Introduction

Several previous cases with deviation of the aortic arch in patients with esophageal carcinoma have been documented (1). Almost all previous cases were due to a scarce congenital anomaly: right-side aortic arch (RAA). However, we observed an acquired “RAA-like” patient resulting from a large esophageal cancer which was pre-operatively misdiagnosed with a right-side aortic arch. Through the left thoracotomy and cervical anastomosis, complete resection of the tumor and pertinent mediastinal lymph nodes was performed and favorable short-term outcomes were achieved during the follow-up period.

Case presentation

In March 2016, a 54-year-old male was admitted to our hospital with esophageal squamous cancer and the chief complaint of progressive dysphasia which had persisted for 3 months. Before admission, esophageal endoscopy and concomitant biopsy were performed at a local hospital which revealed a squamous cancer in the middle third of the esophagus. In addition, his thoracic enhanced computed tomography (CT) on admission suggested a large esophageal cancer (Figure 1). The three-dimensional CT of the thoracic great arteries showed a possible RAA and a curved descending aorta. After preoperative evaluation, the approach of using a left thoracotomy with cervical anastomosis was successfully performed and favorable short-term outcomes were achieved. According to previous reports, and the experience of the presented case, we emphasize clear recognition of the anatomical situation in the upper mediastinum and the importance of an optimal surgical approach for esophagectomy.

Abstract: The present study is the first reported case of a patient undergoing esophagectomy with ectopic aortic arch secondary to a large esophageal cancer, which was pre-operatively misdiagnosed with a right-side aortic arch (RAA). The patient, a 54-year-old male, was first admitted to our hospital for esophagectomy owing to esophageal squamous cancer and had complained of progressive dysphasia for 3 months. Chest computed tomography (CT) revealed a mass in the middle thoracic esophagus. Furthermore, the three-dimensional CT of the thoracic great arteries showed a possible RAA and a curved descending aorta. After preoperative evaluation, the approach of using a left thoracotomy with cervical anastomosis was successfully performed and favorable short-term outcomes were achieved. According to previous reports, and the experience of the presented case, we emphasize clear recognition of the anatomical situation in the upper mediastinum and the importance of an optimal surgical approach for esophagectomy.

Keywords: Esophageal cancer; right-side aortic arch (RAA); left thoracotomy
of 80° in order to dissect the esophagus. The left recurrent laryngeal nerve was exposed and thoracic duct was ligated. Unexpectedly, when mobilizing the thoracic esophagus, we found a very large exogenous esophageal neoplasm, up to 15 cm in length and 4 cm in diameter (Figure 3), involving its near tissues and compressing the aortic arch and descending aorta to the right. The pre-operative diagnosis of RAA was corrected to an acquired transposition of the aortic arch. No malformations of the branches originating from the aortic arch were found. Without incident, subtotal esophagectomy and dissection of the regional lymph nodes was performed using open surgery in thoracic phase.

Subsequently, a diaphragmatic incision was made to expose the abdominal cavity. Through the incision, the stomach was completely mobilized and then delivered into the thoracic cavity. Afterwards, a left cervical incision was performed and a gastric conduit was reconstructed and then sent to left side of the neck. This cervical anastomosis was fashioned with a circular mechanical stapler and reinforced by hand-sewn sutures in the neck phase (2). The post-operative course was uneventful with no sign of vocal cord paralysis. Via the post-operative histopathological examination, negative surgical margin was found and clinical stage IIIC (T4bN0M0G2) was determined. Nine months after the esophagectomy, the patient died due to local recurrence after two cycles of chemotherapy.

Informed consent was obtained from the patient for publication of this case report and any accompanying images.

Discussion

RAA is most commonly due to congenital anomaly which is about 0.14% prevalent and often associated with malformations of large arteries (3). However, acquired transposition of the aortic arch has been seldom reported in patients with esophageal cancer. In the present case report, we describe an extremely unusual ectopic aortic arch attributing to a large esophageal cancer.

We misdiagnosed this case as RAA associated with esophageal cancer before surgery, a complicating condition which has been documented repeatedly (4,5). Notably, with respect to surgical intervention, comprehensive identification of the mediastinal anatomical situation should be closely focused in esophageal cancer cases with congenital or acquired anomaly of the aortic arch. If esophageal cancer is associated with vascular malformations in the thorax, a surgical plan should be made predominantly depending on the anatomic situation of the lesion and adjacent structures, particularly the recurrent laryngeal nerves. In such cases, minimally invasive surgery is difficult
to carry out. Several surgical techniques of esophagectomy have been successfully performed in RAA patients with esophageal cancer including left and right thoracotomy plus sternotomy or laparotomy (3,6-8). In this case, we performed a left thoracotomy plus cervical anastomosis due to the large and advanced lesion. Fortunately, this approach reached a promising short-term outcome.

Accordingly, for optimal management during the peri-operative period of esophagectomy, it is very important to recognize the anatomical situation in the upper mediastinum to perform a safe and curative operation.

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None.

Footnote

Conflicts of Interest: The authors have no conflicts of interest to declare.

Informed Consent: Informed consent was obtained from the patient for publication of this manuscript and any accompanying images.

References


