Thymic mucoepidermoid carcinoma: A case report

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Abstract
Mucoepidermoid carcinoma of the thymus is a rare malignant neoplasm of the anterior mediastinum. There are less than 30 cases described in the English literature. We report a case of a 77-year-old man with right chest pain. Thorax CT scan showed a lobulated cystic mass having a focal solid portion with direct invasion of the adjacent pericardium in the anterior mediastinum. He underwent surgical resection for his anterior mediastinal mass via right thoracoscopic surgery without any complication. He had an uneventful recovery and was discharged on the sixth postoperative day.

Keywords: Thymus, Mucoepidermoid carcinoma, Mediastinum

Introduction

Mucoepidermoid carcinoma is a rare tumor of the lung, which arises from the minor salivary glands in the tracheo-bronchial tree. Thymic carcinomas are unusual malignant neoplasms that have wide variety of morphologic appearances. Mucoepidermoid carcinomas of the thymus are extremely rare malignant mediastinal neoplasms and account for 2% of thymic carcinomas [1,2]. We report a case of mucoepidermoid carcinoma of the thymus with radiologic and histologic findings.
Case presentation

In this case, a 77-year-old male, admitted to our hospital complaining of right chest pain was reported. The physical examination and laboratory studies were unremarkable. A 50x40 mm-sized lobulated cystic mass with focal heterogeneously solid portion in the anterior mediastinum was observed with thorax computed tomography (CT) scan of the chest (Figure 1). The patient did not show a mass with typical manifestations of myasthenia gravis. He had not ptosis and progressive muscle weakness. On thorax CT images, tumor was observed to adhere to the pericardium and infiltrated it (Figure 2). Neither pericardial/pleural effusion nor enlarged lymph nodes were detected.

Thorax CT guided fine needle biopsy of the anterior mediastinal mass was not performed. Thymectomy and resection of the adherent tissues were performed with the right video-thoracoscopic surgery. There were no intraoperative complications. Grossly, resected tumor measured 5x4 cm in size. It was a solid gray-red well-circumscribed mass with cystic compartments. Under the microscopical examination, mucous lake surrounded with atypical squamous epithelium and cylindric mucinous epithelium was seen in the thymus (Figure 3). Tumor has low mitotic activity without necrosis. Immunohistochemically, the epidermoid cells of the tumor were positive to pan-CK and p63 (Figure 4). The mucus-secreting cells are demonstrated by staining with mucicarmine (Figure 5). This part of tumor was diagnosed as low-grade mucoepidermoid carcinoma of the thymus according to the histopathological criteria of WHO classification [3].

Postoperative period was uneventful. Chemotherapy and radiotherapy of the main tumor bed were performed on the patient. Since there was a possibility of tumor metastasis to another anatomical location, the patient was referred to a whole body positron emission tomography positron emission tomography–computed tomography (PET-CT) study to search for the potentially secondary tumor, but there was not observed any abnormality. The fiberoptic bronchoscopy did not identify. Endobronchial lesion is located in tracheobronchial tree. So, the sign of the mucoepidermoid carcinoma of the lung was not detected. The patient was on regular follow-up for 4 months after discharging from hospital. He had remained asymptomatic, and there was no evidence of tumor recurrence during the period of postoperative follow-up. Informed patient consent was obtained for all procedures.
Discussion

Thymic carcinoma is a rare neoplasm of the thymus, and unlike a thymoma, it exhibits malignant features of obvious cytologic atypia, having a more aggressive propensity for early local invasion and wide spread metastases. It includes the following subtypes: Squamous cell carcinoma, lymphoepithelioma-like carcinoma, neuroendocrine carcinoma, adenosquamous carcinoma, clear cell carcinoma, papillary carcinoma, basaloid carcinoma, sarcomatoid carcinoma, and mucoepidermoid carcinoma [4,5]. The most common histologic type is squamous cell carcinoma, usually occurring in middle-aged men [6,7]. Our case was old-aged.

The prognosis of thymic carcinoma is poor because of early metastatic involvement of the pleura, lungs, lymph nodes of the mediastinum, bone and liver [8,9]. In our case, there was only pericardial invasion and the tumor was in pathological stage III according to the staging system of Masaoka. Thus, the present tumor was judged to be potentially completely resectable. Local recurrences and distant metastases are frequent, occurring at 35% and 50%, respectively. The overall survival rate at 5 years for thymic carcinoma is reported to be approximately 35% [6,7]. Thymic carcinomas have been roughly classified as either low or high-grade malignancy [6]. The low-grade tumors with relatively favorable prognoses include squamous cell carcinoma, mucoepidermoid carcinoma and basaloid carcinoma, whereas the high-grade tumors include the others with poor prognosis. Although it appears that mucoepidermoid carcinoma of the thymus is extremely rare, its histological characteristics have been well analyzed: It is composed of variably sized cysts, lobules, sheets and nests of such tumor cells as epidermoid cells, mucus-secreting cells, and occasionally intermediate type cells in a variable admixture [10,11]. In our case, similar histological and gross features were observed, and this tumor was finally diagnosed as mucoepidermoid carcinoma of the thymus. As described above, thymic mucoepidermoid carcinoma is generally considered to be a low-grade malignancy with a good prognosis [6].

The well-differentiated type, in which mitosis was virtually absent, was not fatal, while the poorly differentiated type was clinically aggressive enough to threaten life, and cellular atypia, mitosis, necrosis, and predominance of an epidermoid component were often observed. The present primary tumor regarded as the poorly-differentiated type because of the mitosis and cellular atypia. Accurate histologic subtyping, namely tumor differentiation as well as clinical staging are highly recommended for the diagnosis and treatment of this disease. Especially, when the tumor is anatomically located in contact with the pleura cavity, more accurate and careful staging may be clinically necessary in the long-term follow-up. If complete resection is possible, this type of tumor should be removed by surgery, as with other types of thymic carcinoma. Complete resection was performed in our case. Takahashi et al [12] reported that in a limited early stage of thymic carcinoma, surgical resection was adequate for treatment, although the rate of complete resection is only 20–35% [7,13,14]. However, if the tumor is advanced or recurrent, a multimodality approach, including surgical resection, radiation, chemotherapy, or a combination selected [14,15]. In our case, after surgery, multimodal therapy including chemotherapy and radiotherapy were performed.

In conclusion, we here in described a rare case of thymic mucoepidermoid carcinoma undergoing multimodality treatment, including surgical resection, irradiation and chemotherapy. Although clinical aspects of this disease are little known, it was assumed that such aggressive therapy contributed to long-term survival.

References


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