

# SNMMI Comment on the 2016 Society of Surgical Oncology “Choosing Wisely” Recommendation on the Use of PET/CT in Colorectal Cancer

Katherine Zukotynski<sup>1</sup>, Hossein Jadvar<sup>2</sup>, Thomas Hope<sup>3</sup>, Rathan M. Subramaniam<sup>4</sup>, Katherine Van Loon<sup>5</sup>, Madhulika Varma<sup>6</sup>, and Ryan D. Niederkoher<sup>7</sup> for the PET Center of Excellence

<sup>1</sup>Departments of Radiology and Medicine, McMaster University, Hamilton, Ontario, Canada; <sup>2</sup>Department of Radiology, University of Southern California, Los Angeles, California; <sup>3</sup>Department of Radiology, UCSF, San Francisco, California; <sup>4</sup>Departments of Radiology and Clinical Sciences, University of Texas Southwestern Medical Center, Dallas, Texas; <sup>5</sup>Department of Medicine, UCSF, San Francisco, California; <sup>6</sup>Department of Surgery, UCSF, San Francisco, California; and <sup>7</sup>Department of Nuclear Medicine, Kaiser Permanente, Santa Clara, California

**T**here has been recent debate about where PET/CT with <sup>18</sup>F-FDG fits into the evaluation of patients with colorectal cancer. The PET Center of Excellence of the Society of Nuclear Medicine and Molecular Imaging (SNMMI) would like to comment on the “Choosing Wisely” list recently published by the Society of Surgical Oncology, item 4 of which reads as follows: “Don’t perform routine PET-CT in the initial staging of localized colon or rectal cancer or as part of routine surveillance for patients who have been curatively treated for colon or rectal cancer. A CT of the chest, abdomen and pelvis with IV and PO contrast provides excellent staging and standard PET imaging does not significantly improve diagnostic accuracy or outcomes as part of the initial workup or surveillance testing. Use of PET does not eliminate the need for recommended staging CT with IV and PO contrast but does increase costs.” We are concerned about this statement because it is broad and does not allow for consideration of the utility of PET/CT on a case-by-case basis. Further, in an era in which insurance companies derive unwavering policies from consensus statements such as this one, the effect of such a broad-sweeping statement may be that providers are denied the opportunity to exercise clinical judgment on the most appropriate imaging modality—“choosing right.” Indeed, there are several instances in which PET/CT is clinically valuable both for initial staging and for subsequent treatment planning in patients with colorectal cancer.

The current colon cancer guidelines of the National Comprehensive Cancer Network (1) state that PET/CT should be considered at the time of initial staging in two specific instances. The first is when there is a need to characterize equivocal or indeterminate findings on staging contrast-enhanced (CE) CT or when a patient has contraindications to intravenous contrast for CT (e.g., a contrast allergy or impaired renal function), and the second is when there is a need to evaluate patients with potentially curable stage M1 disease (with intent to exclude other sites of occult disease that might potentially

render the disease unresectable). The guidelines also highlight the usefulness of PET/CT for restaging in the setting of a serially elevated carcinoembryonic antigen level, negative results on conventional imaging, and potentially resectable metachronous metastases documented by CT, MRI, or biopsy (2).

PET/CT performed at the time of initial staging changes the treatment strategy in a substantial number of patients with colorectal or anal cancer. A prospective study by Ozis et al. on 97 patients with rectal cancer who underwent CE CT and PET/CT at initial staging found that PET/CT detected more sites of disease than CE CT alone (particularly distant metastases) and had an impact on treatment strategy in 14% of the patients (3). Petersen et al., in a retrospective review of 67 patients with colorectal cancer who underwent CE CT and PET/CT at initial staging, concluded that compared with CE CT alone, PET/CT changed the management plan in 30% of the patients (4). Jones et al., in a systematic review of 12 studies comparing PET or PET/CT with conventional imaging during the initial staging of anal carcinoma, reported that the PET findings altered the stage in 41% of the patients (5). Further, Shi et al. and Byun et al. reported on retrospective series that found PET/CT could be prognostic of patient survival (6,7).

PET/CT can reliably assess therapeutic response, with particularly compelling data in patients undergoing treatment for locally advanced rectal cancer (7). Among recently published papers, Calvo et al. prospectively evaluated PET/CT in 38 patients with rectal carcinoma before and after adjuvant therapy and found that a metabolic response was associated with significantly higher survival (9). Leccisotti et al. prospectively studied PET/CT in 126 patients with rectal cancer before and after neoadjuvant chemoradiation, and their findings suggested that early assessment of response using PET/CT could predict an incomplete pathologic response, thus opening the door to earlier therapy modification if needed (10). Schneider et al. retrospectively studied 199 patients with rectal cancer at the time of restaging after neoadjuvant chemoradiation and found that the PET results brought about a change in clinical management in up to 32% of the patients (11).

PET/CT often detects recurrent colorectal cancer when conventional anatomic imaging (e.g., CT or MRI) does not. In particular, in a retrospective study by Choi et al. on 245 colorectal cancer patients who underwent CE CT and PET/CT as part of routine follow-up after resection with curative intent, PET/CT was found to detect more sites of recurrent disease than CE CT (12). Further,

Received Aug. 14, 2016; revision accepted Sep. 7, 2016.

For correspondence or reprints contact: Hossein Jadvar, Keck School of Medicine of USC, University of Southern California, 2250 Alcazar St., CSC 102, Los Angeles, CA 90033.

E-mail: jadvar@med.usc.edu

Published online Sep. 29, 2016.

COPYRIGHT © 2017 by the Society of Nuclear Medicine and Molecular Imaging.

DOI: 10.2967/jnumed.116.182584

Gade et al., Mittal et al., and Metser et al. found that in patients with previously resected colorectal cancer and clinically suspected recurrence, lesion detection was significantly higher with PET/CT (on the order of 15%–30%) than with CE CT (13–15).

As the current guidelines of the National Comprehensive Cancer Network—as well as a growing body of literature—suggest, PET/CT plays an important role in determining both the initial treatment strategy and the subsequent treatment strategy for certain patients with colorectal cancer. Studies using PET/CT to assess therapeutic response, particularly in rectal cancer, have had promising findings. Ultimately, we need to partner with our clinical (medical and surgical) colleagues to adopt a personalized approach for our patients and ensure appropriate, effective use of PET/CT in their care. In a manner of speaking, “routine” is, at least to some extent, becoming a word of the past and probably should not be used in our recommendations. We should be “choosing right” for each patient to provide optimal clinical care.

## DISCLOSURE

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

## REFERENCES

1. Five things physicians and patients should question. Society of Surgical Oncology website. <http://www.surgonc.org/docs/default-source/default-document-library/sso-five-things-physicians-and-patients-should-question-7-11-2016.pdf>. Published July 12, 2016. Accessed September 21, 2016.
2. Clinical practice guidelines in oncology (NCCN guidelines®): colon cancer—version 2.2016. National Comprehensive Cancer Network website. [https://www.nccn.org/professionals/physician\\_gls/pdf/colon.pdf](https://www.nccn.org/professionals/physician_gls/pdf/colon.pdf). Published November 24, 2015. Accessed September 21, 2016.
3. Ozis SE, Soydal C, Akyol C, et al. The role of <sup>18</sup>F-fluorodeoxyglucose positron emission tomography/computed tomography in the primary staging of rectal cancer. *World J Surg Oncol*. 2014;12:26–33.
4. Petersen RK, Hess S, Alavi A, Høilund-Carlsen PF. Clinical impact of FDG-PET/CT on colorectal cancer staging and treatment strategy. *Am J Nucl Med Mol Imaging*. 2014;4:471–482.
5. Jones M, Hruby G, Solomon M, Rutherford N, Martin J. The role of FDG-PET in the initial staging and response assessment of anal cancer: a systematic review and meta-analysis. *Ann Surg Oncol*. 2015;22:3574–3581.
6. Shi D, Cai G, Peng J, et al. The preoperative SUVmax for <sup>18</sup>F-FDG uptake predicts survival in patients with colorectal cancer. *BMC Cancer*. 2015;15:991–998.
7. Byun BH, Moon SM, Shin US, et al. Prognostic value of <sup>18</sup>F-FDG uptake by regional lymph nodes on pretreatment PET/CT in patients with resectable colorectal cancer. *Eur J Nucl Med Mol Imaging*. 2014;41:2203–2211.
8. Maffione AM, Marzola MC, Caprici C, Colletti PM, Rubello D. Value of <sup>18</sup>F-FDG PET for predicting response to neoadjuvant therapy in rectal cancer: systematic review and meta-analysis. *AJR*. 2015;204:1261–1268.
9. Calvo FA, Sole CV, de la Mata D, et al. <sup>18</sup>F-FDG PET/CT-based treatment response evaluation in locally advanced rectal cancer: a prospective validation of long-term outcomes. *Eur J Nucl Med Mol Imaging*. 2013;40:657–667.
10. Leccisotti L, Gambacorta MA, de Waure C, et al. The predictive value of <sup>18</sup>F-FDG PET/CT for assessing pathological response and survival in locally advanced rectal cancer after neoadjuvant radiochemotherapy. *Eur J Nucl Med Mol Imaging*. 2015;42:657–666.
11. Schneider DA, Akhurst TJ, Ngan SY, et al. Relative value of restaging MRI, CT, and FDG-PET scan after preoperative chemoradiation for rectal cancer. *Dis Colon Rectum*. 2016;59:179–186.
12. Choi EK, Yoo IR, Park HL, et al. Value of surveillance <sup>18</sup>F-FDG PET/CT in colorectal cancer: comparison with conventional imaging studies. *Nucl Med Mol Imaging*. 2012;46:189–195.
13. Gade M, Kubik M, Fisker RV, Thorlacius-Ussing O, Petersen LJ. Diagnostic value of <sup>18</sup>F-FDG PET/CT as first choice in the detection of recurrent colorectal cancer due to rising CEA. *Cancer Imaging*. 2015;15:11–18.
14. Mittal BR, Senthil R, Kashyap R, et al. <sup>18</sup>F-FDG PET-CT in evaluation of post-operative colorectal cancer patients with rising CEA level. *Nucl Med Commun*. 2011;32:789–793.
15. Metser U, You J, McSweeney S, Freeman M, Hendler A. Assessment of tumor recurrence in patients with colorectal cancer and elevated carcinoembryonic antigen level: FDG PET/CT versus contrast-enhanced 64-MDCT of the chest and abdomen. *AJR*. 2010;194:766–771.