

A Unique Presentation of Esophageal Adenocarcinoma Metastasized to Muscle

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Background	We describe the case of a 68-year-old man with a history of esophageal adenocarcinoma presenting with metastases in the soft tissue of his back.
Summary	A 68-year-old male with a remote history of esophageal adenocarcinoma status postdistal esophagectomy presented to a clinic noting soft tissue masses. These masses were removed; on pathological review, they were consistent with esophageal adenocarcinoma. The patient's disseminated cancer was treated with chemotherapy, but his disease progressed, and he ultimately expired.
Conclusion	Esophageal adenocarcinoma metastasizing to skeletal muscle is a rare finding consistent with a poor prognosis. While it is tempting to attribute soft tissue swelling to common diagnoses, a careful physical examination and full body scan are appropriate at every stage.
Key Words	esophageal adenocarcinoma; metastasis; soft tissue metastasis; skeletal muscle

DISCLAIMER

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Case Description

A 68-year-old male with a history of achalasia presented with progressive dysphagia and weight loss over a six-month period. Diagnostic workup included an esophagogastroduodenoscopy (EGD) demonstrating esophageal narrowing without mass, and manometry confirmed high lower esophageal sphincter tone, consistent with achalasia.

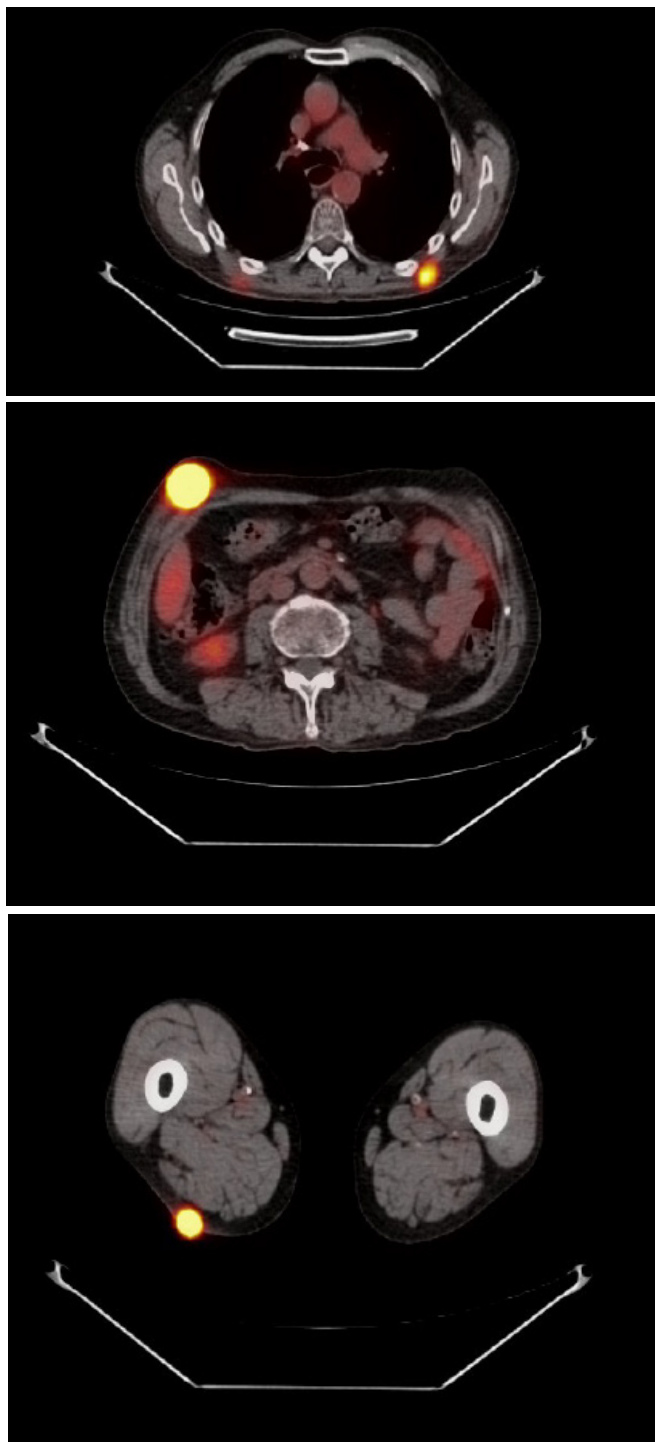
The patient initially underwent laparoscopic Heller myotomy with Toupet fundoplication without complication in March 2014. This procedure did not fully address his symptoms; despite multiple esophageal dilations, he continued to experience significant dysphagia. He then underwent laparoscopic distal esophagectomy with intrathoracic esophagogastrostomy in February 2015, which was completed without complication. The distal esophagus and proximal stomach were removed and sent to pathology, which revealed invasive moderately differentiated adenocarcinoma, pT3N1. A subsequent PET/CT was not suggestive of metastases. He received adjuvant treatment with carboplatin and paclitaxel with radiation for six weeks and demonstrated improvement, tolerating oral intake, and gaining weight. He completed his oncologic treatment and had surveillance EGD, which was negative for recurrence. At his subsequent follow-up visits, he reported that his dysphagia was resolved and his activity tolerance was improving.

In July 2019, during a routine primary care appointment, he mentioned soft-tissue masses on his back, leading to a referral to general surgery for assessment. In August, he visited the general surgery clinic at the Fargo Veterans Affairs Medical Center, noting several masses in the soft tissue of his back and arms. The masses in his back were bothersome, and he desired removal. On exam, firm and mobile masses were observed in the subscapular region on both sides of the spine, with the left side being larger than the right. They were nontender and lacked any sinus. Clinically, they appeared consistent with lipomas, leading to a referral to the minor procedure clinic for removal. Subsequently, he underwent removal of the right-sided mass under local anesthesia without any immediate complications.

During the procedure, a transverse incision was made over the right-sided mass, and dissection progressed sharply to the paraspinous muscle. The fascia was then divided along its natural tension lines. The mass, which was firm and spiculated, was observed to have multiple adhesions to the surrounding muscle but was successfully removed completely. Due to the depth and appearance of the first mass,

the decision was made to leave the larger mass until the pathology results were available. The soft tissue mass that was removed from the right subscapular region was consistent with metastatic adenocarcinoma, which aligned with the previously diagnosed gastroesophageal junction primary from four years prior.

Figure 1. PET/CT Demonstrating FDG-avid Activity in Soft Tissues. Published with Permission



Following a minor procedure, the patient received pathology results. A tumor board discussion at the VAMC determined the need for further staging, leading to a PET/CT scan in which multiple foci of FDG activity were identified in the chest, abdominal wall, and bilateral thighs. To facilitate chemotherapy, a port was placed in September 2019. The patient then underwent palliative FOLFOX chemotherapy, achieving an initial response. Unfortunately, the disease progressed, and the patient passed away in August 2020.

Discussion

Esophageal carcinoma is among the most aggressive types of cancer worldwide, with distant metastases being a major contributor to mortality rates.¹ While lymph node, liver, and lung involvement are well documented, subcutaneous tissue and skeletal muscle are rare metastatic sites. Studies report skeletal muscle metastases occurring in less than 2% of hematogenous metastases from solid tumors.^{2,6-8} Despite ongoing research efforts to elucidate the underlying mechanisms,² a comprehensive understanding of the pathophysiology behind esophageal cancer's metastatic patterns remains unclear.

Unfortunately, this lack of clarity extends to the optimal surveillance strategy for this high-risk cancer. For patients who underwent definitive chemoradiation, current recommendations suggest contrast-enhanced CT scans of the chest and abdomen every six months for up to two years, alongside esophagogastroduodenoscopy (EGD) every three to six months.³ The utility of tumor markers like carcinoembryonic antigen remains unknown. These recommendations are based on findings from the literature suggesting a predominance of locoregional relapses, typically occurring within the first 24 months posttreatment.

Esophageal cancer patients who have undergone trimodality therapy, including surgery, chemotherapy, and radiation, such as the one described in this case, are recommended to undergo CT scans of the chest and abdomen every six months for two years by the National Comprehensive Cancer Network, with additional EGD performed as needed. This surveillance strategy prioritizes detection of distant metastases, which are the most frequent relapse pattern and typically occur within 36 months of surgery.⁵

Limited case reports suggest that the most common clinical presentation of distant esophageal metastases is a soft, bothersome swelling. These lesions can appear malignant, as reported by Chand et al., or benign, as in our case. Regardless of presentation, diagnosis usually involves positron emission tomography and computed tomography (PET-CT) and tissue biopsy.^{6,8} Unlike the diagnostic protocol for many other soft tissue masses, ultrasound is not described in diagnosing esophageal metastases. With the escalating use of high-resolution modalities, more of these metastatic foci will likely be discovered. Local intervention is appropriate for symptomatic relief,⁸ but prognosis remains poor due to the systemic nature of disease.

In our case, the skeletal muscle metastases were the first indication that his esophageal adenocarcinoma, which had been treated almost four and a half years earlier, had recurred. His subsequent PET/CT scan demonstrated additional areas of FDG-avidity that were not clinically apparent, underlining the role of these advanced imaging modalities. We elected to remove the symptomatic lesions as a palliative measure.

Conclusion

Skeletal muscle metastases are an exceptionally rare complication of esophageal adenocarcinoma and represent a poor prognosis. A high suspicion, thorough physical exam, and use of PET/CT are necessary to diagnose these lesions. As diagnostic imaging improves, the incidence of these rare metastases will likely increase.

Lessons Learned

While esophageal adenocarcinoma metastasizing to skeletal muscle is rare, keeping a high index of suspicion can allow for appropriate workup and identification of this serious clinical finding.

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