**18F-Fluciclovine–Avid Axillary Lymph Nodes After COVID-19 Vaccination on PET/CT for Suspected Recurrence of Prostate Cancer**

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Abnormally increased 18F-FDG avidity of axillary lymph nodes has become a frequent diagnostic dilemma on PET/CT in the current climate of global vaccinations directed against severe acute respiratory syndrome coronavirus 2. This avidity is due to the inflammatory response evoked by vaccines and the nonspecific nature of 18F-FDG uptake, which is increased in both malignant and inflammatory processes. Similarly, 18F-fluciclovine, an amino acid analog indicated for the assessment of biochemical recurrence of prostate cancer, may also demonstrate nonspecific inflammatory uptake. We report a case of 18F-fluciclovine PET/CT obtained for concern about prostate cancer. In this case, isolated avid lymph nodes were seen in the left axilla. A screening questionnaire revealed that the patient had recently received the second dose of the Pfizer-BioNTech coronavirus disease 2019 vaccine in his left shoulder, and hence, the uptake was determined to be reactive.

**Key Words:** PET/CT; COVID-19 vaccine; 18F-fluciclovine; axillary lymph nodes; inflammation; prostate cancer

**CASE REPORT**

A 65-y-old man underwent surgery for prostate cancer. Subsequent pathologic evaluation revealed a Gleason score of 7 (3 + 4) with evidence of perineural invasion and 1 of 5 local lymph nodes positive for spread. There was no seminal vesicle invasion or extracapsular extension, and the surgical margins were negative. The overall stage was IVA (pT2, N1, M0). At 4 mo after surgery, there was a persistent detectable prostate-specific antigen level of 1.0 ng/mL, and 18F-fluciclovine (Axumin; Blue Earth Diagnostics) PET/CT was performed for further assessment. The examination revealed 4 foci of avidity (SUVmax, 3.2) in the left axilla localizing to mildly enlarged but morphologically normal lymph nodes (Fig. 1). The remainder of the uptake was physiologic, with no other sites of pathologic radiotracer avidity. A review of the intake questionnaire showed that the patient had received the Pfizer-BioNTech COVID-19 vaccine in his left shoulder 17 d previously, and hence, the uptake was determined to be reactive. Subsequent clinical follow-up revealed an undetectable prostate-specific antigen level at 6 mo and again at 9 mo, confirming the benign nature of the uptake and the absence of residual malignancy.

**DISCUSSION**

Although 18F-FDG is by far the most frequently used radiotracer in PET/CT, several additional radiotracers are in clinical use, including 18F-fluciclovine for imaging in men with suspected prostate cancer recurrence. 18F-fluciclovine is an amino acid analog and is taken up by prostate cancer (4). However, like 18F-FDG, 18F-fluciclovine can show increased uptake in inflammatory processes due to uptake by white blood cells. Given this fact, false-positive uptake has been reported in nonmalignant entities such as pneumonia, lymphadenitis, and ring worm infection (4,5). As with 18F-FDG, 18F-fluciclovine uptake by inflammatory processes tends to be mild compared with that by malignancy and most commonly has an intensity less than that in normal bone marrow (as in this case, with the L3 vertebra...
having an SUV$_{\text{mean}}$ of 3.5). Hence, it is to be expected that, as with $^{18}$F-FDG, low-grade avidity by axillary nodes will be a potential finding on $^{18}$F-fluciclovine PET/CT after COVID-19 vaccination.

Recently, several prostate-specific membrane antigen PET tracers have received approval for use in the United States as an additional means for detecting prostate cancer. Despite their excellent performance, these, too, can demonstrate nonspecific uptake by inflammatory diseases such as pneumonia, and hence, it is likely only a matter of time before a case of COVID-19 vaccination–related uptake in the axillary lymph nodes is reported with these agents (6).

**CONCLUSION**

The case underlines the importance of correlating examination findings with disease pathophysiology, radiotracer mechanism of action, and clinical history to optimize the accuracy of PET/CT interpretation.

**DISCLOSURE**

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

**REFERENCES**