Migration of esophageal self-expandable metal stent to the pleural cavity

A 41-year-old man presented with dysphagia secondary to squamous cell carcinoma of the esophagus, staged as T4N1M0. A covered self-expandable metal stent (SEMS) (Choostent; Solco Intermed, Seoul, Korea) was inserted for symptomatic relief, chemoradiotherapy was performed with carboplatin and 5-fluorouracil, and this was followed by radiotherapy to deliver 39.6 Gy in 22 daily fractions.

The patient presented with cough, thoracic pain, and vomiting 3 months after SEMS placement. The chest radiograph revealed mediastinal enlargement, right pleural effusion, and infiltrates (Fig. 1). Upper endoscopy revealed patency of the stent with the distal end of the stent ending in a closed cavity (Fig. 2a), which after aspiration had an appearance compatible with that of the pleural cavity (Fig. 2b; Video 1). Computed tomography (CT) confirmed migration of the esophageal SEMS to the pleural space, secondary to tumor growth, with formation of a 105 × 42-mm cavity (Fig. 3). Multiple hepatic metastases were detected.

After medicosurgical discussion, a derivational lateral esophagostomy and jejunostomy were performed, and broad spectrum antibiotics were given. CT performed after 15 days revealed resolution of the cavity. SEMS placement provides effective palliation for patients with esophageal cancer [1]. However, its safety in patients undergoing chemoradiotherapy is uncertain [1,2]. In fact, SEMS migration is common following down-staging of esophageal carcinoma with chemoradiotherapy, with several published case reports of prosthesis migration in this context [3–5]. However, to the best of our knowledge, this is the first report of migration of an esophageal SEMS to the pleural cavity. This case is also original for the fact that migration of the prosthesis was not related to down-staging of the tumor after chemoradiotherapy. We hypothesize that fragility of the esophageal wall due to radiation injury might have favored prosthesis migration in the setting of tumor growth. This hypothesis is supported by the previously described association between SEMS placement with chemoradiotherapy and esophageal perforation [2].

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Bibliography

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