standard treatment guidelines, neoadjuvant chemotherapy and surgical resection, or without local radiation therapy, followed by adjuvant chemotherapy appears to be the best option. In conclusion, although primary fissural PNET is an extremely rare soft tissue sarcoma, this possibility must be taken into consideration in the differential diagnosis of a primary fissural mass.

References

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Positive Positron Emission Tomography/computed Tomography Scan and Multiple Pulmonary Nodules: An Unusual Diagnosis

Tomografía por emisión de positrones/tomografía computarizada positiva y nódulos pulmonares múltiples: un diagnóstico poco habitual

To the Editor:

Primary amyloidosis is a rare disease caused by deposits of amyloid fibrils formed by insoluble protein in different tissue types. No genetic or environmental factors that might make an individual more susceptible to amyloid deposits have been identified.

Primary pulmonary amyloidosis, when deposits are limited to the lungs and associated structures with no involvement of other organs, is a very rare manifestation of the disease.

Amyloid nodules in pulmonary parenchyma are usually an incidental finding that must be distinguished from other diseases. Pulmonary amyloid nodules with uptake on positron emission tomography (PET) are very rare. To date, only 7 cases have been published in the English literature.

An 80-year-old man, a former smoker, presented with a few months’ history of dyspnea and systemic signs (asthenia, anorexia and loss of weight). Physical examination and laboratory tests results did not reveal any significant changes. Chest X-ray showed multiple dissever nodular lesions in both lung fields. Chest and abdominal computed tomography (CT) found dissever nodular pulmonary lesions with lobulated borders and no calcifications, in both lungs (Fig. 1). No microorganisms or malignant cells were identified in bronchoscopy specimens. PET-CT results suggested that some of the lesions in both lungs had significantly higher affinity for FDG than others, and there were no lesions with increased uptake in any other part of the body. Lung function tests showed moderate airway obstruction. Transthoracic and transbronchial biopsies showed amorphous amyloid-type cells. Two years after diagnosis, the patient remains in good clinical condition under treatment with inhaled long-acting beta-antagonist bronchodilator and corticosteroid.

Primary pulmonary amyloidosis occurs in 4 forms: tracheobronchial amyloidosis, nodular pulmonary amyloidosis, infiltrating interstitial amyloidosis and amyloidosis of the lymph nodes. The anatomical and functional evaluation of respiratory tract amyloidosis should be evaluated using standard X-ray, CT, endoscopy and lung function testing.

From a radiological point of view, primary nodular parenchymal pulmonary amyloidosis occurs in the form of single or multiple nodules in any lobe, and must be taken into account in the differential diagnosis of primary or metastatic pulmonary malignancies, pulmonary tuberculosis, pulmonary sarcoidosis, and connective tissue diseases. Nodules are generally peripheral and subpleural and 4 typical features are observed on CT: well-defined lobulated borders, calcification that is often central or forms in an irregular pattern within the nodule, multiple shapes and sizes, and slow growth, often over years, with no remission.

F-FDG PET-CT is widely used in the detection of cancer. As with other diagnostic procedures, some results may be false positives, and some non-malignant diseases, such as tuberculosis, sarcoidosis, fungal disease, interstitial lung disease, arthrosis, vascular thrombosis, osteoporosis and rheumatoid nodules, can show increased metabolic activity. In our case, multiple pulmonary amyloidosis nodules with moderate F-FDG uptake were observed, an observation also reported by Khan et al. and Zhang et al.

Pathological examination of a tissue sample is the standard reference for diagnosing amyloidosis.

On optical microscopy, hematoxylin and eosin staining shows homogeneous, pinkish, amyloid substances with no cell structure.

Fig. 1. Chest computed tomography showing dissever pulmonary nodules with lobulated borders.
often associated with typical fibrosis, and a red-green birefrin-
gence on polarized light microscopy of Congo-red stained tissue.\textsuperscript{4} Information on prognosis and treatment of amyloidosis in the lung is
taken from testimonial reports. To date, no fully effective treatment
is available.\textsuperscript{1} Nodular disease may progress slowly, with lesions
increasing in size and number.\textsuperscript{1} Some patients have an excellent
prognosis,\textsuperscript{2} while in others, disease progresses until death from
respiratory failure.\textsuperscript{1}

Although primary pulmonary amyloidosis is a very rare dis-
order, a definitive diagnosis may mean that the patient can be
correctly monitored and unnecessary interventions and treatments
be avoided.

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Recurrent Tracheoesophageal Fistula in
Children With Repaired Esophageal Atresia
and the Usefulness of Flexible Bronchoscopy\textsuperscript{2}

Fistula traqueoesofágica recurrente en niños con atresia
esofágica reparada y utilidad de la broncoscopia flexible

To the Editor:

Diagnosing recurrent tracheoesophageal fistula (TEF) in a child
with congenital esophageal atresia repair during the neonatal
period who has recurrent or persistent respiratory symptoms is
a real challenge, since medium leakage in the gastrointestinal tract
may be missed on standard contrast studies.\textsuperscript{1,2} We present a case
that illustrates the importance of high-pressure administration
of contrast medium in the esophagus via a catheter, for visualization
of recurrent TEF, and the additional utility of flexible bronchoscopy
for diagnosis.

A 3 \(2\frac{1}{2}\)-year-old boy was evaluated for recurrent lower respira-

tory tract infections. He had a history of congenital esophageal
atresia with distal TEF repair in the neonatal period. At the age of
2 \(2\frac{1}{2}\) years, the child was hospitalized for complicated pneumonia
with pneumothorax and pneumomediastinum, requiring admission
to the intensive care unit and chest tube placement. Chest CT
showed diffuse ground glass opacities and areas of atelectasis in
both lungs. In view of the seriousness of the clinical signs, recur-
rent TEF was considered in the differential diagnosis. The patient
had a history of anastomotic leakage, a known risk factor for fistula
recurrence,\textsuperscript{2} yet there were no signs of contrast medium leak or
esophageal stenosis in the gastroduodenal tract.

Nonetheless, the patient was admitted on 4 occasions in the fol-
lowing year for fever, cough and shortness of breath. Evaluation by
a pediatric pulmonologist was requested by the attending physi-
cian. Chest auscultation revealed crepitations and wheezing. Chest
X-rays showed areas of atelectasis in both lungs and mild gastro-
esophageal reflux was detected on pH-metry. The most important
finding on flexible bronchoscopy was a pouch in the posterior trae-
chal wall, with air bubbling during bag mask ventilation adminis-
tered by the anesthesiologist, suggesting a communication between
the esophagus and the pouch (Fig. 1 and Supplementary video 1). Lipid-laden macrophages were found in bronchoalveolar

lavage (BAL) fluid and \textit{Pseudomonas aeruginosa} was isolated from
BAL culture. Both of these data suggest chronic aspiration

and lower respiratory tract infection. Since no diagnostic pro-
cedures had been performed in the gastroduodenal tract, contrast
medium was injected via a catheter into the esophagus, confirming

concurrent TEF (Fig. 1 and Supplementary video 2).\textsuperscript{2}

Approximately 50% of children with history of esophageal
atresia and TEF repair have recurrent or persistent respiratory

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ica recurrente en niños con atresia esofágica reparada y utilidad de la broncoscopia

Fig. 1. (A) A tracheal pouch in the posterior tracheal wall was observed using flexible bronchoscopy (white arrow), near the carina. (B) Air bubbling was then seen during
bag valve mask ventilation administered by the anesthesiologist, suggesting a communication between the esophagus and the pouch.