Case Report

Spontaneous thoracolumbar epidural hematoma in an apixaban anticoagulated patient

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ABSTRACT

Background: Spontaneous spinal epidural hematomas (SSEHs) are often attributed to anticoagulation. Although they are rare, they may contribute to significant morbidity and mortality.

Case Description: An 83-year-old female with a history of atrial fibrillation on apixaban, presented with 4 days of back pain, progressive lower extremity weakness and urinary retention. When the patient's MRI showed a dorsal thoracolumbar SSEH, the patient underwent a T10–L3 laminectomy for hematoma evacuation. Within 2 postoperative months, her neurological deficits fully resolved.

Conclusion: Apixaban is associated with SSEH resulting in severe neurological morbidity and even mortality. Prompt MRI imaging followed by emergency surgical decompressive surgery may result in full resolution of neurological deficits.

Keywords: Apixaban, Spontaneous spinal epidural hematoma, SSEH

INTRODUCTION

Spontaneous spinal epidural hematoma (SSEH) is rare (affecting 0.1/100,000 persons/year) and can potentially result in life-altering neurological disability and/or mortality.²⁴ Based on numerous studies, patients on anticoagulation (i.e. warfarin or heparin) are susceptible to developing SSEH.³ However, there are only scarce reports regarding the impact of newer oral anticoagulants (NOAC) on the incidence of SSEH.³

Here, an 83-year-old female on apixaban developed a thoracolumbar SSEH (levels T10–L3). Following an urgent MRI, the patient underwent a T10–L3 laminectomy. Within 2 months postoperatively, her preoperative lower extremity paresis and urinary retention fully resolved.
CASE DESCRIPTION

An 83-year-old female with atrial fibrillation on apixaban (Eliquis 2.5 mg bid) presented with back pain (of sudden onset 4 days before). Her symptoms included worsening lower extremity paresis (left 0/5 and right 3/5), an L3 sensory level, and urinary retention. Although all laboratory studies were normal, the noncontrast thoracolumbar CT and MR studies showed a slightly hyperdense, posterior, extradural lesion, extending from T10 to L3 with maximal compression at T12/L1, resulting in significant cord compression [Figure 1a-e].

Figure 1: (a) Sagittal CT depicts the ventral borders of the epidural hematoma (blue arrows). (b and c) Sagittal STIR (b) and T2W (c) MRI clearly delineating the hematoma in different stages of hemoglobin degradation. (d) Axial T1W MR at the level of maximal compression (T12–L1) showing hematoma occupying almost the entire spinal canal. (e) Axial T2 MRI at the level of maximum compression (T12–L1). (f) Postoperative sagittal CT showing the extent of the laminectomy and complete hematoma evacuation.
Surgery

The patient underwent a T10–L3 laminectomy for evacuation of the SSEH. Immediately postoperatively, the patient was noted to have recovered distal motor strength of 1/5 on the left (ankle plantar flexion and dorsiflexion). Notably, the postoperative CT scan demonstrated complete evacuation of the hematoma, with no postoperative bleeding complications [Figure 1f].

Postoperative course

The patient was transferred to a Spinal Rehabilitation Center on the 2nd postoperative day; still largely paraplegic. Bridging anticoagulation with enoxaparin 4000 IU/0.4 ml was initiated on the 3rd postoperative day and was replaced by the patient’s previous apixaban regimen (2.5 mg bid) on the 20th postoperative day. When discharged after 45 days, she was neurologically normal, with only a residual mildly reduced left tibialis anterior muscle strength (4/5).

DISCUSSION

SSEH is rare and carries the risk of significant morbidity and/or mortality. Patients typically average 63 years of age and typically present with sudden onset back pain, radicular pain, sensorimotor deficits, and sphincter dysfunction reflecting the level of the bleed.1,5

The most common etiology is spontaneous rupture of the dorsal internal vertebral venous plexus, which is a valveless system susceptible to abrupt pressure changes.5,8 Bleeding of arterial origin is less frequently encountered.8

Most SSEH occur in the cervicothoracic or the thoracolumbar region, with two-thirds extending over 2–5 vertebral segments.5

Diagnostic studies

MRI is the gold standard in assessing SSEH. Hematomas are usually hypointense. Although high-resolution CT may also identify these lesions, MRI is clearly the superior diagnostic study.1,4,6

Apixaban anticoagulation predisposing to SSEH

Anticoagulation is a major predisposing factor to SSEH.5 Apixaban was the latest of the commonly prescribed NOACs to be correlated with SSEH.1,6 There have recently been three reported cases of SSEH associated with apixaban, all of which were located on the cervical spine, in elderly patients and were treated by laminectomy for clot evacuation [Table 1].1,6 In our case, surgical hematoma evacuation resulted in complete resolution of preoperative neurological deficits.

Management of SSEH

Symptomatic SSEH should be managed based on emergency MRI studies; many will require prompt laminectomy.3,5 Initial neurological status at presentation is an important determinant of outcome, as patients with incomplete neurological deficits tend to fare better than those with complete injuries; better initial ASIA score predicts better outcomes.7

CONCLUSION

SSEH, recently associated with apixaban, is a rare medical emergency requiring emergency MRI studies and operative decompression to achieve the best outcomes.

Declaration of patient consent

Patient’s consent not required as patients identity is not disclosed or compromised.

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Conflicts of interest

There are no conflicts of interest.
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