

Ga-68 DOTA-NOC PET/CT for the Detection of Residual/Recurrence in a Rare Case of Sacral Spinal Canal Paraganglioma

Dear Editor,

A 40-year-old male patient of sacral paraganglioma, who underwent S1-S2 laminectomy with excision of the sacral subdural paraganglioma 2 years ago, presented with ache in the lower back since 3 months with no history of numbness or weakness in both his limbs. Magnetic resonance imaging (MRI) of his hip revealed a subcentimetric heterogeneous nodule in the T1-weighted (T1W) and T2-weighted (T2W) short tau inversion recovery (STIR) images in the sacral region. Ga-68 DOTA-NOC positron emission tomography/computed tomography (PET/CT) was performed to characterize the lesion. Ga-68 PET/CT images [Figure 1] revealed a tracer avid (the maximum standardized uptake value being 4.7) subcentimetric (8.0 mm × 8.0 mm) soft

tissue nodular lesion in the spinal canal at the level of the S1 vertebra. The fat planes with the adjacent nerve roots appeared to be lost and so, local radiotherapy was planned for the patient.

Spinal paraganglioma is a very rare tumor, accounting for around 0.07% of all paragangliomas and 3–4% of all spinal tumors.^[1] The common locations of spinal paraganglioma are the cauda equina and filum terminale regions at the end of spinal column below the 1st lumbar vertebra (L1 vertebra) level.^[2] The clinical presentation of spinal paragangliomas depends on the level of the affected spinal cord and the degree of compression. The sacral spinal paraganglioma commonly presents with back pain radiating to the lower limbs, sensory and motor deficits, and bowel-bladder incontinence.^[3] Spinal paraganglioma is difficult to distinguish radiologically from other spinal lesions.^[1] Other intradural lesions such as lipoma, schwannoma, meningioma, ependymoma, and dermoid tumor should be included in differential diagnosis, which shows the same MRI imaging feature as paraganglioma.^[4,5] Ga-68 DOTA-NOC PET/CT shows tracer avidity in neural crest tumors, such as paraganglioma, due to the expression of somatostatin receptors.^[6] Spinal paragangliomas rarely metastasize; however, in the literature, metastasis cases are documented.^[7] Surgery is a primary treatment modality and even after gross tumor excision, only 4% shows local recurrence.^[8] Radiotherapy is reserved for surgically

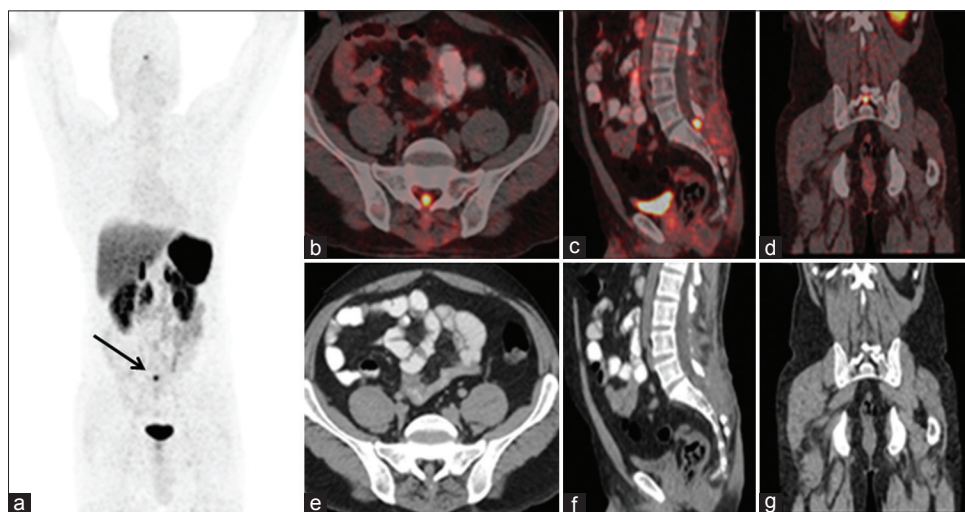


Figure 1: Ga-68 PET/CT maximum intensity projection (MIP) image (a) revealed solitary tracer avid focus in the sacral region. PET/CT and CT transaxial, sagittal, and coronal images (b-g) revealed a tracer avid (the maximum standardized uptake value being 4.7) subcentimetric sized (8.0 mm × 8.0 mm) soft tissue nodular lesion in the spinal canal at the level of S1 vertebra. The fat plane between the lesion and the nerve root, and the filum terminale appears to be nonpreserved

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nonassessable lesion(s) or recurrence; however, resistance to that treatment is also documented.^[9] DOTA-NOC imaging is also able to facilitate curative treatment by the identification of surgically nonresectable disease or metastatic spread of the disease.^[10] In the present case, DOTA-NOC scan showed tracer avidity in spinal canal paraganglioma, which is surgically not assessable due to the proximity of the nerve roots and can be considered for radiotherapy or Lu-177 DOTA-NOC therapy. To the best of my knowledge, this is the first case that reported somatostatin receptor expression in sacral spinal paraganglioma.

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Conflicts of interest

The authors declare no conflicts of interest.

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