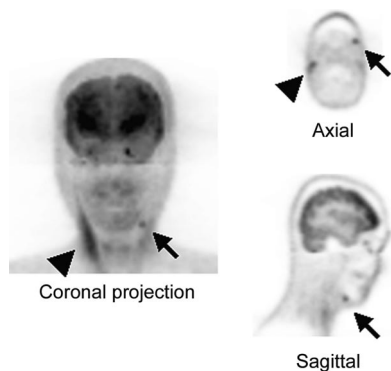


THIS MONTH IN JNM

Tulchinsky and Peters outline the development of leukocyte receptor-binding radiotracers and describe requirements for ideal infection and inflammation imaging agents. **Page 718**

Sciagrà and colleagues report on the use of gated SPECT to assess the effect of abciximab therapy on outcomes after myocardial reperfusion in patients with acute myocardial infarction. . . . **Page 722**



Slomka and colleagues apply delayed-enhancement MRI as a validation tool to measure the accuracy of a novel automated myocardial perfusion ²⁰¹Tl SPECT quantification method, based on normal limits, for detection and sizing of infarcts. **Page 728**

Fricke and colleagues evaluate the enhanced accuracy of a SPECT/low-dose CT device in myocardial perfusion scintigraphy and compare results with those from ¹³N-ammonia PET. **Page 736**

van Dyck and colleagues use ¹²³I-β-CIT to investigate the relationship between striatal dopamine transporter (DAT) protein availability and a polymorphism of the gene that codes for DAT and discuss the implications of their findings for neuropsychiatric studies. **Page 745**

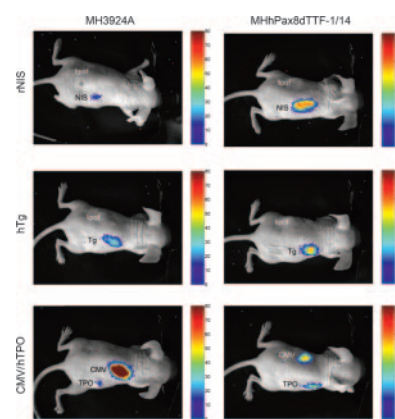
Ishimori and colleagues report on the detection of unexpected ¹⁸F-FDG-avid

additional primary malignant tumors, as well as false-positive results, in patients being evaluated by whole-body ¹⁸F-FDG PET/CT for known or suspected malignancies. **Page 752**

Israel and colleagues study the nature and significance of unexpected focal ¹⁸F-FDG uptake localized by PET/CT within the gastrointestinal tract in a retrospective review of more than 4,000 patients. **Page 758**

Henze and colleagues describe the pharmacokinetic modeling of the high uptake of ⁶⁸Ga-DOTA-TOC in meningiomas and offer data that may serve as a basis for the use of PET in radiotherapy planning and subsequent monitoring of somatostatin receptors. **Page 763**

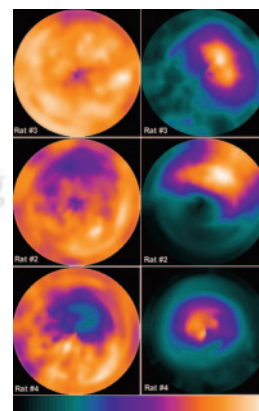
Yen and colleagues evaluate the sensitivity and prognostic significance of whole-body ¹⁸F-FDG PET for follow-up evaluation of recurrences and metastases in patients with nasopharyngeal carcinoma. **Page 770**



Yen and colleagues detail a study evaluating the utility of PET in staging untreated squamous cell carcinoma of the buccal mucosa and compare the results with those from CT and MRI and histopathology. **Page 775**

Zitzmann and colleagues report on a novel bacterial peptide display system as an alternative to phage peptide display for the isolation of new prostate tumor-specific peptides. **Page 782**

van Eerd and colleagues study the pharmacodynamics of ¹¹¹In-DPC11870 to determine the mechanism of accumulation of the radiolabeled LTB4 antagonist in infectious and inflammatory foci in an animal model. **Page 786**



Takei and colleagues elucidate the relationship between apoptosis and glucose utilization during cancer chemotherapy using ^{99m}Tc-annexin V and ¹⁸F-FDG and compare the uptake of these tracers in a rat tumor model. **Page 794**

Semnani and colleagues assess the efficacy of targeting of lung metastases in mice with various formulations of the radioiodinated thymidine analog ¹²³I-IUDR/¹²⁵I-IUDR. **Page 800**

Tait and colleagues look at ways in which changes in membrane-binding affinity, molecular charge, and method of labeling affect the biodistribution of ^{99m}Tc-annexin V in normal and apoptotic tissues. **Page 807**

Zhou and colleagues describe the potential of dual-tracer small-animal SPECT in simultaneous imaging of ^{99m}Tc-sestamibi to

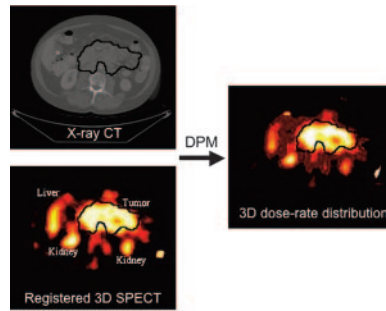
assess myocardial perfusion and of ^{111}In -labeled stem cells to delineate stem cell grafts in regions of infarct. . . . **Page 816**

Maina and colleagues compare the biologic profiles of $^{99\text{m}}\text{Tc}$ -demobesin 1 and ^{111}In -Z-070 using various gastrin-releasing peptide receptor (GRP-R)-expressing tissues of human and animal origin and discuss the implications for the development of bombesin analogs for GRP-R-targeting applications. . . . **Page 823**

Altmann and colleagues visualize functional protein-to-protein interaction as thyroid transcription factors induce reporter gene expression in rat hepatoma cells and report on the ability of these factors to induce endogenous thyroid-specific gene expression and iodide accumulation in non-thyroid tumor cells. . . . **Page 831**

Dewaraja and colleagues describe the accuracy of ^{131}I activity quantification and absorbed dose estimation when patient-specific, 3-dimensional methods are used for SPECT reconstruction and ab-

sorbed dose calculation and discuss the implications for advances in radioimmunotherapy. . . . **Page 840**



Vallabhajosula and colleagues evaluate the potential value of several factors in predicting myelotoxicity associated with ^{90}Y - and ^{131}I -labeled monoclonal antibody therapy in patients with prostate cancer. . . . **Page 850**

Brasse and colleagues use 3-dimensional whole-body PET simulations and phantom studies to investigate how gains in noise equivalent count rates

from a noiseless random coincidence estimation method are translated to improvements in signal-to-noise ratio, with accompanying potential for reduction in scan duration. . . . **Page 859**

Seo and colleagues evaluate an iterative reconstruction algorithm using SPECT/CT data from phantoms and ^{111}In -capromab pentetide studies to assess the utility of a commercial SPECT/CT system in imaging of postprostatectomy patients at risk for residual or recurrent disease. . . . **Page 868**

Cheng and colleagues describe the use of matrix-assisted laser desorption/ionization time-of-flight mass spectrometry as a potential tool for high-throughput screening and characterization of molecular imaging probes or drug design and development. . . . **Page 878**

Froidevaux and colleagues evaluate the receptor-binding potency and biodistribution of radiolabeled analogs of α -melanocyte-stimulating hormone in a mouse melanoma model. . . . **Page 887**

ON THE COVER

In patients with known cancer, work-ups often focus on the primary disease, and incidental coexistence of another primary malignant lesion can be missed. The prevalence of additional primary neoplasms is substantial. Whole-body PET with ^{18}F -FDG has been used successfully and with increasing frequency in the evaluation and clinical management of an expanding number of neoplasms. Reports also indicate that ^{18}F -FDG PET has the potential for cancer screening and can detect new malignant tumors in a small fraction of asymptomatic individuals. Because PET/CT allows precise determination of the location of ^{18}F -FDG uptake, whole-body studies may be of value in the detection of unexpected additional primary malignant tumors in patients with known or suspected malignancies.

