Case Report

Spontaneous sacral fracture with associated acrometastasis of the hand

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ABSTRACT

Background: Acrometastases, secondary tumors affecting oncological patients with systemic metastases, are associated with a poor prognosis. In rare cases, acrometastases may precede establishing the primary tumor diagnosis.

Case Description: A 72-year-old female heavy smoker presented with low back pain, and right lower extremity sciatica/radiculopathy. X-rays, CT, MR, and PET-CT scans documented primary lung cancer with multi-organ metastases and accompanying pathological fractures involving the sacrum (S1) and right 4th digit. She underwent a S1 laminectomy and amputation of the distal phalanx of the right fourth finger. The histological examination documented a poorly differentiated pulmonary adenocarcinoma infiltrating bone and soft tissues in the respective locations. The patient was treated with a course of systemic immunotherapy (i.e. pembrolizumab). At 6-month follow-up, the patient is doing well and can stand and walk without pain.

Conclusion: Spontaneous sacral fractures may be readily misdiagnosed as osteoporotic and/or traumatic lesions. However, in this case, the additional simultaneous presence of a lytic finger lesion raised the suspicion that these were both metastatic tumors. Such acrometastases, as in this case attributed to a lung primary, may indeed involve the spine.

Keywords: Acrometastases, Elderly, Hand metastases, Immunotherapy, Sacral fracture, Spine surgery

INTRODUCTION

Acrometastases comprise the 0.1% of all bone metastases, including those found in the spine.[1] They may precede establishing the primary diagnosis of an underlying malignant tumor in 10% of cases.[2] These lesions are often detected in oncological patients with systemic metastases, and are poor prognostic findings (i.e. correlating with short life expectancies typically of <6 months).[3] Here, we describe a 72-year-old female with an unknown primary lung cancer who presented with 2 spontaneous pathological fractures involving the sacrum, and 4th digit of the right hand.
CASE ILLUSTRATION

A 72-year-old heavy female smoker, with no history of trauma, presented with 20 days of low back pain, right lower extremity sciatica, and a lesion of the right 4th digit. On examination, the patient had right S1 motor/sensory deficits, and ecchymosis/swelling of the distal phalanx of the right fourth digit [Figure 1]. The lumbar MR showed a S1 pathological fracture/lytic lesion (i.e. hyperintense at S1/S2 on T2-weighted images), and a second lytic lesion of the right 4th digit.

Diagnostic evaluations

**Chest X-ray**

The chest X-ray revealed a radiopaque lesion in the right mid-upper lung, with atelectasis and slight retraction of the superior mediastinum [Figure 2].

**Hand X-ray**

The hand X-ray showed soft tissue thickening of the right ring finger’s distal phalanx, with a distal phalanx fracture and bone loss/lytic process.

**Full body CT**

The full-body CT scan documented a large thoracic mass with inhomogeneous contrast enhancement. There were also hypodense areas in the anterior segment of the right upper lobe (90 × 84 mm), with infiltration of the bronchial and vascular branches. An adjacent nodular lesion (11 × 11 mm) and an additional opacity were distinguished in the lateral basal segment of the left lower lobe (5 × 5 mm; considered satellite metastases). Several lymphadenopathies were appreciable in multiple thoracic locations. Osteolytic areas with fractures were detected in the right pubic ramus and at S1 [Figure 3]. Additional metastases were located in the orbit, in the pterion, and in the spleen.

**PET-CT**

The 18F-FDG PET-CT scan showed accumulation of tracer at multiple sites: in the right adrenal gland (SUV max 11.2), in the middle and upper lung lobes (SUV max 15.8), and in several thoracic lymph nodes (SUV max 16.6). Tracer uptake was also found at T7, T8, T10, L3, the sacrum, the right acetabulum and ischiopubic branch, and the fourth finger of the right hand (SUV max 18.3).

![Figure 1: Anterior (a) and lateral view (b) of the right fourth finger with ecchymosis and swelling of the distal phalanx.](image1)

![Figure 2: Anterior-posterior chest X-ray showing a radiopaque area in the right mid-upper lung, with atelectasis and slight retraction of the superior mediastinum.](image2)

![Figure 3: Thoracic CT scan showing a large thoracic mass with inhomogeneous contrast-enhancement and hypodense areas, probably colliquative, in the context of the anterior segment of the right upper lobe (90 × 84 mm), with infiltration of the bronchial and vascular branches.](image3)
Lumbar MR

The lumbar MRI study revealed morphological alterations involving the L3, S1, and S2 levels that markedly homogeneously enhanced with contrast [Figure 4].

Surgery at S1 and removal of distal phalanx right fourth digit

The patient underwent a S1 decompressive laminectomy with concurrent biopsy of the S1 vertebral body, lumbopelvic fixation (L4-L5-ileum), and thermal ablation of the L3 lesion. The distal phalanx of the right fourth finger was also amputated.

Postoperative course

The patient recovered postoperatively, without any complications, and was able to stand/walk after 2 weeks of rehabilitation. The postoperative CT scan documented adequate neural decompression and screw placement [Figure 5].

Histology

The histological examination revealed a poorly differentiated pulmonary adenocarcinoma infiltrating the bone and soft tissues at S1 and the 4th digit. Immunohistochemical stains were positive for: CKAE1/AE3, CK7, EMA, vimentin, and TTF1 [Figure 6]. Tumor Proportion Score was positive (neoplastic cells ≥50%). The examination for EGFR, BRAF exons mutation was negative. No rearrangements of the ROS1 and ALK genes were detected.

Immunotherapy and radiotherapy

The patient received a course of immunotherapy (pembrolizumab 400 mg IV/42 days) and targeted radiotherapy (20 Gy/4 Gy/fraction) to treat the osteolytic lesions in the right pubic ramus.

Clinical outcome at 6 postoperative months

At 6-month follow-up, the patient can stand and walk without any pain. However, she reports the new onset of transient diplopia, related to the lesion in the left orbit, which regresses with corticosteroids. The last full-body PET scan showed reduced tracer uptake in previous lesions, but also new secondary lesions in the left lung, frontal lobe, parietal lobe, and C6 vertebral body.

DISCUSSION

Acrometastases characterize a heterogeneous group of rare secondary tumors.\(^1,2\) Lung cancer represents the most frequent primary tumor. A recent systematic review\(^3\) described the 247 cases of acrometastases that mostly originated from
primary lung cancer (91; 36.8%), gastrointestinal cancers (62; 25.1%), and urinary tract cancers (33; 13.4%). While X-ray often shows non-specific osteolytic lesions of the affected finger, PET scans have a 90% sensitivity and 78% specificity in detecting benign and malignant tumor. As acrometastases most commonly occur in the terminal phase of oncological patients, the invasiveness of the surgical treatments should be carefully balanced with patient's life expectancy.

CONCLUSION

The present case suggests that, despite the poor prognosis of patients with acrometastases, systemic immunotherapy may prolong survival, thus enabling surgical spine decompression and fixation with the goal of improving the patient's quality of life.

Declaration of patient consent

The authors certify that they have obtained all appropriate patient consent.

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Nil.

Conflicts of interest

There are no conflicts of interest.

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