68Ga-DOTA-D Phe1-Tyr3-Octreotide (DOTATOC)-PET/CT in a Suspected Case of Recurrent Meningioma

Sir,

We present a case of 45-year-old female who presented with headache for last 1 month. She was a follow-up case of left frontal meningioma that was operated 4 years back. CEMRI of the brain suggested post-surgical changes with no obvious enhancing residual/recurrent lesion. The patient was then referred for 68 Ga DOTANOC PET/CT to rule out recurrent disease. PET/CT revealed a focus of somatostatin receptor expressing lesion in the left frontal region at the post-operative site suggesting recurrent disease.

Meningioma is the most common non-glial brain tumor. It arises from the cap cells of the arachnoid membrane. However, 90% of meningiomas are benign. Surgery often with adjuvant radiotherapy is the usual treatment.^[1] MRI is the imaging method of choice for diagnosis and radiotherapy planning. Somatostatin receptors (SSTR) are present in normal leptomeninges^[2] and have been shown to be over expressed in meningiomas.^[3] The major receptor subtype over expressed is SSTR 2.^[4] This fact has been exploited for SSTR scintigraphy (SRS) and more recently for 68Ga-DOTA-peptide PET/CT in meningioma. Even after complete removal, meningiomas tend to recur in 10% to 32% of the cases within 10 years.^[5,6] MRI fails to differentiate between post-therapy radiation necrosis and recurrent disease in majority of the cases. Recurrent brain tumors are typically characterized by intravenous contrast enhancement, mass effect, and associated vasogenic edema. However, treatment necrosis also presents with similar characteristics, making it difficult to reliably distinguish from tumor recurrence. Because of the high tumor to background ratio and expression of the somatostatin receptor by the tumor, 68Ga-DOTA-peptide PET/CT helps in differentiation between the post-operative scar and recurrent disease in meningioma. It also helps in selection of patients for somatostatin-based analog-based therapies.

Financial support and sponsorship

Nil

Conflicts of interest

There are no conflicts of interest

Sachin Jain, Ashwani Gupta, Anurag Jain¹

Department of Nuclear Medicine, ¹Department of Radiology, Action Cancer Hospital, Paschim Vihar, New Delhi, India

Address for correspondence:

Dr. Sachin Jain, Department of Nuclear Medicine and PET/CT Action Cancer Hospital, A4, Paschim VIhar, New Delhi, India E-mail: sachinnpatni@gmail.com

References

- Stafford SL, Perry A, Suman VJ, Meyer FB, Scheithauer BW, Lohse CM. Primarily resected meningiomas: outcome and prognostic factors in 581 Mayo Clinic patients, 1978 through 1988. Mayo Clin Proc 1998;73:936-42.
- 2. Reubi JC, Maurer R, Lamberts SWJ. Somatostatin binding sites in human leptomeninx. Neurosci Lett 1986;70:183-6.
- Reubi JC, Maurer R, Klijn JGM, Stefanko SZ, Foekens JA, Blaauw G. High incidence of somatostatin receptors in human meningiomas: biochemical characterization. J Clin Endocrinol Metab 1986;63:433-8.
- Barresi V, Alafaci C, Salpietro F, Tuccari G. Sstr2A immunohistochemical expression in human meningiomas: Is there a correlation with the histological grade, proliferation or microvessel density? Oncol Rep 2008;20:485-92.
- Adegbite AB, Khan MI, Paine KWE, Tan LK. The recurrence of intracranial meningiomas after surgical treatment. J Neurosurg 1983;58:51-6.
- Mirimanoff RO, Dosoretz DE, Linggood RM, Ojemann RG, Martuza RL. Meningioma: analysis of recurrence and progression following neurosurgical resection. J Neurosurg 1985;62:18-24.

This is an open access article distributed under the terms of the Creative Commons Attribution-Non Commercial-Share Alike 3.0 License, which allows others to remix, tweak, and build upon the work non-commercially, as long as the author is credited and the new creations are licensed under the identical terms.

Access this article online
Website: www.indjsp.org
DOI: 10.4103/0972-3919.202241
Quick Response Code:

How to cite this article: Jain S, Gupta A, Jain A. 68Ga-DOTA-D Phe1-Tyr3-octreotide (DOTATOC)-PET/CT in a suspected case of recurrent meningioma. Indian J Nucl Med 2017;32:164.