A case study of rare lympho-epithelioma like esophageal carcinoma

Abhishek Mukherjee¹, Jaydeep Nath¹, Srimanti Sinha¹ and Tridip Chatterjee²*
¹Suraksha Diagnostic Centre, Salt Lake, Kolkata, West Bengal, India
²Suraksha Genomics (R&D Division of Suraksha Diagnostic), Salt Lake, Kolkata, West Bengal, India

Abstract

Sub classification of esophageal carcinoma is important due to differences in prognosis and management. Esophageal Lymphoepithelioma-Like Carcinoma (LELC) is extremely rare, with only a few cases reported to date. Carcinoma with lymphoid infiltration in the stomach, breast or nasopharynx has a good prognosis, but in the esophagus this histological type is extremely rare and its characterization is unclear.

Review of the literature revealed case reports describing lesions with similar histology. We present a 69-year-old man with a giant pedunculated-polyoid lesion of esophagus this histological type is extremely rare and its characterization is unclear. Carcinoma with lymphoid infiltration in the stomach, breast or nasopharynx has a good prognosis, but in the esophagus this histological type is extremely rare and its characterization is unclear.

Introduction

Lymphoepithelioma-like Carcinoma (LELC) can be defined as a tumor with histological similarity to undifferentiated nasopharyngeal carcinoma, which has lymphoid stroma (lymphoepithelioma) that occurs outside the nasopharynx. LELC has been usually detected in organs like the thymic gland [1], breast [2], stomach [3], salivary gland [4] and lungs [5]. But, tumors of the esophagus other than squamous cell carcinoma and adenocarcinoma are quite rare [6]. The prognosis of patients suffering from this type of cancer has been reported to be favorable [7]. Lymphoepithelioma-like carcinoma (LELC) has been described as an un- or poorly-differentiated form of squamous cell carcinoma associated with reactive lymphoplasmyctatic infiltration. The stomach is the most common site for gastrointestinal LELC; however, esophageal involvement is occasionally observed [8]. In the stomach and breast, undifferentiated medullary carcinoma with lymphoid infiltration has been shown to have a good prognosis [9]. Esophageal LELC are primarily submucosal lesions with normal-appearing epithelial coverage and rarely can have polypoid, ulcerative, or reddish mucosal irregularity [8]. The prognosis of patients suffering from this type of cancer has been reported to be favorable [10].

Case report

The patient was a 69-year-old man presented with weight loss, loss of appetite, and dysphagia. Routine blood laboratory test results indicated no abnormality. Thoracic computed tomography (CT) examination with intravenous contrast agent injection revealed a nodular, esophageal soft tissue mass at the level of the carina. The size of the lesion was 13×10 mm axially, Paratracheal, subcarinal, and bilateral hilar lymphadenopathies were detected in the mediastinal region. Upper endoscopy revealed a giant pedunculated-polyoid lesion partially shrinking the esophageal lumen at 23 cm from the incisor. Yellowish-white exudative secretion over the polyoid lesion was present. The esophageal mucosa was irregular, reddish colored, and contained erosion from the level of the lesion to the cardiosophageal junction.

Endoscopic guided biopsy was performed and tissue material sent in 10% buffered formalin for histological opinion. Histologically, the large-sized carcinoma cells formed small focal nests and infiltrated into the submucosal layer. Prominent infiltration of T lymphoid cells, was observed between and around the carcinoma cells. Based on these histopathological features, this lesion was considered to be an LELC with poorly differentiated squamous cells. The patient’s complaint of dysphagia ended immediately after polypectomy.

Methodologies

Endoscopic guided biopsy was received in Histopathology Department. Multiple tissue fragments procured went through routine tissue processing on automated tissue processor and H & E staining was performed for microscopic examination (Figures 1-3).

Discussion

LELC was first reported by Bégin et al. in 1987 [10]. LELC is defined as a tumor with histological similarity to undifferentiated nasopharyngeal carcinoma with lymphoid stroma, and this condition has been described in various organs. However, an esophageal LELC is extremely rare. Burke first detected EBV DNA in gastric cancer that histologically resembled nasopharyngeal lymphoepithelioma and after that several reports also demonstrated a close relationship between that type of gastric cancer and EBV. However, there does not appear to be a relationship between esophageal cancer and EBV [10].

LELC tumors can arise in a multitude of locations such as the thyroid and breast, as well as gastrointestinal sites such as the biliary...
The most characteristic endoscopic appearance of esophageal LELC is submucosal tumor covered with intact or ulcerative esophageal mucosa and a depressed middle portion. Exceptional cases may have simple ulcerative morphology, polypoid lesion, or solely reddish mucosal irregularity in our case, the lesion was polypoid and the accompanying reddish mucosal irregularity was established from the lesion to the level of the cardioesophageal junction [10]. The role of diffuse infiltrating lymphocytes, consisting of a large number of T lymphocytes and a small number of B lymphocytes, has not yet been clarified in LELC. Two hypotheses have been proposed. In one, the presence of diffuse lymphocytes is explained by the immune response of the host against the carcinoma. In the other, the diffuse lymphocytes are explained in terms of a cell reaction caused by the cytokines produced by the carcinoma cells.

Generally, the prognosis of patients suffering from poorly differentiated esophageal squamous cell carcinoma is extremely poor. However, esophageal LELC seems to have a relatively good prognosis [7]. The prognosis is indicated by the survival curves and recurrence rates after treatments [10].

References