Case report

Wilson’s disease and hepatocellular carcinoma: possible protective role of copper

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SUMMARY A male patient with Wilson’s disease developed a hepatocellular carcinoma after treatment for nine years with D-penicillamine. Examination at necropsy showed that excess liver copper had been effectively removed. As copper has been shown to protect against chemically induced hepatocellular carcinoma in rats, this may be the reason for the extreme rarity of hepatocellular carcinoma in patients with Wilson’s disease and possibly in other liver diseases with hepatic copper overload.

Hepatocellular carcinoma is a common complication of longstanding cirrhosis and, overall, the likelihood of this in any particular aetiological group of patients has been said to depend on the proportion of men within that group and the duration of cirrhosis. At one end of the spectrum are patients with haemochromatosis or HBsAg positive cirrhosis who are predominantly male and at the other end are those with primary biliary cirrhosis or HBsAg negative chronic active hepatitis with very few men affected. It is therefore suprising that in Wilson’s disease (hepatolenticular degeneration) which has an even sex incidence and a substantial proportion of patients surviving for long periods, there have only been two proven cases and one possible further instance. In this paper we describe a fourth case and discuss the relationship between the development of hepatocellular carcinoma and the protective role of copper overload which has been shown experimentally.

Case history

A 31 year old male chemical engineer was referred to the Liver Unit in December 1971 after an episode of melaena. There had been previous episodes in 1968 and 1969. He had recently noticed mental fuzziness and difficulty in concentrating at work and on examination Kayser-Fleischer rings and hepatosplenomegaly were present. Investigations showed a serum bilirubin concentration of 10.3 μmol/l (0.6 mg/dl); aspartate aminotransferase 34 IU (normal <50 IU); alkaline phosphatase 55 IU (normal <100 IU); albumin 30 g/l; total protein 53 g/l; and prothrombin time 17 seconds (control 12 seconds). Immunoglobulins, autoantibodies, HBsAg, and alpha fetoprotein were normal or negative.

Hepatic arteriogram and splenic venogram showed gastric and oesophageal varices with patent splenic and portal veins. Electroencephalogram was normal.

The diagnosis of Wilson’s disease was confirmed by the finding of a zero serum caeruloplasmin concentration on two occasions, a low serum copper concentration of 0.8 μmol/l (5 μg/dl) and 1.3 μmol/l (8.1 μg/dl). Urinary copper was 0.94 μmol (58.9 μg) and 1.15 μmol (72 μg) per 24 hours (normal <0.8 μmol (50 μg)/24 h) before and 6.65 μmol (410 μg) per 24 hours after D-penicillamine. Investigations of the family revealed a low caeruloplasmin and raised urinary copper excretion in his sister (who was subsequently treated with D-penicillamine) with normal concentrations in both parents.

In January 1972 a lienorenal anastomosis was performed and he was started on D-penicillamine therapy. This was continued for the next nine and a half years at dosages ranging from 725 to 900 mg daily (Fig. 1). In 1973 he had a minor episode of encephalopathy and in 1980 he developed fluid retention which was thereafter controlled by spironolactone. He was otherwise well and in full time

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copper-associated protein in large groups of cells patchily and randomly distributed. The wedge biopsy specimen showed the same features and confirmed the presence of micronodular cirrhosis. Follow up biopsy showed a more collapsed liver with broader bands of inflamed connective tissue surrounding small hyperplastic nodules. Perls’ stain revealed grade I siderosis with an irregular distribution. Orcein staining was negative.

At necropsy the lienorenal shunt was patent and the inferior vena cava was normal. There were dilated tortuous veins in the lower oesophagus but no blood within the gastrointestinal tract. The surface of the liver showed micronodular cirrhosis. It weighed 1502 g and on sectioning there were large haemorrhagic nodules up to 5 cm in diameter (Fig. 3). There was no evidence of tumour in any other organ. Histological examination showed nodules composed of a well-differentiated hepatocellular carcinoma of trabecular pattern with evidence of bile formation (Fig. 4). On orcein staining of the surrounding liver in contrast to the initial biopsy but similar to the one taken one year before death, very few cells or groups of cells contained copper-binding protein, usually those cells enmeshed in fibrous tissue. Liver copper level was 48 mg/100 g dry weight (normal <50 mg/100 g) as determined by flame photometry at the National Hospital for Nervous Diseases, Queen Square, London.

Discussion

One possible explanation for the low incidence of hepatocellular carcinoma in patients with Wilson’s disease is that the presence of copper in the hepatocytes protects them from the oncogenic consequences of cirrhosis. There have been several reports showing protection by increased dietary copper from chemically included hepatic carcinogenesis in rats.5-8 If this mechanism operates in Wilson’s disease protection could be reversed once the excess copper has been removed by treatment but the time needed to remove excess copper by chelation therapy is not accurately known. Sherlock9 stated that complete decoppering is unusual even after many years of treatment and according to Walshe10 the speed of decoppering depends on the dose of penicillamine, many patients managing on 900 mg/day but some requiring 2–3 g/day to remain in negative copper balance. Another factor is dietary intake which is usually from 2–5 mg/day but with water supplied in copper pipes, shellfish, nuts, chocolate, and some other foods this figure may be much higher. The only certain indication that ‘decoppering’ is adequate is a repeat chemical estimation of copper in liver tissue.
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Fig. 2  H&E stained section of percutaneous liver biopsy in 1972 showing prominent fibrous septum, cellular and nuclear pleomorphism, ballooned cells with frequent nuclear vacuolation, and peripheral cytoplasmic microvesicular transformation. Mild patchy inflammation is seen disrupting the limiting plate (H&E x300).

Fig. 3  Cut surface of the liver at necropsy showing multiple haemorrhagic tumour nodules.
Fig. 4 Sections of the tumour showing well differentiated hepatocellular carcinoma of trabecular pattern (H&E x130).

Thus our present patient had no histochemical evidence of copper overload 21 months before death (although he was still excreting over 5 μmol/24 h in his urine (Fig. 1)) and a tissue copper estimation performed after death showed the concentration to be at the upper end of the normal range. This suggests that a normal tissue copper concentration was only achieved towards the end of the nine year treatment period.

Of the three previous reports of hepatocellular carcinoma in Wilson’s disease, the first, from Norway in 1957, is the least certain. A 14 year old boy with a distant cousin who had had Wilson’s disease and a brother with possible Wilson’s disease had died 18 months after first presentation after the development of jaundice, ascites, and several haematemeses. He had no Kayser-Fleischer rings nor neurological abnormality; furthermore, serum copper-oxidase activity was raised and he had a serum copper concentration of 30 μmol/l (199 μg/100 ml). The second was a Frenchman with a 41 year history of involuntary movements and at least 22 years of hepatomegaly and Kayser-Fleischer rings. He was treated with D-penicillamine for five years before dying in hepatic coma. Necropsy showed cirrhosis and nodules of pseudoglandular type hepatocellular carcinoma. The third case was a 32 year old Japanese man with a six year history of tremor and Kayser-Fleischer rings who had been treated with D-penicillamine therapy for 18 months. He was admitted with a two-months history of right upper quadrant pain and he died 12 days later after a massive intraperitoneal haemorrhage. Necropsy showed typical pathological changes of Wilson’s disease and diffuse hepatocellular carcinoma of a trabecular pattern.

Some increase in liver copper may occur in the late stages of cirrhosis from any aetiology and the level at which intrahepatic copper deposits are protective in vivo is unknown. Considerable excess hepatic copper is also present from an early stage in certain chronic cholestatic conditions, especially
primary biliary cirrhosis in which hepatocellular carcinoma has been reported only rarely. As it is predominantly a disease of women this also has to be taken into account. The latter does not apply, however, to primary sclerosing cholangitis in which we have been unable to find any reports of associated hepatocellular carcinoma and such considerations may have added relevance now that penicillamine is being used in the treatment of these patients as well as in primary biliary cirrhosis.

References

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