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**Customer satisfaction:** Hicks offers perspective on results of surveys that seek the opinion of referring clinicians about oncologic <sup>18</sup>F-FDG PET/CT reporting and previews an article on this topic in this issue of *JNM*. . . . . *Page 1923*

**Perception of PET/CT misinterpretation:** Karantanis and colleagues detail results of a study designed to collect pertinent information from physicians referring patients for oncologic <sup>18</sup>F-FDG PET/CT, with surprising findings about clinicians' most frequent complaint. . . . . *Page 1925*

**DWI in whole-body PET/MR:** Grueneisen and colleagues assess the diagnostic benefit of diffusion-weighted imaging in an <sup>18</sup>F-FDG PET/MR protocol for whole-body staging of women with primary or recurrent malignancies of the pelvis. . . . . *Page 1930*

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**<sup>99m</sup>Tc-folate M2 macrophage plaque imaging:** Jager and colleagues explore the presence of activated macrophages in human atherosclerotic plaques with <sup>99m</sup>Tc-folate imaging and evaluate whether this technique can discriminate between M1- and M2-like macrophage phenotypes. . . . . *Page 1945*

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**Scintigraphic imaging of endocarditis:** Rouzet and colleagues compare the respective performances of <sup>18</sup>F-FDG PET and leukocyte scintigraphy for diagnosis of prosthetic valve endocarditis in 39 patients. . . . . *Page 1980*

**FACBC physiologic uptake:** Schuster and colleagues explore common physiologic uptake patterns, as well as incidental findings and variants, in patients who have undergone PET imaging with this synthetic amino acid analog. . . . . *Page 1986*

**<sup>11</sup>C-CURB dosimetry:** Boileau and colleagues investigate in humans the whole-body biodistribution and radiation dosimetry of this novel <sup>11</sup>C-labeled suicide irreversible tracer developed as a surrogate measure of activity of the fatty acid amide hydrolase. . . . . *Page 1993*

**Dose reduction using SUV:** Cheng and colleagues explore how the injected <sup>18</sup>F-FDG dose can be lowered while maintaining validity in comparing standardized uptake values between studies. . . . . *Page 1998*

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**<sup>18</sup>F-FDG uptake in CRC with mutated KRAS:** Iwamoto and colleagues explore through in vivo and small-animal studies the mechanisms by which *KRAS* gene mutations, which occur in ~40% of colorectal cancers, increase <sup>18</sup>F-FDG accumulation. . . . . *Page 2038*

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