Large Thoracic Teratoma with Gastro-intestinal Presentation


1 University of Limerick, Limerick, Ireland
2 Division of General Medicine, Pulmonary and Sleep Medicine, University Hospital Limerick, Limerick, Ireland

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Abstract

We report the case of a 27-year-old woman with a large mediastinal teratoma presenting with vague abdominal pain and diarrhea; she was erroneously suspected of having coeliac disease. This is the first such presentation of a mediastinal teratoma. We describe the endoscopic and radiologic presentation, the surgical management, and histologic characteristics.

Keywords: Diarrhea; Tumors; Hernia; Gastritis

Case Report

Intrathoracic teratomas are extremely rare germ-cell tumors. They typically occur in the anterior mediastinum. Intrathoracic teratomas can rupture into the tracheobronchial tree and can undergo malignant transformation. For these reasons, surgical resection is the treatment of choice. Here, we report the case of a 27-year-old woman with an intrathoracic mature teratoma, presenting with abdominal discomfort and diarrhea erroneously considered to be caused by coeliac disease.

A 27-year-old woman with no past medical history was admitted to the hospital for evaluation of nausea, epigastric discomfort and loose stools. Her physical exam was unremarkable. On the suspicion of coeliac disease, the patients underwent upper endoscopy revealing only a hiatal hernia and gastritis. She also underwent colonoscopy which was unremarkable, as were multiple biopsies of the upper and lower gastrointestinal tract. Diagnosis of coeliac disease was excluded. The patient was treated with proton pump inhibitors yet her symptoms persisted. Accordingly, she underwent computer tomography (CT) of the abdomen that incompletely visualized a large mass above the right hemi diaphragm. This finding prompted respiratory team consultation. A dedicated CT of the thorax was performed. The latter demonstrated a large right-sided mediastinal mass (Figure 1).

Figure 1: Computed tomographic scan of the thorax obtained with intravenous contrast demonstrate a large heterogeneous opacity (12 cm × 12 cm × 13 cm) occupying more than two-thirds of the right hemithorax. The mass partially compresses the right atrium and the right pulmonary vein. The presence of air within the mass indicates the presence of air likely arising from a communication between the teratoma and the bronchial system.

The mediastinal mass, arising adjacent to the thymus, was resected (Figure 2), and a mature benign teratoma with components of all three germ-cell layers was diagnosed on pathological evaluation (Figure 3). No immature elements or germ cells or somatic malignancy was identified. A resected mediastinal lymph node had no evidence of malignancy. Following surgery, the patient made an excellent recovery. Epigastric discomfort and loose stools resolved post-operatively.
Figure 2 Gross Specimen of the resected mature teratoma. On cut section the mass, 526 g in weight and 150 mm in maximum dimension, was mainly a multicystic tumor. Some cysts contained yellow material, while other cysts contained tan-white material.

Figure 3 Microscopic view of the mature teratoma, haematoxylin and eosin stain. A) Respiratory mucosa subtended by connective tissue (high magnification). B) Sebaceous glands (low magnification). C) Well-formed hair (low magnification). D) Ganglion nerve (low magnification). The teratoma contained also keratin, bone, and bone marrow and nerve elements (not shown).

Discussion

Teratomas, the commonest type of germ cell tumor, arise during the early stages of embryonic development. Due to their slow growth, teratomas are usually diagnosed between the second and fourth decade of life [1]. Whether the site of primordial origin is the third bronchial cleft or from nests of germinal cells along the urogenital ridge remains controversial [2,3].

Histologically, teratomas can be classified as immature and mature [4]. Immature teratomas, which account for approximately one third of all mediastinal germ cell tumors [5], contain fetal and neuroendocrine tissue [4]. Mature teratomas, which account for approximately two thirds of all mediastinal germ cell tumors [6], contain at least two of the three germ layers found in embryonic life: ectoderm, mesoderm and endoderm [2]. Immature teratomas are usually malignant in nature. In contrast, less than 1% to 2% of mature teratomas undergo malignant transformation [2,7,8], including transformation to sarcomas [9] and adenocarcinomas [8]. Two mechanisms may be involved with malignant transformation. In some patients, particularly if elderly, cells in the teratoma spontaneously become malignant. In other patients, particularly younger ones, the malignant transformation may be triggered by administration of chemotherapy or radiotherapy [10].

Teratomas are usually located in or near the anatomical midline-gonads, retroperitoneum and sacrococcygeal region [11], the path of germinal cell migration during embryogenesis [3]. Less than 3% are located in the mediastinum [4]. Most teratomas in the mediastinum are mature teratomas, and they account for approximately 10% of all mediastinal tumors [4]. Half of all patients diagnosed with a mediastinal teratoma are asymptomatic [12]. When present, symptoms may include dyspnea, back or shoulder pain, bulging of the anterior chest wall, hemoptysis, recurrent pneumonia and, rarely, superior vena cava syndrome [7]. Mature mediastinal teratomas can occasionally invade the surrounding structures or they can rupture into the mediastinum or pleura. Rupture of a mature teratoma into an airway can trigger expectoration of sebum or the pathognomonic expectoration of hair (trichoptysis) [4]. Bronchiectasis secondary to exocrine secretions form the tumor have been described in patients with mature teratomas containing pancreatic acinar tissue [9].

Typical in the presentation of our patient was her age, 27 years, the mean age at diagnosis of mature teratomas [7]. Epigastric pain and diarrhea, however, are unusual. Indeed, to the best of our knowledge, epigastric pain and diarrhea have never been described as the presenting symptom in patients with mediastinal teratomas.

Nearly 40% of mediastinal teratomas rupture, most frequently into the lung and bronchial tree, followed by the pleural space, pericardial space with secondary tamponade, or great vessels [13]. None of this occurred in our patient. Accordingly, none of these complications could have been responsible of the patient’s symptoms. Teratomas tend to accelerate in growth during adolescence and early adulthood [6]. Whether accelerated growth contributed to our patient’s symptoms is impossible to determine considering the lack of chest imaging before her clinical presentation.

In this patient, the close proximity of the mature teratoma to the thymus, which was included in the surgical resection, could be a clue that the teratoma derived from the third endodermal pharyngeal pouch where the initial clustering of embryonic cells or analge for the thymus reside [14].

Typically, as was the case in our patient, CT images of mature teratomas demonstrate a heterogeneous soft-tissue rounded mass with sharp margins in the anterior mediastinum that combines soft tissue, punctate calcification, fat attenuation and fluid-containing cysts. Only occasionally magnetic resonance
imaging, bronchoscopy and CT-guided biopsy are necessary in the diagnostic workup of suspected mature teratomas [4]. When the teratoma ruptures into a bronchus, hair-like material may be seen during bronchoscopy [12]. Laboratory exams are usually noncontributory yet; a Japanese area(s) and may affect only one of the tissue components of an otherwise benign-looking tumor [8,16]. Most mature teratomas can be excised en bloc as local invasion is rare yet described [17]. In the absence of local invasion, recurrences are rare [18-20]. When local invasion occurs, subtotal resection is worth pursuing as it can provide symptomatic relief [2].

For several reasons, aggressive surgical intervention should always be pursued in the treatment of mature teratomas. First, it allows definitive confirmation of the diagnosis. Second, it addresses presenting symptoms. Third, it prevents future complications-including life threatening hemoptyis and malignant transformation. When it occurs, malignant transformation, now referred to as germ cell tumor with somatic-type malignancy [8], can be difficult to diagnose in its early stages because malignancy may be restricted to localized area(s) and may affect only one of the tissue components of an otherwise benign-looking tumor [8,16]. Most mature teratomas can be excised en bloc as local invasion is rare yet described [17]. In the absence of local invasion, recurrences are rare [18-20]. When local invasion occurs, subtotal resection is worth pursuing as it can provide symptomatic relief [2].

References