

Title: Giant intra-abdominal liposarcoma

Authors:

Elena Iglesias Jorquera, Francisco Javier Álvarez-Higueras, Paula Tomás Pujante, Carmen Garre Sánchez

DOI: 10.17235/reed.2018.5176/2017 Link: <u>PubMed (Epub ahead of print)</u>

Please cite this article as: Iglesias Jorquera Elena, Álvarez-Higueras Francisco Javier, Tomás Pujante Paula, Garre Sánchez Carmen. Giant intraabdominal liposarcoma. Rev Esp Enferm Dig 2018. doi:

10.17235/reed.2018.5176/2017.



This is a PDF file of an unedited manuscript that has been accepted for publication. As a service to our customers we are providing this early version of the manuscript. The manuscript will undergo copyediting, typesetting, and review of the resulting proof before it is published in its final form. Please note that during the production process errors may be discovered which could affect the content, and all legal disclaimers that apply to the journal pertain.

Revista Española de Enfermedades Digestivas

IPD 5176 inglés

Giant intra-abdominal liposarcoma

Elena Iglesias-Jorquera, Francisco Javier Álvarez-Higueras, Paula Tomás-Pujante and

Carmen Garre-Sánchez

Digestive Tract Service. Virgen de la Arrixaca University Hospital Clinic. Murcia, Spain

Correspondence: Elena Iglesias Jorquera

e-mail: elena_ij@hotmail.com

CASE REPORT

We report the case of a 67-year-old male with epigastric pain and weight loss during

the last nine months. Physical examination revealed a hard palpable mass in the epi-

mesogastrium. An abdominal ultrasound identified a large, heterogeneous and

hypovascular mass, which compressed the left hepatic lobe and the pancreas (Fig. 1).

An abdominal contrast-enhanced computed tomography (CT) was performed and a

large soft tissue mass, polylobulated, with an intra and retroperitoneal localization was

found (Fig. 2). The diagnosis of dedifferentiated liposarcoma was achieved via a core

needle biopsy (Fig. 3). The patient was referred for surgery and a laparotomy was

performed. A mass of 30 x 30 cm and 3 kg in weight was found, which was resected

together with a segment of infiltrated small bowel. The histological diagnosis was a

dedifferentiated liposarcoma with submucosal infiltration of the duodenum.

DISCUSSION

Soft tissue sarcomas are rare neoplasms representing < 1% of tumors in adults.

Between 12-15% are located in the retroperitoneum (1). They are often incidental

findings and can reach large sizes before symptoms occur (2). The most common type

in adults over 55 years of age is liposarcoma (> 40%). Contrast-enhanced CT is the first-

line investigation and core needle biopsy usually confirms the histological diagnosis.

Differential diagnosis includes tumors with a fatty component such as renal



angiomyolipoma, adrenal myelolipoma, lipomas, hibernomas and extragonadal germ cell tumors. However, the absence of macroscopic fat on CT does not exclude the diagnosis of liposarcoma. Surgery is the definite treatment of these tumors but radiotherapy and/or systemic chemotherapy can also be associated (3).

REFERENCES

- 1. Siegel RL, Miller KD, Jernal A. Cancer statistics, 2017. CA Cancer J Clin 2017; 67(1):7. DOI: 10.3322/caac.21387
- 2. García A, Martín J, Sánchez T, et al. Liposarcoma gigante mixto de la grasa perirrenal. Rev Esp Enferm Dig 2010;102:221-2. DOI: 10.4321/S1130-01082010000300015
- 3. Messiou C, Moskovic E, Vanel D, et al. Primary retroperitoneal soft tissue sarcoma: Imaging appearances, pitfalls and diagnostic algorithm. Eur J Surg Oncol 2017;43(7):1191-8.

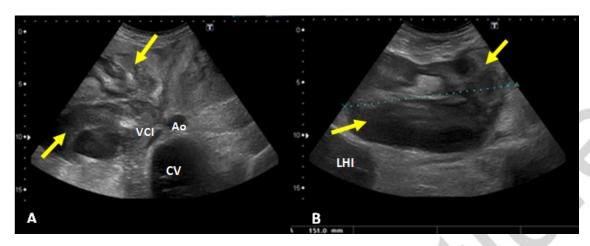


Fig. 1. Abdominal ultrasound (transverse section) showing a heterogeneous hypoechoic mass which is located next to different structures: inferior vena cava (VCI), abdominal aorta (Ao) and vertebral body (CV). B. The mass measures 15 centimeters (yellow arrows) and compresses the left hepatic lobe (LHI) and the pancreas.

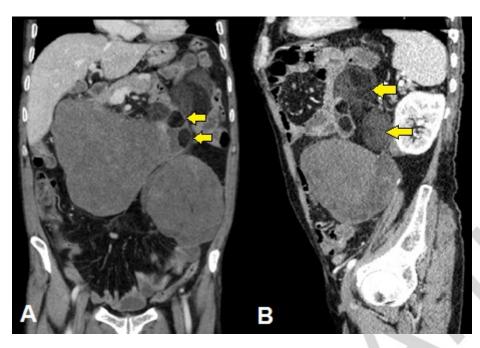


Fig. 2. A. Abdominal contrast-enhanced CT in the portal phase, MPR reconstruction and coronal section. A large bilobed intraperitoneal soft tissue mass that displaces the root of the mesentery is seen with a heterogeneous density and hypodense tracts, compatible with a fatty tumor. The arrows indicate two hypodense nodules with more fat, within the duodenum. B. Left parasagittal section: two large hypodense nodules, with a lot of fatty tissue in the anterior pararenal space and retroperitoneum (arrows).

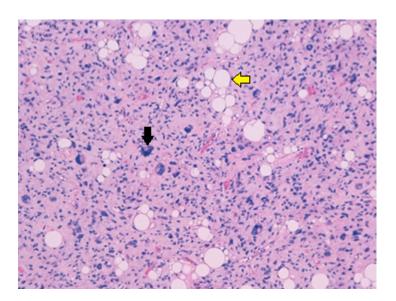


Fig. 3. Hematoxylin-eosin staining. Dedifferentiated liposarcoma: most of the tissue is composed of spindle cells, interspersed with multinucleated "floret type" cells (black arrow). Atypical adipocytes and lipoblasts are identified among them (yellow arrow).