

Role of ¹⁸F-Fluorodeoxyglucose Positron Emission Tomography–Contrast-Enhanced Computed Tomography in Detection of Early Recurrence with Peritoneal Disease in a Case of Adrenocortical Carcinoma

Abstract

Adrenocortical carcinoma (ACC) is an uncommon and highly aggressive malignancy with a poor prognosis. Common sites of metastasis are lung, liver, and lymph nodes. We present a case of ACC in a 54-year-old female with an early disease recurrence of unusual hypervascular peritoneal metastatic abdominal-pelvic deposits detected on ¹⁸F-fluorodeoxyglucose positron emission tomography–contrast-enhanced computed tomography scan.

Keywords: ¹⁸F-fluorodeoxyglucose positron emission tomography–computed tomography, adrenocortical carcinoma, peritoneal metastasis

A 54-year-old female presented with complaints of sudden-onset hypertension, hirsutism, and hot flushes. Initial diagnostic ultrasound and computed tomography (CT) workup detected a left adrenal mass lesion suggesting primary neoplasm. Staging ¹⁸F-fluorodeoxyglucose positron emission tomography-CT (¹⁸F-FDG PET-CT) scan [Figure 1] coronal PET, fused coronal PET-CT [Figure 1a and b], and corresponding fused transaxial images [Figure 1c-h] revealed an FDG-avid (SUVmax: 18.3) large heterogeneously enhancing lobulated left adrenal lesion with areas of necrosis. The lesion is abutting the left kidney and spleen, with tumor thrombus infiltrating the left suprarenal, left renal, left ovarian vein and adjacent inferior vena cava. The patient underwent left adrenalectomy, left nephrectomy, and caval thrombectomy. Histopathology revealed adrenocortical carcinoma (ACC), an uncommon epithelial malignancy of adrenal cortical cells with an unfavorable prognosis. ACC has a female predilection, with the left adrenal being commonly involved.^[1,2] The patient was symptom free until just 5 months post surgery, and she presented with left flank pain. Triple-phase contrast-enhanced CT (CECT)

abdomen-pelvis scan revealed multiple new-onset arterially hyperenhancing small nodular peritoneal deposits scattered in the abdomen-pelvis [Figure 2c, e, g, and i] which appeared isodense to abdominal aorta on subsequent phases. The abovementioned lesions demonstrated low-grade FDG avidity with SUVmax 3.8 as seen on FDG PET-CT scan coronal PET, fused PET-CT [Figure 2a and b], and corresponding fused transaxial images [Figure 2d, f, h, and j]. These scan findings appeared very unusual for ACC metastasis and demonstrated an appearance of splenosis (although the spleen was intact post surgery). Common sites of ACC metastasis include lung, liver, and lymph nodes, with peritoneal involvement being extremely rare.^[3,4] Studies suggest that FDG PET-CT has a greater positive likelihood ratio than CT to identify liver and abdominal ACC recurrences and thus change in management strategy.^[5-7] ACC being an aggressive neoplasm, metastatic disease is known to demonstrate higher FDG avidity, unlike that seen in our case. The patient underwent exploratory laparotomy with excision of multiple peritoneal nodules, and histopathology revealed metastatic ACC deposits. From this case, we learn

**Debdip Roy,
Melvika Pereira,
Divya Shivdasani,
Natasha Singh,
Shreya Dang,
Rachita Rungta**

Department of Nuclear Medicine and Molecular Imaging, P. D. Hinduja National Hospital and MRC, Mumbai, Maharashtra, India

Address for correspondence:

Dr. Debdip Roy,
Department of Nuclear Medicine and Molecular Imaging, P. D. Hinduja National Hospital and MRC, Mumbai, Maharashtra, India.
E-mail: debdiproy9@gmail.com

Received: 09-02-2022

Revised: 23-03-2022

Accepted: 04-04-2022

Published: 02-12-2022

Access this article online

Website: www.ijnm.in

DOI: 10.4103/ijnm.ijnm_33_22

Quick Response Code:



This is an open access journal, and articles are distributed under the terms of the Creative Commons Attribution-NonCommercial-ShareAlike 4.0 License, which allows others to remix, tweak, and build upon the work non-commercially, as long as appropriate credit is given and the new creations are licensed under the identical terms.

For reprints contact: WKHLRPMedknow_reprints@wolterskluwer.com

How to cite this article: Roy D, Pereira M, Shivdasani D, Singh N, Dang S, Rungta R. Role of ¹⁸F-fluorodeoxyglucose positron emission tomography–contrast-enhanced computed tomography in detection of early recurrence with peritoneal disease in a case of adrenocortical carcinoma. Indian J Nucl Med 2022;37:389-91.

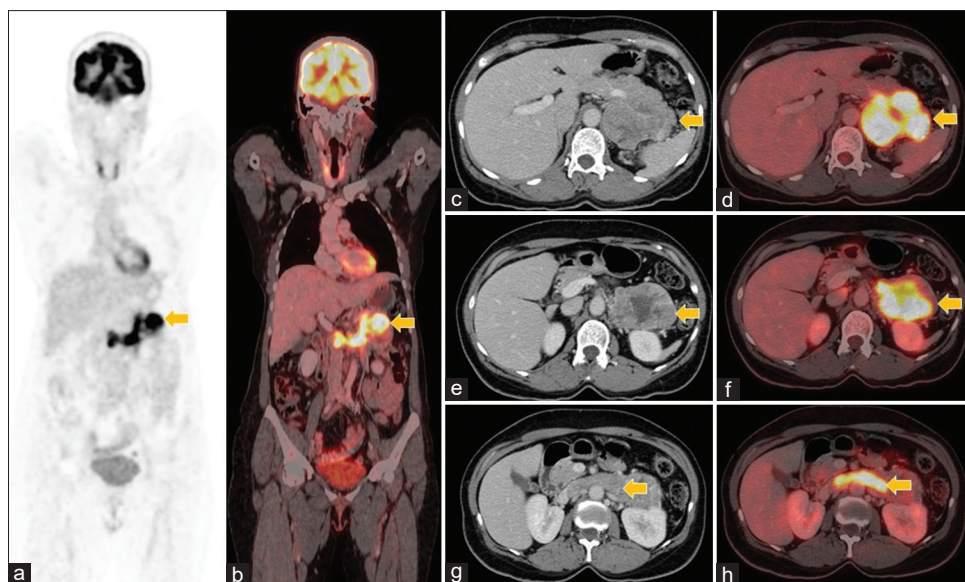


Figure 1: Staging ^{18}F -fluorodeoxyglucose positron emission tomography-computed tomography scan coronal positron emission tomography (a), fused coronal positron emission tomography-computed tomography (b), and corresponding transaxial images (c-h) showing fluorodeoxyglucose-avid large heterogeneously enhancing left adrenal lesion with tumor thrombus infiltrating the left renal vein and adjacent inferior vena cava

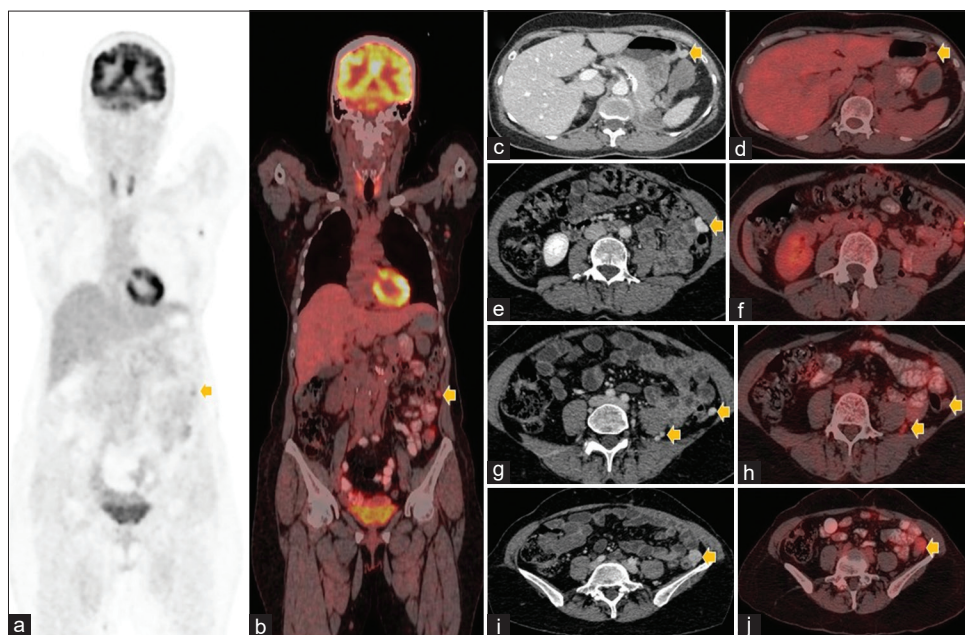


Figure 2: ^{18}F -fluorodeoxyglucose positron emission tomography-computed tomography scan coronal positron emission tomography (a), fused coronal positron emission tomography-computed tomography (b), contrast-enhanced computed tomography abdomen-pelvis (c, e, g, and i) and corresponding fused transaxial positron emission tomography-computed tomography images (d, f, h, and j) showing arterially hyperenhancing small nodular peritoneal deposits with low fluorodeoxyglucose avidity

that ACC metastatic recurrence can involve rarely only the peritoneum quite early post surgery, appearing as hypervascular nodules and presenting low FDG uptake. ^{18}F -FDG PET-CECT (triple-phase protocol) scan would be most appropriate to confidently detect the lesions.

Declaration of patient consent

The authors certify that they have obtained all appropriate patient consent forms. In the form, the patient has given her consent for her images and other clinical information

to be reported in the journal. The patient understands that her name and initials will not be published and due efforts will be made to conceal identity, but anonymity cannot be guaranteed.

Financial support and sponsorship

Nil.

Conflicts of interest

There are no conflicts of interest.

References

1. Ng L, Libertino JM. Adrenocortical carcinoma: Diagnosis, evaluation and treatment. *J Urol* 2003;169:5-11.
2. Takeuchi S, Balachandran A, Habra MA, Phan AT, Bassett RL Jr., Macapinlac HA, *et al.* Impact of ¹⁸F-FDG PET/CT on the management of adrenocortical carcinoma: Analysis of 106 patients. *Eur J Nucl Med Mol Imaging* 2014;41:2066-73.
3. Owen DH, Patel S, Wei L, Phay JE, Shirley LA, Kirschner LS, *et al.* Metastatic adrenocortical carcinoma: A single institutional experience. *Horm Cancer* 2019;10:161-7.
4. Kumar T, Nigam JS, Sharma S, Kumari M, Pandey J. Uncommon metastasizing site of adrenocortical carcinoma. *Cureus* 2021;13:e15267.
5. Ardito A, Massaglia C, Pelosi E, Zaggia B, Basile V, Brambilla R, *et al.* ¹⁸F-FDG PET/CT in the post-operative monitoring of patients with adrenocortical carcinoma. *Eur J Endocrinol* 2015;173:749-56.
6. Leboulleux S, Dromain C, Bonniaud G, Aupérin A, Caillou B, Lumbroso J, *et al.* Diagnostic and prognostic value of ¹⁸-fluorodeoxyglucose positron emission tomography in adrenocortical carcinoma: A prospective comparison with computed tomography. *J Clin Endocrinol Metab* 2006;91:920-5.
7. Mackie GC, Shulkin BL, Ribeiro RC, Worden FP, Gauger PG, Mody RJ, *et al.* Use of [¹⁸F] fluorodeoxyglucose positron emission tomography in evaluating locally recurrent and metastatic adrenocortical carcinoma. *J Clin Endocrinol Metab* 2006;91:2665-71.