

COVID-19–Related Lung Parenchymal Uptake on ^{18}F -PSMA-1007 PET/CT

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Abstract: A 70-year-old man with newly diagnosed prostate cancer underwent ^{18}F -PSMA-1007 PET/CT for staging. PSMA-avid primary prostatic malignancy was identified. Incidental intense patchy peripheral lung uptake was also noted. The patient tested positive for COVID-19 infection.

Key Words: COVID-19, lung uptake, ^{18}F -PSMA-1007 PET/CT

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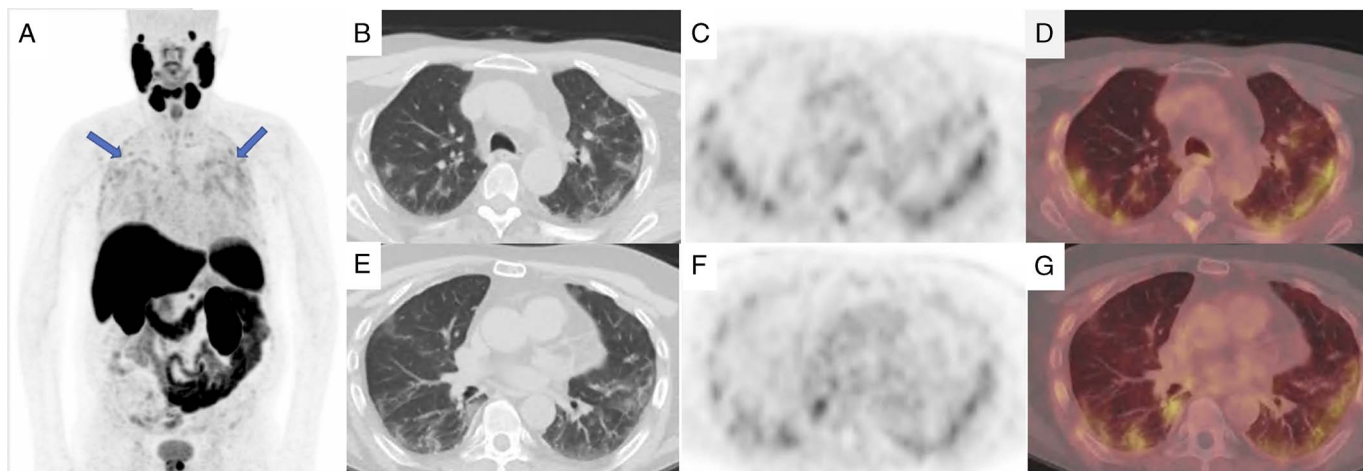


FIGURE 1. A 70-year-old man with prostate cancer. ^{18}F -PSMA-1007 PET/CT. **A**, MIP demonstrates intense focal uptake in the prostate consistent with the known malignancy. Moderate ill-defined uptake in the chest (arrows) was also seen. Axial CT, ^{18}F -PSMA-1007 PET, and fused PET/CT of the chest at 2 different levels (**B–D**) and (**E–G**) demonstrate uptake associated with patchy peripheral/ subpleural opacification. This is associated with moderate uptake on the PET component. The low-dose CT features were in keeping with COVID-19 infection, with a classic pattern of COVID-19 pneumonia bilateral, peripheral ground glass opacities.^{1–3} The presence and pattern of lung parenchymal change depends on the time of imaging during the course of illness. Ground glass opacities commonly develop between days 0 and 4, peaking at days 6 to 13 and may persist beyond 24 days.^{2,4} Sixty-one percent of persistent long-term changes beyond 24 days are ground glass opacities. Also, associated uptake depends on the grade of inflammation, which is associated with variable vascular permeability.⁵ In keeping with this, the patient gave a history of a positive real time-polymerase chain reaction test result for COVID-19 4 weeks before the scan. Inflammatory and infectious lung processes have been documented to demonstrate ^{18}F -PSMA uptake.^{5,6} Similar avid COVID-19-related findings on PET/CT were described but in asymptomatic patients with other tracers including ^{18}F -FDG, ^{68}Ga -DOTANOC, and ^{18}F -flourocholine.^{7–9} This was also described with ^{68}Ga -PSMA, which has the same mechanism of uptake as all PSMA-based PET tracers, including ^{18}F -PSMA-1007.¹⁰ PSMA stands for “prostate-specific membrane antigen,” which could be argued as a misnomer because these are not exclusively expressed on prostate cells and can be found in other tissues and conditions, such as inflammation/infection. There are limited data available explaining the exact mechanism of uptake at sites of inflammation, but it is thought to possibly also be as a result of neovascularization and increased regional blood flow/vascular permeability, delivering more PSMA ligand to the site of inflammation/infection.⁵ This example of nonmalignant PSMA lung uptake highlights a cause, which may be increasingly seen in the context of the current COVID-19 pandemic and may persist even after the initial illness.