lymphangioma, cystic mesothelioma, cystic teratoma, Müllerian cyst, lymphoceles, urinoma, etc. 

The treatment of choice of these retroperitoneal tumours is complete excision with an intact capsule. A less invasive approach like laparoscopy could provide apparently satisfactory results, but it is necessary to study the long-term progress of these cases to define the safest and most appropriate surgical technique. As for prognostic and relapse factors, these include histology grade and capsular integrity during surgical extraction. Tumour size does not seem to be a determinant prognostic factor.

Some authors have used adjuvant chemotherapy and even hysterectomy and double adnexectomy in cases of malignant cystadenocarcinoma, but there is no evidence of their usefulness due to the few cases reported.

To conclude, and in reference to our case, due to the fact that the definitive diagnosis can only be reached with the pathology analysis of the surgical specimen, we therefore believe that surgery should be indicated in future cases at the moment of diagnosis instead of periodical radiological studies.

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Francisco Javier Vicario, Laia Estalella, Judit Hermoso, Franklin Díaz, Pere Gris

Departamento de Cirugía General y del Aparato Digestivo, Parc Sanitari Sant Joan de Déu, Sant Boi de Llobregat, Barcelona, Spain

Departamento de Urología, Parc Sanitari Sant Joan de Déu, Sant Boi de Llobregat, Barcelona, Spain

*Corresponding author.
E-mail address: laia.estalella@pssjd.org (L. Estalella).

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Calcifying Fibrous Tumor of the Small Bowel

Tumor fibroso calcificante de intestino delgado

Although the majority (95%) of gastrointestinal mesenchymal neoplasms are tumors of the gastrointestinal stroma (GIST) and smooth muscle, there are also descriptions of a large variety of other tumor types with a very low incidence, such as schwannomas, desmoid tumors, solitary fibrous tumors, inflammatory fibroid polyps, inflammatory myofibroblastic tumors, etc. The main diagnostic problem presented by these uncommon lesions is that they are easily confused with certain morphological types of GIST, especially with CD117-negative GIST and with no mutations in the KIT or PDGFRA genes (wtKIT/PDGFRA).

Recently, a rare form of gastrointestinal mesenchymal neoplasms have been described, that are clinically benign, with fibro/myofibroblastic differentiation and presence of calcified foci. These have been referred to with the descriptive terms of “childhood fibrous tumor with psammoma bodies” and “calcifying fibrous tumors”. We present the case of a 46-year-old woman who was in treatment for hypertension and had no other history of interest. She consulted with her primary care physician due to recurring abdominal pain and nausea. The patient underwent...
abdominal ultrasound for suspected gallstones. The study
detected thickening of a loop proximal to the terminal ileum,
with a wall that was 5 mm thick and non-painful; there was
also a moderate quantity of fluid in the pelvis, interloop spaces
and pouch of Douglas. Colonoscopy showed evidence of a
submucosal lesion measuring 2.5×3 cm situated in the
terminal ileum, whose biopsy was very superficial and not
diagnostic. Subsequently, computed tomography with con-
trast revealed a soft-tissue density mass measuring 4.5 cm×2.8 cm×2.6 cm located in the region of the terminal ileum and ileocecal valve. The tumor protruded towards the
interior of the cecum and was well defined, with point-like
hyperdensities that were compatible with calcifications. No
dilatation of the bowel loops, locoregional lymphadenopath-
ies, or hepatic or pulmonary involvement were observed
(Fig. 1). With the clinical suspicion of ileal GIST, laparoscopic
ileocolic resection was proposed. The patient developed no
complications and was discharged 3 days after the procedure.

The pathology study of the surgical specimen reported a
well-defined, non-encapsulated mass measuring 2.5 cm at
its maximum dimension that was comprised of a hypo-
cellular fibroblastic proliferation growing within a dense
matrix with abundant collagen. Inside, there was limited
inflammatory infiltrate of plasma cells and lymphocytes as
well as areas of dystrophic calcification and psammoma
bodies (Fig. 1B–D). With immunohistochemistry techniques,
the cells presented positive immunoreaction in occasional
cells to CD34 and negative immunoreaction to CD117, DOG1,
S100 and AML. Given these findings, the diagnosis of ileal
calcifying fibrous tumor (CFT) was reached.

CFT is a soft tissue tumor that is benign and uncommon.
It was originally described in girls aged 2–11 by Rosenthal
and Abdul-Karim in 1988 as a childhood fibrous tumor with
psammoma bodies.7 Initially, these tumors were thought to
represent a reactive process resulting from abnormal
healing, thus the name “calcifying fibrous pseudotumor”.4
The name was finally changed to CFT in the consensus
classification document by the World Health Organisation
in 2002.

Cases of CFT have been reported in multiple locations, such
as the peritoneum, pleurae, mediastinum, lungs, testicles,
suprarenal glands, etc.5–8 Nonetheless, the gastrointestinal
location of CFT is especially uncommon, and only around 10
cases have been reported.9

Histopathologically, CFT are circumscribed, non-encap-
sulated tumors that are comprised of hyalinised collagen,
spindle cells without atypia, lymphoplasmacytic infiltrate,
and psammoma bodies or dystrophic calcifications.10 They
are usually incidental findings that are presented as well-
outlined nodular formations, both on imaging tests as well
as in their later histopathology study, and they show no
metastatic capability. Generally, symptoms are very nons-
ppecific, including abdominal pain and altered bowel habit,
as in the case we have presented. The differential diagnosis,

Fig. 1 – (A) Abdominal CT image showing the ileal mass with calcifications; (B) well-outlined tumour situated in the
submucosa; (C) neoplasm showing psammoma-type calcifications (HE 40×); (D) dense stromal collagen
and lymphoplasmacytic inflammatory infiltrate are observed (HE 40×).
like submucosal spindle cell tumors, includes GIST, leiomyomas or leiomyosarcomas, schwannomas, desmoid tumors and inflammatory myofibroblastic tumors, etc. Diagnosis is easily resolved with an immunohistochemistry study.

The rarity of CFT in the gastrointestinal tract makes its natural history difficult to outline, although there have been no cases of recurrence or metastasis to date. This differs greatly from cases described in other locations, where occasional recurrences have been reported. Therefore, treatment of gastrointestinal CFT should be conservative and involve radiological follow-up, especially for lesions removed by enucleation.

In short, our intention is to call attention to this peculiar entity that has easily been confused with GIST or an evolved inflammatory process.

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Enrique Rodríguez Zarooa, Ana Vallejo Beníteza1, Montserrat Mora Cabezasb, Sofía Pereira Gallardoc

aUGC Anatomía Patológica, Hospital Universitario Virgen Macarena, Sevilla, Spain
bUGC Cirugía General, Hospital Universitario Virgen Macarena, Sevilla, Spain
cCorresponding author.
E-mail address: anvaben@hotmail.com (A. Vallejo Benítez).

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Intrathyroid Parathyroid Adenoma in a Patient With Chronic Lithium Treatment

Adenoma paratiroideo de localización tiroidea en paciente bajo tratamiento prolongado con litio

Lithium carbonate is an effective medication for the treatment of bipolar disorder. Its long-term administration can stimulate the parathyroid cells and induce hyperparathyroidism secondary to parathyroid hyperplasia or adenoma. We present a case of hyperparathyroidism secondary to an adenoma located in the thyroid of a patient treated with long-term lithium carbonate therapy.

The patient is a 49-year-old woman with a medical history of bipolar disorder, for which she had been in treatment with lithium for 20 years. She had been referred to us after the detection of hypercalcaemia during routine lab work. The patient reported no symptoms, and there was no family history of endocrinological disease.

Upon physical examination, no thyroid nodules or cervical lymphadenopathies were palpated. Both the cardiopulmonary auscultation and abdominal examination were normal. Lab work showed: corrected calcium 11.7 mg/dL, phosphorus 3 mg/dL, creatinine 1.9 mg/dL, lithium 0.4 mEq/L (therapeutic