

patients with metastatic carcinoid [3]. It is thus argued that imaging of the brain is justified only when the patient develops symptoms from the CNS, and that even in these cases roentgenography alone may be adequate [3]. Median survival after diagnosis of brain metastases ranges from 4 to 16 months [3]. There is no established staging system and preoperative work up may include chest and abdominal CT scan, somatostatin-receptor scintigraphy, and determination of serum chromogranin-A levels [4]. Surgery is an accepted approach in case of bowel obstruction and for the management of metastatic disease in the liver manifested with carcinoid syndrome [2,5]. The case we presented herein is distinct, given the primary site, the development of brain metastases in a very short period and the unusually rapid disease progression. To our knowledge this is the first case in the literature of extra-epididymal colonic carcinoid with brain metastases.

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A large fibrolamellar hepatocellular carcinoma in a 17-year-old male body builder

Dear Editor,

A 17-year-old boy presented to our emergency department complaining of abdominal distension and bloating after meals for the last two months. Physical examination revealed a palpable liver edge. The patient had no known history of hepatitis, blood transfusion or foreign travel, but reported use of anabolic steroids for body building and weight lifting since the age of 14. The exact dosage of these medications was not known. Laboratory tests were: hemoglobin 13.4 g/dL (13-18), white blood cell count 7,700/mL (4,000-10,000), platelet count 536,000/mL (150,000-400,000), INR 1.08 (0.8-1.2), aspartate aminotransferase 98 U/L, alanine aminotransferase 112 U/L, alkaline phosphatase 160 U/L (55-140), γ -glutamyl transferase 37 U/L (7-40). The serum level of AFP was 25 ng/mL (<10). Tests for hepatitis B surface antigen and anti-HCV antibodies were negative. An abdominal ultrasound revealed a large mass in the right hepatic lobe. An abdominal CT scan confirmed a very large mass of the right liver lobe, well-defined and lobulated, measuring 14 cm in maximum diameter. Differential diagnosis included fibrolamellar hepatocellular carcinoma (FLHCC), focal nodular hyperplasia (FNH), and hepatocellular carcinoma (HCC). An abdominal MRI that followed didn't solve the diagnostic problem and surgical exploration was decided. Right hepatectomy was performed and the pathological report was: right hepatectomy containing a neoplasm of 14 cm in maximum diameter, consistent with FLHCC; surgical margins were free of disease. The patient was discharged on the 7th postoperative day and remains free of disease for 5 years as indicated by serial abdominal ultrasound.

FLHCC is an uncommon tumor affecting mostly adolescents and young adults on the second and third decade of life, lacking underlying liver disease in contrast with HCC [1]. The use of anabolic steroids in the development of hepatic neoplasms is controversial, although various studies point out to an increased cancer risk [2]. Also animal studies over the use of anabolic steroids suggest reduced lifespan due to development of tumors in the liver and kidney [3]. Clinical symptoms and findings, if any, are non specific for the disease. Serum

AFP levels are usually not elevated, but elevations in vitamin B12 binding capacity have been reported [4]. Tomographic imaging modalities usually reveal a solitary, hypervascular, heterogeneous mass that demonstrates well-defined, lobulated margins and lacks large areas of necrosis and hemorrhage that are typical of conventional HCC. Differential diagnoses for the imaging appearance of FLHCC include FNH, a large cavernous hemangioma, and HCC, because of the presence of a central scar most of the times [5]. The treatment of choice when possible is complete surgical resection; chemotherapy and radiotherapy are not effective. The median 5-year survival rate for patients with resectable FLHCC is 76%, which is much better compared with HCC patients. However, late metastasis and local recurrence have been reported [4,5].

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