lymphangioma, cystic mesothelioma, cystic teratoma, Mülle-
rian cyst, lymphocele, urinoma, etc.\textsuperscript{6}

The treatment of choice of these retroperitoneal tumours is
complete exeresis with an intact capsule. A less invasive
approach like laparoscopy could provide apparently satisfac-
tory results,\textsuperscript{8} 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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Calcifying Fibrous Tumor of the Small Bowel\textsuperscript{3,4}

\textbf{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 schwanno-
mas, desmoid tumors, solitary fibrous tumors, inflammatory
fibroid polyps, inflammatory myofibroblastic tumors, etc.\textsuperscript{1,2} 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 \textit{KIT} or \textit{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”\textsuperscript{3}
and “calcifying fibrous tumors”.\textsuperscript{4}

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
recurrent abdominal pain and nausea. The patient underwent

\textsuperscript{3} Please cite this article as: Rodríguez Zarco E, Vallejo Benítez A, de Soto Cardenal B, Mora Cabezas M, Pereira Gallardo S. Tumor fibroso
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 contrast 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 lymphadenopathies, 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 hypocellular 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. Initially, these tumors were thought to represent a reactive process resulting from abnormal healing, thus the name “calcifying fibrous pseudotumor”.

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, pleura, mediastinum, lungs, testicles, suprarenal glands, etc. Nonetheless, the gastrointestinal location of CFT is especially uncommon, and only around 10 cases have been reported.

Histopathologically, CFT are circumscribed, non-encapsulated tumors that are comprised of hyalinised collagen, spindle cells without atypia, lymphoplasmacytic infiltrate and psammoma bodies or dystrophic calcifications. 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 nonspecific, including abdominal pain and altered bowel habit, as in the case we have presented. The differential diagnosis,

![Fig. 1](https://www.elsevier.es/)

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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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