The spinal epidural abscess (SEDA) is a medical and surgical emergency whose evolution may involve functional prognosis and/or even vital prognosis. This rare pathology is seen most often in the context of immunosuppression and the clinical picture of classical triad of pain, fever and neurological deficit.

Most of the spinal epidural abscesses are thought to result from the haematogenous spread of bacteria usually from a cutaneous or mucosal source. The sources of bacteraemia are thought to be from furuncles, pharyngitis, and dental abscesses. Direct spread of infection into the epidural space from a source adjacent to the spine-for example, spondylitis or discitis-has also been described. Those having spread from hematogenous route usually are located in posterior aspect of the spinal canal while those from direct source has their locations are in anterior spect of spinal canal. Direct spread from a paraspinal abscess and Vertebral osteomyelitis are also source of abscess. blunt trauma, epidural catheter insertion and postoperative are also source of abscess. MRI plays a decisive role in its diagnosis (Figure 1).

While antibiotic therapy is the key to its treatment, an early surgical decompression is sometimes necessary to preserve the functional prognosis and to prevent neurological deficits.

We report below the cases of patients admitted for management of spinal epidural abscess in the neurosurgery department at Hassan II University Hospital of Fez.
Material and Methods:

This is retrospective study conducted at Hassan II University Hospital of Fez, of five cases of spinal epidural abscess that were treated in the neurosurgery department between 2004 and 2017.

Results:

All patients were male with a mean age of 52.4 years, ranging from 18 years to 67 years. Type 2 diabetes was found in two of our patients representing a percentage of 40% of this cohort. The average admission time was 39.6 days with extremes ranging from 3 to 60 days? Fever and spinal syndrome had been found in all of our patients while the neurological deficit was found in 4 of our patients. Spinal MRI was emergently performed in all of our patients and the thoracic region was the most common location (Table 1).

Discussion:

Spinal epidural abscess is a rare pathology (0.2-2.8 per 10 000 person / year), preferentially affecting those over 50 years of age usually in an immunocompromised persons1,9,10,11,12,13,14. The most offending organism being Staphylococcus aureus.3,8,9,15,16 Mycobacterium tuberculosis, Streptococcus milleri, Haemophilus parainfluenzae, Brucella species, and Actinomyces israelii are amongst the many other isolates described. Disseminated fungal infections such as cryptococcosis, aspergillosis, and blastomycosis are rare causes and usually arise in immunocompromised patients. Aspergillus species are known to cause spinal epidural abscess in patients with AIDS.13,14,15

The described sites of spinal epidural abscess are variable. Only 20% occur anterior to the spinal cord. Abscesses in the cervical or upper thoracic region are not uncommon. Larger studies, however, describe a preponderance of lower thoracic and lumbar abscesses. The patients with lumbar abscesses may be misdiagnosed as having a herniated lumbar disc.3,13,14

Most of the posterior SEDA have a location remote from a site of infection (dental abscess, pharyngitis, skin infection) and is most often associated prior osteomyelitis or spondylodiscitis.3,14

The classic triad of fever, spinal syndrome and neurological deficits remain unspecific. Heusner described 4 stages of clinical evolution of the spinal epidural abscess5. The 1st stage is characterized by a spinal syndrome and fever, 2nd stage by spinal irritation, the 3rd stage with sensory loss and / or sphincter dysfunction and the 4th stage by a heavy motor deficit.

MRI is the examination of choice, and helps in the positive diagnosis of the SEDA and search of the initial infection site. Spinal CT myelogram has a similar sensitivity to MRI but entails a significant risk of spread of infection.6

<table>
<thead>
<tr>
<th>Case</th>
<th>Spinal location</th>
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<tbody>
<tr>
<td>Patient 1</td>
<td>Lumber L2-L5</td>
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<tr>
<td>Patient 2</td>
<td>Cervico-thoracic C4-T4</td>
</tr>
<tr>
<td>Patient 3</td>
<td>Lumber L5-S1</td>
</tr>
<tr>
<td>Patient 4</td>
<td>Thoracic T3-T10</td>
</tr>
<tr>
<td>Patient 5</td>
<td>Cervico-thoraco-lumber</td>
</tr>
</tbody>
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Table 1 : Location of SEDA in our series (5 cases)
SEDA is a medico-surgical emergency, early treatment based on antibiotics for (4-6 weeks) and a surgical intervention for compression is essential for a good functional prognosis. Surgical treatment entails emergency evacuation of the pus with decompression of the spinal cord and nerve roots. In the presence of paraparesis, the outcome depends on the time that elapses from its onset to surgery. The longer the lapse the greater is the usual long term legacy. The appearance of a motor neurological deficit exceeding the 36h generally entails poor prognosis for functional recovery.7

**Conclusion:**

The SEDA is a rare pathology, most often seen in a context of immunosuppression. The clinical picture is nonspecific and requires biological and radiological investigations. The therapeutic treatment must be started early for good functional and vital prognosis. The appreciable neurological recovery seen in some patients reflects the value of prompt diagnosis and early treatment. The key to successful management is early diagnosis, which requires that involved clinicians consider the diagnosis. Repeated spinal and neurological examinations are essential in any patient with an unknown focus of infection and when there is spinal pain or tenderness full investigation is warranted. The increasing availability of MRI may mean that earlier diagnosis and lower morbidity from this condition should become a reality in the near future.

**References:**


