Primary pericardial extragastrointestinal stromal tumor: A case report and literature review

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Abstract. Gastrointestinal stromal tumors (GISTs) are the most prevalent mesenchymal tumors of the gastrointestinal tract. GISTs are considered to originate from the interstitial cells of Cajal, the pacemakers of the peristaltic activity of the gastrointestinal tract. More than 95% of GISTs express KIT protein and discovered on GIST-1. GISTs may also be encountered in locations outside the gastrointestinal tract, in which case they are referred to as extra-GISTs (EGISTS) and often behave more aggressively. This is the case report of a primary pericardial EGIST in a 53-year-old male patient, confirmed by immunohistochemistry. To the best of our knowledge, this is the third case of EGIST diagnosed above the diaphragm. Therapy with adjuvant imatinib (400 mg/day) was recommended. At the beginning of treatment, the patient responded well, however, after a further 6 months the disease progressed and the patient succumbed to the disease 18 months later. To the best of our knowledge, this is the third reported case of EGIST occurring above the diaphragm without being associated with the esophageal wall. Two cases of primary EGIST originating from the pleura were reported previously (1,2). In addition, this is the first reported case of an EGIST originating from the pericardium.

Introduction

Gastrointestinal stromal tumors (GISTs) are the most prevalent mesenchymal tumors of the gastrointestinal tract (1). GISTs are considered to originate from the interstitial cells of Cajal (ICC), which are the pacemakers of the peristaltic activity of the gastrointestinal tract. GISTs mainly emerge in the stomach (60%), jejunum and ileum (30%), colorectum, duodenum, esophagus and appendix (10%). However, they may also be encountered in locations outside the gastrointestinal tract, such as the omentum, mesentery and retroperitoneum, in which case they are referred to extra-GISTs (EGISTS) (2). More than 95% of GISTs express KIT protein and discovered on GIST-1 (DOG-1), which are considered to be the most specific and sensitive diagnostic markers for GISTs. Approximately 80% of GISTs harbor a KIT mutation and 8-10% express mutations in the platelet-derived growth factor receptor, α polypeptide (PDGFRα). Imatinib, a selective inhibitor of KIT and PDGFRα, is a treatment agent effective against EGISTS (1,2).

Case report

A 53-year-old male patient with severe dyspnea was admitted to the Acibadem Adana Hospital (Adana, Turkey). A chest computed tomography (CT) scan revealed a massive pericardial effusion and pericardiocentesis was performed. A positron emission tomography (PET)/CT scan performed following recurrence of pericardial effusion revealed right pericardial hypermetabolic mass lesions [maximum standardized uptake value (SUV), 12.1], pericardial effusion, right hilar lymphadenopathy (maximum SUV, 4.9) and multiple metastatic nodules in both lungs. There was no evidence of a primary malignancy originating from the abdominopelvic cavity. The chest CT and coronary CT angiography revealed that the pericardial mass involved the right atrium and exhibited peripheral contrast enhancement. There was no association between the pericardial mass and esophagus (Fig. 1). The lesion was considered to be inoperable; a right thoracotomy and wedge resection were performed and a small number of lung nodules were resected. Microscopically, the tumor was composed of spindled or ovoid cells with eosinophilic cytoplasm and pleomorphic nuclei with conspicuous nucleoli. Mitotic figures were frequently observed. Immunohistochemically, the tumor cells were strongly positive for cluster of differentiation 117 (CD117; c-Kit) and CD34 and exhibited heterogeneous moderate positivity for DOG-1 (Fig. 2). The Ki-67 labeling index was 61%. The pathological diagnosis was malignant EGIST. Therapy with adjuvant imatinib (400 mg/day) was recommended. At the beginning of treatment, the patient responded well, however, after a further 6 months the disease progressed and the patient succumbed to the disease 18 months later. To the best of our knowledge, this is the third reported case of EGIST occurring above the diaphragm without being associated with the esophageal wall. Two cases of primary EGIST originating from the pleura were reported previously (1,2). In addition, this is the first reported case of an EGIST originating from the pericardium.

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Figure 1. Images of (A and B) coronary CT angiography, (C) axial and (D) reformatted coronal chest CT, showing right pericardial rim-enhancing hypodense mass lesions (white arrows) involving the right atrium. The esophagus (curved arrow), aorta (arrowheads), coronary arteries (short arrows) and the remaining cardiac chambers are normal. Diffuse pericardial thickening and effusion are also present (black-on-white arrow). CT, computed tomography.

Figure 2. Immunohistochemical analysis of the resected tumor nodules. (A) The tumor was composed of spindle cells with eosinophilic cytoplasm (hematoxylin and eosin staining; magnification, x20). The tumor cells exhibited (B) strong diffuse cytoplasmic immunoreactivity for CD34; (C) intense cytoplasmic staining for CD117; and (D) moderate heterogeneous staining for discovered on gastrointestinal stromal tumor 1. CD, cluster of differentiation.
Discussion

EGISTs are considered to originate from the common precursor cells that differentiate into this ICC-derived neoplasm during development outside of the gastrointestinal tract. Another theory is that this tumor may originate from the pluripotent mesenchymal stem cells located outside of the gastrointestinal tract (3). Contrary to GISTs, the clinical findings of EGISTs are not comprehensively known and are often associated with adverse prognostic factors, such as high proliferative index, large size, lymph node involvement and distant metastasis. The emergence of EGIST outside the gastrointestinal tract may lead to a delay in the occurrence of clinical symptoms. A number of these cases are diagnosed at a late stage, making surgical resection difficult (1,2). A mutational study on EGISTs demonstrated that this type of neoplasm exhibits a lower frequency (41.4%) of KIT mutation at exon 11, which may be the reason underlying the good response to imatinib (4). Radiologically, these tumors are often identified in the bowel wall, involving either the mucosal or serosal surface. The tumor size ranges between 1 and 35 cm, with a median size of 5 cm. In approximately two-thirds of the cases the tumor margins are well-defined. Large tumors, in particular, may display areas of hemorrhage and necrosis, leading to a heterogeneous appearance on imaging. The enhancement pattern may be homogeneous or heterogeneous (5-7).

References