Transplantation of haemochromatosis liver and intestine into a normal recipient

Background—Haemochromatosis is a common genetic disease leading to iron overload. Although the gene had been identified (HFE), the pathogenesis had not been fully elucidated. The inadvertent transplant of a C282Y homozygous liver and intestine provided a unique opportunity to study this problem.

Methods—A 19 year old man underwent orthotopic liver and intestinal transplantation in January 1997 for the treatment of short bowel syndrome secondary to a mid gut volvulus with resection and cholestatic liver disease resulting from total parenteral nutrition. The organ donor was an 18 year old woman posthumously discovered to be a C282Y homozygote for haemochromatosis.

Results—Preoperative recipient blood tests included a transferrin saturation of 10% (normal 20–55%). Transferrin saturation at 94% with a normal serum ferritin of 34 µg/l (normal range 15–300 µg/l) and a ferritin level of 56 µg/l. Hepatic iron concentration at four months was 20 µmol/g and at 22 months was 22.3 µmol/g (normal 0–35 µmol/g). Genetic testing for haemochromatosis on a peripheral blood sample at 21 months was normal (wild type) for the C282Y mutation of the HFE gene. Genetic testing on a peripheral blood sample at 21 months was normal (wild type) for the C282Y mutation. The donor family was investigated for haemochromatosis (fig 1). A brother of the donor was an iron loaded homozygote and her mother appeared to be a non-expressing homozygote. The mother had a non-identical twin that was homozygous for the C282Y mutation with abnormal iron studies.

Conclusions—This case suggests that the genetic defect of haemochromatosis has been transplanted into the recipient with the donor intestine and that iron accumulation will probably occur with time. Within 21 months of transplantation the recipient is showing evidence of the typical biochemical abnormality seen in a young patient with haemochromatosis, namely an increase in transferrin saturation with a normal hepatic iron concentration. Although the serum ferritin is normal, it is likely that if untreated it will continue to rise with time. Therefore, we have identified a predisposition to future iron overload rather than iron overload at 21 months. The concomitant transplantation of the haemochromatosis liver is less likely to be contributing to the abnormal iron metabolism. Transplantation of a haemochromatosis liver alone into a normal recipient has been previously documented at this centre with a progressive decline in hepatic iron concentration and a normal radioulex absorption study. This supports the hypothesis that the fundamental defect in haemochromatosis is site specific at the level of the intestine rather than a systemic abnormality.

P C ADAMS
G JEFFREY
Department of Medicine,
University of Western Ontario,
London, Ontario, Canada

R ALANEN
S CHAKRABARTI
Department of Pathology,
University of Western Ontario,
London, Ontario, Canada

R PRESHAW
Department of Surgery,
University of Calgary,
Calgary, Alberta, Canada

W HOWSON
Department of Laboratory Medicine,
University of Western Ontario,
London, Ontario, Canada

D GRANT
Department of Surgery,
University of Western Ontario,
London, Ontario, Canada

Correspondence to: Paul C Adams, MD, Department of Medicine, London Health Sciences Centre, 339 Windermere Road, London, Ontario, Canada N6A 5A5. Email: padams@julian.uwo.ca

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