Hepatic sinusoidal ectasia in association with Hodgkin's disease

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Summary
A patient developed abnormal liver function tests in association with hepatic sinusoidal ectasia concurrently with relapse of Hodgkin's disease. Tests returned to normal following successful chemotherapy for the Hodgkin's disease. Sinusoidal ectasia and the related condition, peliosis hepatis, should be recognized as rare non-infiltrative manifestations of Hodgkin's disease.

Introduction
Hodgkin's disease may be associated with a variety of pathological lesions in the liver other than lymphomatous infiltration; lesions include granulomas, cholestasis which may rarely progress to cirrhosis, amyloidosis, and portal tract accumulations of histiocytes, eosinophils, lymphocytes and granulocytes. The patient to be reported, and several previous reports, suggest that sinusoidal ectasia and peliosis hepatis should be recognized as non-infiltrative manifestations of Hodgkin's disease.

Case report
A 30-year-old man presented in January 1976 with three enlarged lymph nodes in the left anterior triangle of the neck. Biopsy showed mixed cellularity Hodgkin's disease. Detailed investigations including a staging laparotomy with splenectomy gave no evidence of disease elsewhere and he was therefore staged as IA. Liver function tests (bilirubin, alkaline phosphatase, and aspartate aminotransferase) were normal. Wedge and needle-liver biopsies at laparotomy showed only a slight increase in chronic inflammatory cells in the portal tracts. Radical radiotherapy to a mantle field was given (3500 rad in 20 fractions over 4 weeks with boosting to 4000 rad over the involved nodes).

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Follow-up was uneventful until May 1977, when routine liver function test showed the alkaline phosphatase to be 244 u/l (normal 20–85). The patient felt well and had no abnormal physical findings. In June, the alkaline phosphatase was 207 u/l, 5-nucleotidase (5-NT) 18 u/l (normal 2–11) and γ-glutamyl transpeptidase (γ-GT) 173 u/l (normal 5–65). Bilirubin and aspartate aminotransferase remained normal. In September the patient continued to feel well with no abnormal physical findings and a normal chest X-ray, but admitted to a 12 kg weight-loss in the preceding twelve months. Further investigations showed haemoglobin 9.5 g/dl, erythrocyte sedimentation rate 84 mm/hr, platelet count 1098 x 10⁹/l; alkaline phosphatase 291 u/l; 5-NT 398 u/l; γ-GT 260 u/l and albumin 37 g/dl (normal 35–45); α₁- and γ-globulins were elevated and the iron binding capacity was reduced.

It was considered likely that the Hodgkin's disease had recurred. Lymphangiography showed a diffuse enlargement of many para-aortic and external iliac nodes with a bubbly texture and large filling defects. A percutaneous liver biopsy showed a slight increase in portal tract lymphocytes; there were also some clusters of blood spaces, the significance of which was not appreciated. It was considered important for choice of therapy to discover if the patient had liver infiltration, or alternatively nodal disease only, with reactive changes in the liver. The patient was therefore submitted to a second staging laparotomy in October 1977. Nodal biopsies confirmed recurrent mixed cellularity Hodgkin's disease. At surgery, the liver was noted to be enlarged and vascular with a dark mottled appearance. A wedge biopsy and two needle biopsies were taken.

The wedge biopsy showed marked sinusoidal ectasia with the lining endothelial cells being benign in appearance. Some dilated blood spaces lacked an endothelial lining (Fig. 1). The changes
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FIG. 1. Wedge biopsy of liver showing hepatic sinusoidal ectasia (HE, ×160).

were most marked in the subcapsular zone but were also present deeper in the liver. The dilated sinu-
soids had no constant relationship with central
veins and portal tracts. Central veins were mildly
engorged with no evidence of perivenous fibrosis.
Portal tracts were normal except for a few lympho-
cytes and eosinophils in some tracts. Liver cells were
swollen in some areas and granular brown pigment
was seen in some Kupffer cells. There were no
granulomas, nor any evidence of Hodgkin's disease.
The two needle biopsies had the same appearance
although blood spaces were less prominent.

Postoperatively the patient was given radical
radiotherapy (3500 rad) to an inverted Y field,
followed by six courses of combination chemo-
therapy (MOPP regime). By February 1978, liver
function tests had returned to normal and sub-
sequently have remained so. The patient has been
followed-up until June 1981 and has had no clinical,
radiological or laboratory evidence of further recur-
rence. At no time did he take any androgenic
or anabolic steroids.

Discussion

A young man with Hodgkin's disease developed
abnormal liver function tests associated with
sinusoidal ectasia concurrently with the relapse of
his disease. There had been no evidence of dilatation
of sinusoids at staging laparotomy 15 months
earlier. Following successful retreatment of his
Hodgkin's disease the liver function tests returned to
normal.

Sinusoidal dilatation has been previously noted
in patients with Hodgkin's disease (Bruguera et al.,
1978) being found in 6 of 16 hepatic biopsies, only
one of which showed lymphomatous infiltration. The
patient now reported demonstrates that sinusoi-
dal ectasia may be associated with marked
abnormality of liver function tests. The return of
such tests to normal with successful treatment of
the Hodgkin's disease suggests that the sinusoidal
ectasia may have regressed with remission of the
primary disease.

Sinusoidal ectasia may be the initial stage, or
may represent a minor degree, of peliosis hepatis.
Sinusoidal ectasia may be seen in livers showing the
phlebitcetatic form of peliosis hepatis and both
lesions are seen following the administration of
anabolic and androgenic steroids, and in association
with malignant disease and tuberculosis. The
macroscopic appearance of the liver of the present
patient suggests that he may have had early phlebi-
ectatic peliosis hepatis in addition to marked
sinusoidal dilatation but no definite blood cysts were
seen in the biopsy.

There have been seven patients reported with
peliosis hepatis in association with Hodgkin's disease. Three of the patients were taking anabolic steroids (Bagheri and Boyer, 1974; McDonald and Speicher, 1978; Taxy, 1978) but four were not (Taxy, 1978; Duffaud-Thierman et al., 1977; Weickel, 1957). Sinusoidal ectasia and peliosis hepatis are rare, non-infiltrative complications of Hodgkin's disease. Peliosis hepatis may be responsible for the initial presentation of Hodgkin's disease (Duffaud-Thierman et al., 1977) and similarly, as in this patient, sinusoidal ectasia may produce the first evidence of relapse.

References
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