

[PICTURES IN CLINICAL MEDICINE]

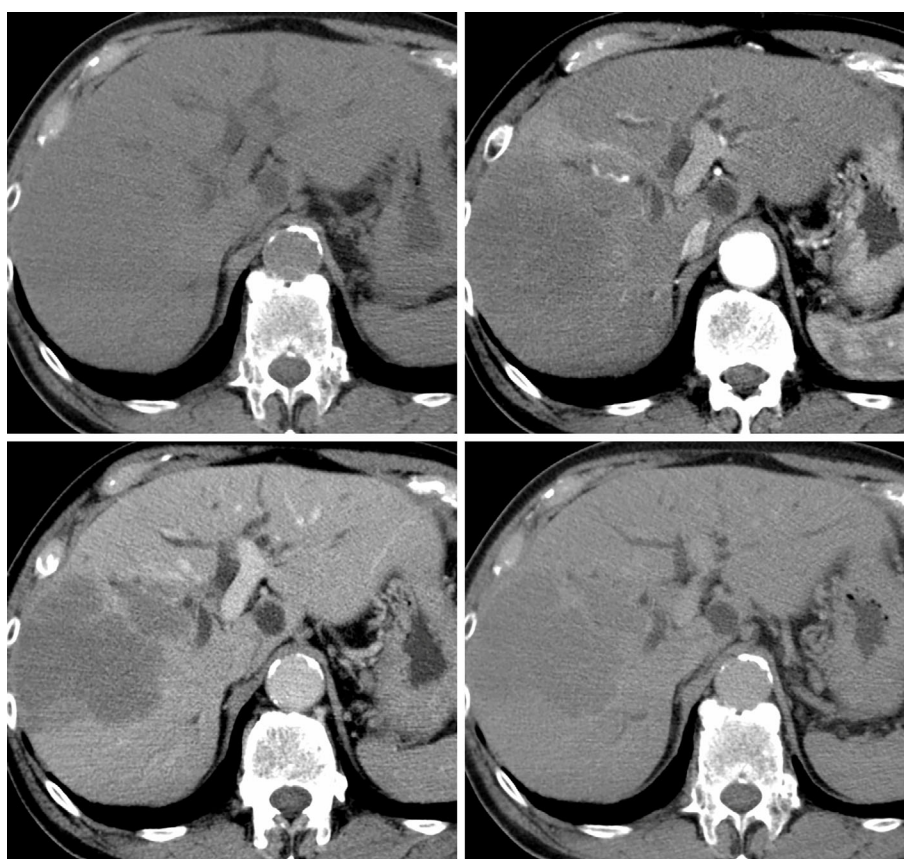
Primary Hepatic Diffuse Large B-cell Lymphoma Mimicking Intrahepatic Cholangiocarcinoma

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Key words: primary hepatic lymphoma, cholangiocellular carcinoma, obstructive jaundice

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Picture 1.

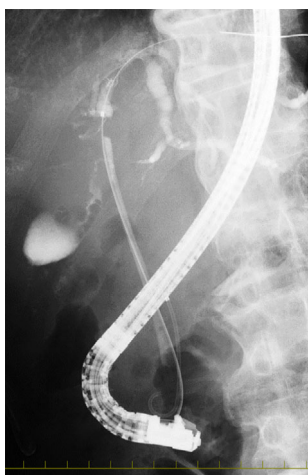
A 78-year-old man with obstructive jaundice and osteolytic left scapular tumor was referred to our hospital. Serum carcinoembryonic antigen (CEA), CA19-9, and soluble interleukin 2 receptor were 3.6 IU/L (reference range: 0-5), 128 IU/L (reference range: 0-37), and 2,940 IU/L (reference range: 145-519), respectively. Dynamic contrast-enhanced computed tomography showed a low-density mass in the

right anterior segment of the liver and dilation of the left intrahepatic bile duct (Picture 1). We performed a transpapillary biliary biopsy and drainage (Picture 2). Histologically, the tumor was composed of cleaved lymphocytes. A liver needle biopsy confirmed diffuse large B-cell lymphoma (DLBCL) that was positive for CD20 (Picture 3), BCL6, MUM1, and MYC/BCL2 and negative for CD5, CD10, and

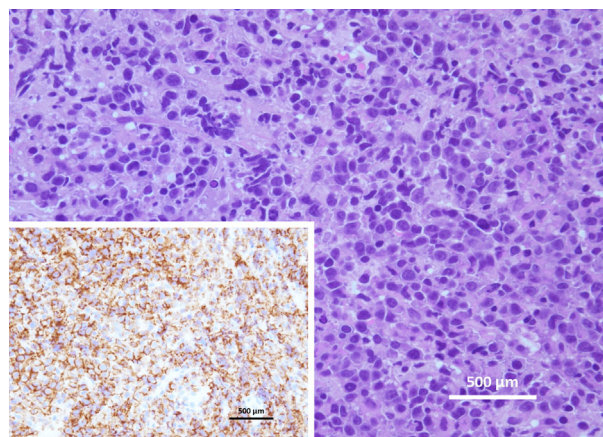
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Picture 2.



Picture 3.

TdT. The Ki-67 index was approximately 90%. ^{18}F -Fluorodeoxyglucose-positron emission tomography showed no abnormal uptake except for in the liver [maximum standardized uptake value (SUV max) 28] and left scapular tumor (SUV max 3.3) after palliative radiotherapy for bone metastasis. The findings of a bone marrow biopsy were normal. We diagnosed the tumor as primary hepatic DLBCL (NOS, ABC, MYC/BCL2 double expressor). The patient was treated with combination chemotherapy, including pirarubicin, cyclophosphamide, vincristine, and prednisolone following rituximab. Chemotherapy resulted in a partial response via a reduction in the overall tumor size by more than 50% (89.8×58.7 mm to 42×28 mm). The patient was then transferred to another hospital in order to receive further treatments. Primary hepatic lymphoma (PHL) complicated with obstructive jaundice is rare and can mimic a diagnosis of intrahepatic cholangiocellular carcinoma (1). PHL should be considered in the differential diagnosis of patients

presenting with biliary obstruction.

The authors state that they have no Conflict of Interest (COI).

Acknowledgement

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Reference

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