reinfused intravenously. This rapidly corrected the coagulopathy but the volume of ascites remained the same (fig 1).

After 18 days it was decided to concentrate the ascitic fluid using a Gambro Haemofilter FH22 (Gambro Dialysatoren) before reinfusion. In a closed circuit the ascitic fluid was drained through a peritoneal catheter, then pumped across the haemofilter membrane. The protein concentrate was then reinfused through a blood transfusion filter (Pall) into a femoral vein (fig 2). The volumetric infusion pump (IMED) was set at a rate between 100 and 600 ml/hour, starting with a high rate initially while the abdomen was grossly distended and decreasing as the abdominal girth diminished. The filtrate, which varied in volume from 100 to 150 ml/hour, was discarded. The haemofilter was perfused with heparin at a dose of 100 units/hour to prevent blockage of the filter and clot formation in the ascitic fluid. No prophylactic antibiotic treatment was used and ascitic fluid sent daily for culture remained sterile. Transient hypoponatraemia and oliguria occurred on the third day of the procedure but the serum urea and creatinine concentrations remained stable. After six days the abdominal girth had decreased from 80 cm to 65 cm and ultrafiltration was stopped. Subsequently there was only slight ascitic drainage, and this ceased entirely after two days.

Figure 1 Blood product requirements and changes in prothrombin time in relation to ascitic fluid losses.

Figure 2 Circuit for ultrafiltration and reinfusion of ascites.
Ultrafiltration for intractable ascites after liver transplantation

Discussion
In liver cirrhosis ascites formation is a consequence of progressive intrahepatic venous obstruction, leading to a rise in intrasinusoidal pressure and increased hepatic lymph production. Another important mechanism is the hyperaldosteronism and urinary sodium retention which occurs in response to the depletion of the vascular space. Even when there has been gross ascites preoperatively it generally resolves promptly after successful liver transplantation. Among 120 recipients of paediatric liver transplant, only three children have developed severe postoperative ascites, on each occasion in association with acute graft rejection. Ultrasound Doppler studies have proved extremely useful in the early diagnosis of acute rejection by demonstrating a reduction in hepatic venous pulsatility, indicating a swollen and non-compliant liver. These changes reflect reduced hepatic venous outflow and ascites formation may follow. Depletion of the intravascular space leads to renin, angiotensin, and aldosterone secretion and so to urinary sodium retention. This secondary hyperaldosteronism probably contributes to prolonged ascitic loss even after a rejection episode had resolved. It seems probable also that the severe preoperative ascites may have opened channels for ascitic fluid production.

Thus the rapid development of ascites after liver transplantation may indicate the onset of acute graft rejection. When venous obstruction has been excluded a liver biopsy is needed. Management should include control of the rejection and treatment of the ascites. Replacement of large volumes of plasma proteins may be required to replenish the intravascular volume, to maintain normal coagulation, and to maintain humoral immunity. Diuretics, especially aldosterone antagonists such as spironolactone, are usually given to counteract the secondary hyperaldosteronism. When these forms of treatment prove inadequate it seems logical to replace the ascitic fluid loss, and various methods of reinfusion and ultrafiltration have been described. The ultrafiltration device most commonly described has been the Rodia scint system, which allows continuous ultrafiltration and reinfusion of the concentrated ascites. We used a less complicated and more readily available method for ultrafiltration of ascites. Continuous arteriovenous haemofiltration is a well established technique and is frequently used for patients with oliguria and fluid overload. The haemofilter removes water and crystalloids with a molecular weight below 15 000. Filtration is usually driven by arterial pressure alone but can be enhanced by a pump and this was of course needed to circulate the ascitic fluid. By recycling the ascites in this way protein losses were avoided and the clotting deficit was rapidly corrected. Immunoglobulins and complement are also lost in ascitic fluid and their retention reduces the risk of infection. To our knowledge this method of ultrafiltration of ascites has not been used before and it proved remarkably simple, practical, and effective.

We thank the nursing staff of the paediatric intensive care unit for their technical advice and expertise in caring for this patient.

Ultrafiltration for intractable ascites after liver transplantation.

G Noble-Jamieson, N Jamieson and N D Barnes

Arch Dis Child 1991 66: 988-989
doi: 10.1136/adc.66.8.988

Updated information and services can be found at:
http://adc.bmj.com/content/66/8/988

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/