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3.1 Clinical Presentation

Sacral tumors are generally diagnosed late and can present as large, advanced neoplastic masses because of mild initial symptoms. The clinical pattern depends on the anatomic location of the lesion within the sacrum, its extension, and whether it compresses or invades neighboring structures [1]. The pain may initially be nonspecific and as clinical examination is usually poor, these tumors may remain clinically silent for long periods of time. The most common initial symptom of a sacral tumor is local pain due to its mass effect and compression. Occasionally smaller lesions could become symptomatic secondary to involvement of critical structures, such as nerves or ureters, or because of pathologic fractures. Generally, however, these tumors remain asymptomatic until they are quite large, and lower sacral tumors can grow large enough for their anterior portion to be palpated during a rectal examination [1–3]. While lateral extension of sacral tumors across the sacroiliac joints causes local pain at the joint, invasion of the origin of the gluteus maximus and piriformis muscles leads to local pain and subsequently decreases hip extension and external rotation strength [3–7].

Subsequently, as nerve roots become increasingly compressed or infiltrated by tumor, multiradicular sensory deficits develop and can include radicular pain radiating uni- or bilaterally into the buttocks, posterior thigh or leg, external genitalia, and/or perineum (Fig. 3.1).

As this continues to progress, motor deficits, and eventually, bladder, bowel, and/or sexual dysfunction from anterior extension of the tumor into the presacral space can be noted [1].

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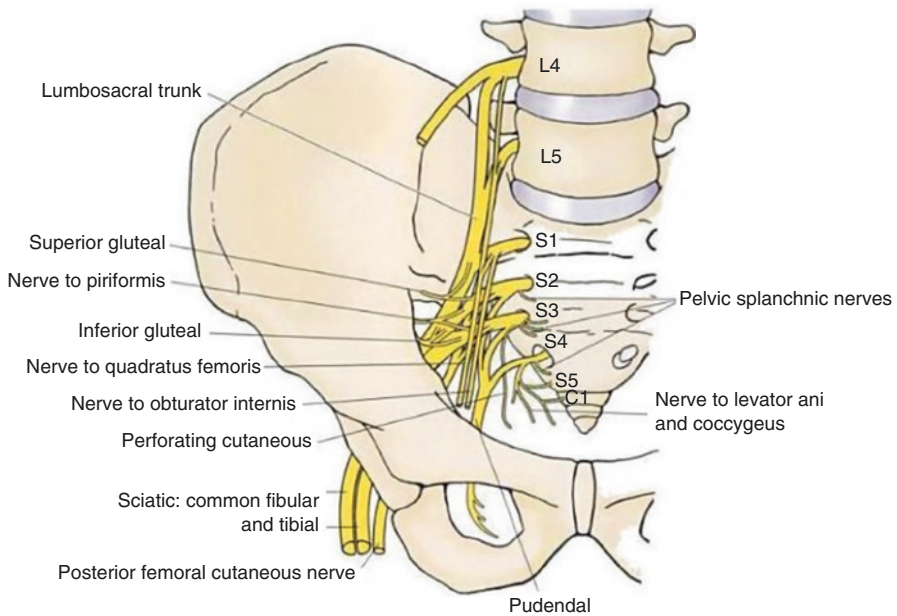


Fig. 3.1 Sacral nerve roots (2016, June). Retrieved July, 2016, from <http://wiki.ahuman.org/index.php/HumanNervesSpinalRoots>

Involvement of lumbosacral nerve roots in sacral lesions leads to certain specific deficits. A lesion involving the L-5 nerve root, commonly in its L5-S1 foraminal or extraforaminal course, may cause radicular pain and hypesthesias in the lateral thigh and calf as well as dorsum of the foot to the great toe [1, 2, 4]. Motor weakness of the L-5 nerve root may result in weakened ankle dorsiflexion, great toe extension, knee flexion, and hip abduction. The straight-leg raise test, or Lasegue's sign, which involves raising the patient's leg with a straight knee while the patient is supine, would result in sciatic pain and render a positive result. A lesion involving the S-1 nerve root, in its canalicular, S1-2 foraminal or extraforaminal course, typically causes radicular pain and hypesthesias in the posterior thigh and calf as well as at the lateral and plantar face of the foot and the small toe. A motor deficit due to an S-1 lesion may result in weakened ankle plantar flexion, knee flexion, and hip extension. A unilateral lesion to the S2 or S3 nerve root usually leads to mild or moderate bladder, bowel, and/or sexual dysfunction [8, 9]. A bilateral lesion of the S2 or S3 roots almost always results in complete bladder, bowel, and sexual dysfunction, and although a unilateral lesion at the same nerve root may cause symptoms, they are generally more nonspecific. However, unilateral or even bilateral lesions of the S4 and/or S5 roots do not result in autonomic dysfunction, although anatomical work has shown some S4 and S5 root contribution to bladder and bowel function [10]. Performance of a thorough physical exam in such patients is critical and can significantly aid in diagnosis and ancillary testing.

Conflict-of-Interest Statement No benefits have been or will be received from a commercial party related directed or indirectly to the subject matter of this article.

References

1. Payer M. Neurological manifestation of sacral tumors. *Neurosurg Focus*. 2003;15(2):E1.
2. Deutsch H, Mummaneni PV, Haid RW, Rodts GE, Ondra SL. Benign sacral tumors. *Neurosurg Focus*. 2003;15(2):E14.
3. Chandawarkar RY. Sacrococcygeal chordoma: review of 50 consecutive patients. *World J Surg*. 1996;20(6):717–9.
4. Cheng EY, Ozerdemoglu RA, Transfeldt EE, Thompson Jr RC. Lumbosacral chordoma. Prognostic factors and treatment. *Spine*. 1999;24(16):1639–45.
5. Yonemoto T, Tatezaki S, Takenouchi T, Ishii T, Satoh T, Moriya H. The surgical management of sacrococcygeal chordoma. *Cancer*. 1999;85(4):878–83.
6. Lin PP, Guzel VB, Moura MF, et al. Long-term follow-up of patients with giant cell tumor of the sacrum treated with selective arterial embolization. *Cancer*. 2002;95(6):1317–25.
7. York JE, Kaczaraj A, Abi-Said D, et al. Sacral chordoma: 40-year experience at a major cancer center. *Neurosurgery*. 1999;44(1):74–80.
8. Althausen PL, Schneider PD, Bold RJ, Gupta MC, Goodnight Jr JE, Khatri VP. Multimodality management of a giant cell tumor arising in the proximal sacrum: case report. *Spine*. 2002;27(15):E361–5.
9. Unni KK. Dahlin's bone tumors: general aspects and data on 11,087 cases. 5th ed. Philadelphia: Lippincott-Raven; 1997.
10. Bergh P, Kindblom LG, Gunterberg B, et al. Prognostic factors in chordoma of the sacrum and mobile spine: a study of 39 patients. *Cancer*. 2000;88(9):2122–34.