

Chronic Osteomyelitis of the Lumbar Transverse Process

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Pyogenic spondylitis involving only the posterior element of a vertebra is rare. To the best of our knowledge, there have been no reports of osteomyelitis of the transverse process. We report here on a 45-year-old male with a one month history of swelling associated with lower back pain. The magnetic resonance imaging showed a paraspinal soft tissue mass, and computed tomography revealed a fine osteolytic lesion in the right transverse process of the 5th lumbar spine, and this was all consistent with chronic osteomyelitis. A mixed staphylococcal infection was identified. Open drainage, resection of the transverse process and intravenous injection of anti-staphylococcal antibiotics resolved the back pain and reduced the erythrocyte sedimentation rate to normal. Pyogenic osteomyelitis of the transverse process is extremely rare, which can cause a misdiagnosis or a delayed diagnosis. Careful consideration of this disease is needed when evaluating patients who complain of back pain.

Keywords: *Osteomyelitis, Transverse process, Lumbar spine*

Primary spondylitis of a vertebral body rarely progresses to involve the posterior element.¹⁾ Less than 5% of the cases are known to involve only the posterior element. Most pyogenic vertebral infections involving the posterior element occur at the articular facets. There are only a few case reports of involvement of the pedicle, lamina and the spinous process.²⁻⁴⁾

To the best of our knowledge there has been no report of osteomyelitis involving the transverse process of the spine. Our report is the first report of osteomyelitis involving the transverse process of the spine.

CASE REPORT

A 45-year-old male, who was transferred from a local clin-

ic, presented with a one month history of a swollen painful back. He had experienced intermittent back pain over the previous 2-year period. He denied a history of acupuncture or local injections. A mass-like swelling was palpated on the right side of his back. The range of motion of his back was slightly limited, but there was no radicular pain. The motor, sensory and deep tendon reflexes were normal in both lower extremities.

The white blood cell count and the erythrocyte sedimentation rate were 9,000/mm³ and 76 mm/hour, respectively. The plain radiographs indicated only subtle changes in the right transverse process of the 5th lumbar vertebra (Fig. 1). T2 weighted axial magnetic resonance imaging (MRI) revealed a space-occupying lesion, which was located behind the right paraspinal muscle, and it was connected to the paraspinal muscle (Fig. 2). Enhanced computed tomography (CT) revealed a 7 × 4 × 1 cm-sized space-occupying lesion with a rim enhancement pattern and swelling of the surrounding soft tissues. CT also revealed an osteolytic lesion (2 × 1 cm in size), which was located in the right transverse process of L5, and it was similar to Brodie's abscess in the long bones (Fig. 3).

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Open drainage of the abscess and a resection of the right transverse process of L5 were performed under general endotracheal anesthesia (Fig. 4). Grossly, the osteolytic lesion was surrounded by a sclerotic bony margin and there was pus in the abscess cavity, and the pus had gravitated anteriorly and posteriorly to the paraspinal muscle; this suggested chronic osteomyelitis. The pus culture demonstrated a mixed infection with *Staphylococcus aureus* and coagulase-negative staphylococcus. The histological examination revealed an infiltration of chronic inflammatory cells into the dense lamellar bone spaces, which was all consistent with chronic osteomyelitis (Figs. 5 and 6).

The patient received an intravenous injection of 1st generation cephalosporin (cefotiam hydrochloride 1.0 g

bid) for 4 weeks, which resulted in the complete resolution of symptoms and a decrease in the erythrocyte sedimentation rate to normal and this lasted during follow-up. At the two year follow-up, the patient had normal spine functions.

DISCUSSION

Many spinal infections occur as a result of the hematogenous seeding of bacteria from various distant sites. Batson⁵ first proposed the valveless venous system as a route

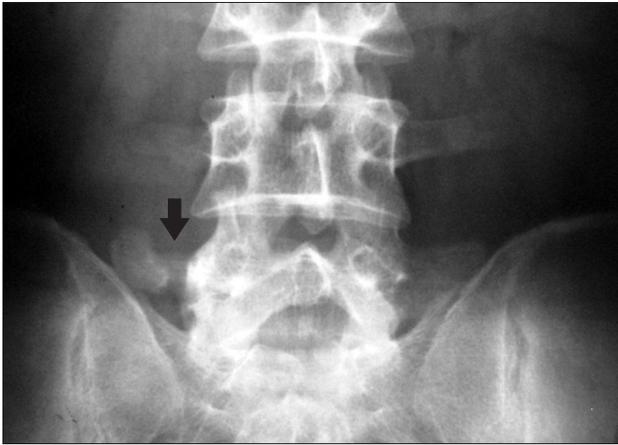


Fig. 1. The plain radiograph shows a subtle osteolytic lesion with surrounding sclerosis of the right transverse process of L5 (arrow).

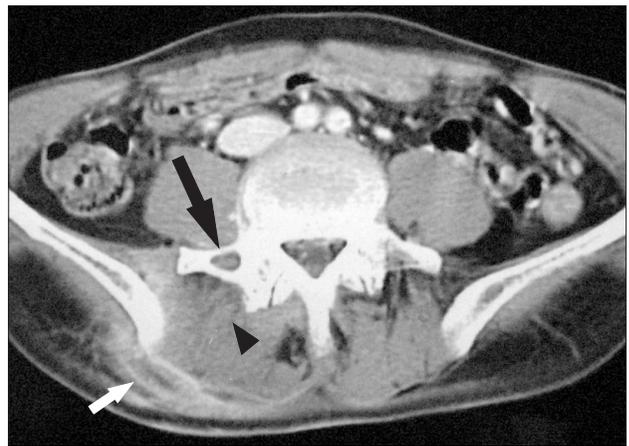


Fig. 3. Axial enhanced computed tomography (CT) reveals a space-occupying lesion that showed a rim enhancement pattern (white arrow) and swelling and signal change of the paraspinal muscle (arrow-head). CT also reveals a 2 × 1 cm sized osteolytic lesion (black arrow) located in the right transverse process of L5.

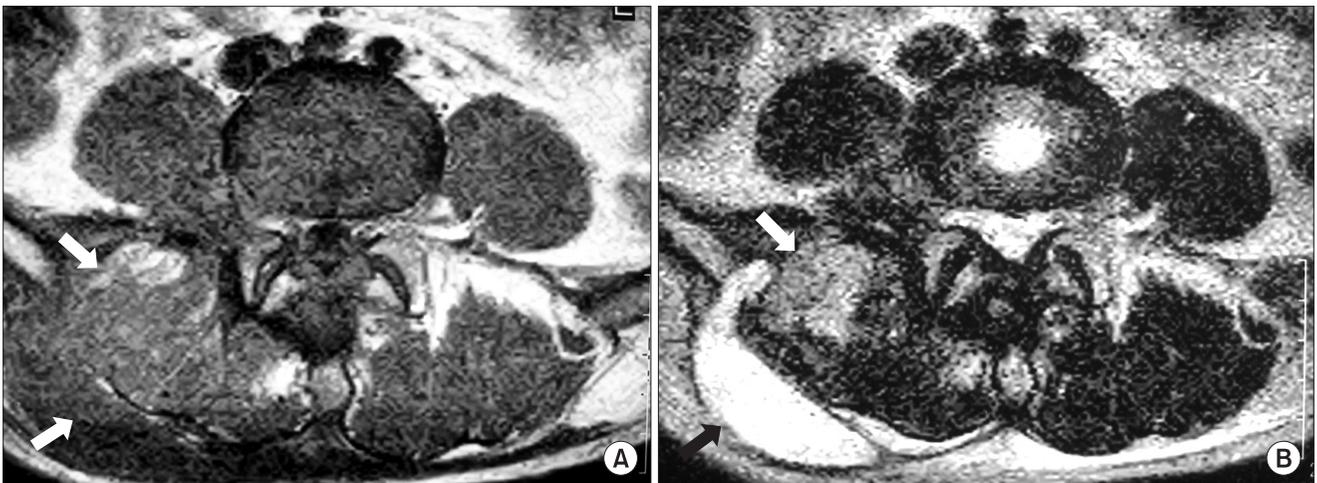


Fig. 2. (A) Axial T1-weighted magnetic resonance imaging (MRI) reveals an intermediate signal intensity lesion (white arrow). (B) Axial T2-weighted MRI reveals high signal intensity in the space-occupying lesion (black arrow), which was located behind the right paraspinal muscle, and intermediate signal intensity in the muscle (white arrow).



Fig. 4. The postoperative plain radiograph shows the complete resection of the right transverse process of L5 (arrow).

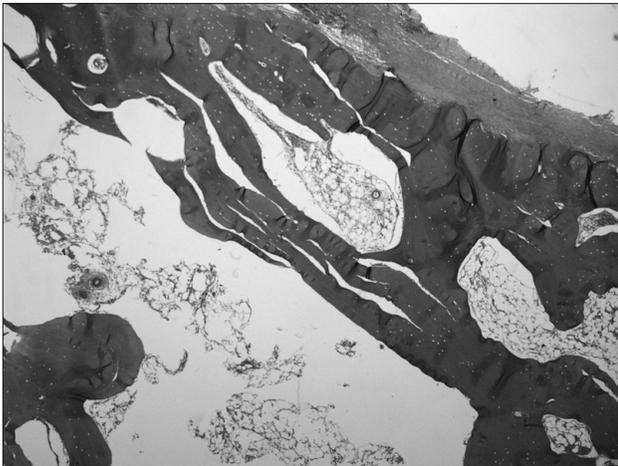


Fig. 5. Histologically, the dense lamellar bone spaces are infiltrated by chronic inflammatory cells (H&E, $\times 100$).

for the spread of a pelvic or genitourinary infection to the spine. Wiley and Trueta⁶⁾ suggested the arterial network as the most likely source of bacterial spread to the spine. The transverse process contains end-arterioles and sinusoidal vessels that are similar to but less vascularized than the metaphyseal area of the vertebral body. An infecting organism enters the end-arterioles via the dorsal spinal plexus arising from a segmental artery. Once established, an infection within the transverse process can extend through the cortex into the paraspinal soft tissue.^{2,6)}

Invasive diagnostic and treatment efforts have been associated with iatrogenic infections of the spine. In addition, direct bacterial inoculations of the spine via a penetrating trauma such as stab and gunshot wounds are other sources of infection. In our case, there was no evidence of any pelvic or genitourinary infections, nor was there any

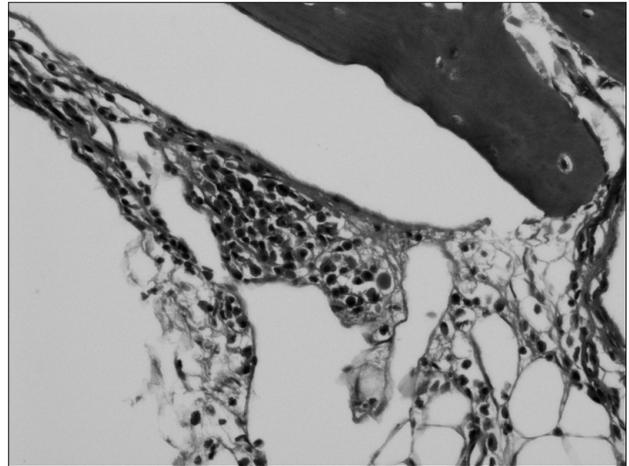


Fig. 6. The high power microscopic study shows a mixture of inflammatory cells that are mainly lymphocytes in the lamellar bone spaces (H&E, $\times 400$).

evidence of an iatrogenic infection and direct inoculation. Therefore, the arterial hematogenous route is the most likely mechanism of the osteomyelitis in this case.

Staphylococcus aureus is clearly the main organism in hematogenous vertebral osteomyelitis, and it accounts for more than 55% of all the reported cases. A polymicrobial cause is quite unusual and it accounts for less than 2.5% of the cases.⁷⁾ In our case, 2 staphylococcus species, *Staphylococcus aureus* and coagulase-negative staphylococcus were identified. Our patient denied a history of acupuncture or local injections. But one may consider the possibility of iatrogenic infection.

Imaging studies are important for the diagnosis, the treatment and its planning, and for monitoring the treatment outcome of a spinal infection. The sensitivity of CT is high, but it lacks specificity. MRI is sensitive, specific and accurate and it is the method of choice for examining spinal infections.⁸⁾ In our case the osteolytic lesion was not detected by MRI, while an accurate diagnosis was made by CT. Therefore, we believe that CT is more sensitive to the subtle changes in the bone than MRI, and MRI combined with CT is a more useful modality for evaluating transverse process osteomyelitis.

The treatment modality is determined by the extent of the infection and the degree of neurological compromise. In our case, the paraspinal abscess was surgically drained, and the primary osteolytic lesion was resected in order to eradicate the infection focus. Although 6 to 8 weeks of antimicrobial therapy is the generally accepted regimen, a recent report recommended 4 weeks of an appropriate high-dose, parenteral antimicrobial therapy,⁷⁾ and this was adopted in our case.

King and Mayo⁹⁾ reported that subacute osteomyelitis of the spine is a subtype of subacute hematogenous osteomyelitis. However, they did not introduce transverse process osteomyelitis.

Transverse process osteomyelitis is extremely rare, and this can result in a misdiagnosis or a delayed diagnosis. This report shows the transverse process is a possible location of vertebral osteomyelitis, and transverse process osteomyelitis with paraspinal abscess is an unusual cause

of back pain. Therefore, transverse process osteomyelitis should be borne in mind as part of the differential diagnosis when conducting an examination of back pain.

CONFLICT OF INTEREST

No potential conflict of interest relevant to this article was reported.

REFERENCES

1. Calderone RR, Larsen JM. Overview and classification of spinal infections. *Orthop Clin North Am.* 1996;27(1):1-8.
2. Buoncristiani AM, McCullen G, Shin AY, Bathgate B, Akbarnia BA. An unusual cause of low back pain: osteomyelitis of the spinous process. *Spine (Phila Pa 1976).* 1998;23(7):839-41.
3. Ehara S, Khurana JS, Kattapuram SV. Pyogenic vertebral osteomyelitis of the posterior elements. *Skeletal Radiol.* 1989;18(3):175-8.
4. Garcia A Jr, Grantham SA. Hematogenous pyogenic vertebral osteomyelitis. *J Bone Joint Surg Am.* 1960;42(3):429-36.
5. Batson OV. The vertebral vein system: Caldwell lecture, 1956. *Am J Roentgenol Radium Ther Nucl Med.* 1957;78(2):195-212.
6. Wiley AM, Trueta J. The vascular anatomy of the spine and its relationship to pyogenic vertebral osteomyelitis. *J Bone Joint Surg Br.* 1959;41(4):796-809.
7. Sapico FL. Microbiology and antimicrobial therapy of spinal infections. *Orthop Clin North Am.* 1996;27(1):9-13.
8. Tali ET. Spinal infections. *Eur J Radiol.* 2004;50(2):120-33.
9. King DM, Mayo KM. Subacute haematogenous osteomyelitis. *J Bone Joint Surg Br.* 1969;51(3):458-63.