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# Can the Inflammatory Response Be Evaluated Using $^{18}\text{F}$ -FDG Within Zones of Microvascular Obstruction After Myocardial Infarction?

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Inflammation that occurs after acute myocardial infarction plays a pivotal role in healing by facilitating the creation of a supportive scar.  $^{18}\text{F}$ -FDG, which is taken up avidly by macrophages, has been proposed as a marker of cell-based inflammation. However, its reliability as an accurate indicator of inflammation has not been established, particularly in the early postinfarction period when regional myocardial perfusion is often severely compromised. **Methods:** Nine adult dogs underwent left anterior descending coronary occlusion with or without reperfusion. Animals were imaged between 7 and 21 d after infarction with PET/MR imaging after bolus injection of gadolinium-diethylenetriaminepentaacetic acid (DTPA), bolus injection of  $^{18}\text{F}$ -FDG, bolus injection of  $^{99\text{m}}\text{Tc}$ -DTPA to simulate the distribution of gadolinium-DTPA (which represents its partition coefficient in well-perfused tissue), and injection of  $^{111}\text{In}$ -labeled white blood cells 24 h earlier. After sacrifice, myocardial tissue concentrations of  $^{18}\text{F}$ ,  $^{111}\text{In}$ , and  $^{99\text{m}}\text{Tc}$  were determined in a well counter. Linear regression analysis evaluated the relationships between the concentrations of  $^{111}\text{In}$  and  $^{18}\text{F}$  and the dependence of the ratio of  $^{111}\text{In}/^{18}\text{F}$  to the apparent distribution volume of  $^{99\text{m}}\text{Tc}$ -DTPA. **Results:** In 7 of 9 animals,  $^{111}\text{In}$  increased as  $^{18}\text{F}$  increased with the other 2 animals, showing weak negative slopes. With respect to the dependence of  $^{111}\text{In}/^{18}\text{F}$  with partition coefficient, 4 animals showed no dependence and 4 showed a weak positive slope, with 1 animal showing a negative slope. Further, in regions of extensive microvascular obstruction,  $^{18}\text{F}$  significantly underestimated the extent of the presence of  $^{111}\text{In}$ . **Conclusion:** In the early post-myocardial infarction period,  $^{18}\text{F}$ -FDG PET imaging after a single bolus administration may underestimate the extent and degree of inflammation within regions of microvascular obstruction.

**Key Words:** myocardial infarction; myocardial inflammation; PET/MRI;  $^{18}\text{F}$ -FDG;  $^{111}\text{In}$ -labeled white blood cells; late gadolinium enhancement (LGE)

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**T**here has been significant effort to develop noninvasive imaging methods to quantify in time and location the inflammatory response after myocardial infarction toward the goal of potentially intervening with therapies that may prevent or limit the degree of adverse left ventricular remodeling. The goal of understanding the role of the inflammatory response in the progression of myocardial remodeling is not new (1), but to date the techniques have been crude and limited in both spatial and temporal resolution.

Whereas the uptake of  $^{18}\text{F}$ -FDG in regions of recent myocardial infarction was found to be an unreliable indicator of myocardial viability, recent studies have suggested that the uptake of  $^{18}\text{F}$ -FDG by macrophages could be used to identify the degree and extent of myocardial inflammation in the early postinfarction period. Here, we have focused on validating the reliability of  $^{18}\text{F}$ -FDG as a marker of postinfarction inflammation, particularly in light of the markedly compromised flow in areas of recently infarcted myocardium. Adding another layer of complexity to the use of  $^{18}\text{F}$ -FDG in this setting is the need to suppress competing uptake in the normal myocardium and its nonspecificity with respect to inflammation cell type (2,3). Despite these limitations, the combination of PET and MR imaging in 1 device, hybrid PET/MR imaging, would allow one to serially track in 3 dimensions the relationships between scarring and macrophage-based inflammation using late gadolinium enhancement (LGE) and  $^{18}\text{F}$ -FDG, respectively. The potential role of this approach to assess myocardial scarring and inflammation has already been shown in another disorder associated with inflammation, sarcoidosis (4).

Here, using both in vivo imaging and ex vivo tissue analysis, we have investigated in a well-characterized canine model of myocardial infarction the uptake of  $^{18}\text{F}$ -FDG into infarcted myocardium and its distribution into the central core areas of microvascular obstruction (MO) in comparison to the distribution of  $^{111}\text{In}$ -labeled white blood cells.

## MATERIALS AND METHODS

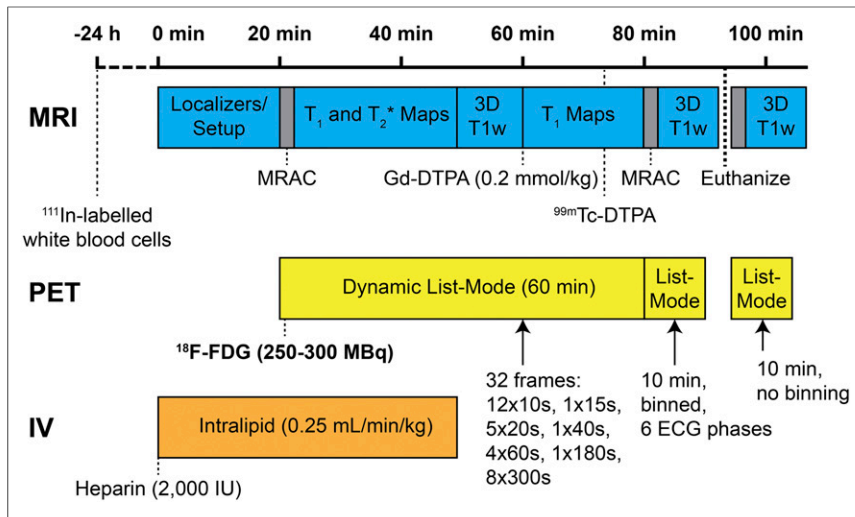
### Animals and Surgical Preparation

The study was approved by the Animal Care Committee of the University of Western Ontario (Animal Use Protocol 2011-67).

Nine adult female, bred-for-research hounds (19–24 kg) were used. Anesthesia was induced using propofol and maintained with isoflurane (2%). After a left thoracotomy, a myocardial infarction was induced by placing a snare ligature around the left anterior descending coronary

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**FIGURE 1.** Timing of entire experimental protocol. ECG = electrocardiogram; IV = intravenous; MRAC = magnetic resonance attenuation correction; T1w = T1-weighted.

artery. In 5 animals, this snare was released after 3 h, and in 4 it was left in place for the duration of the study. Animals underwent PET/MR imaging at a single imaging session in both the occlusion reperfusion group (OR), at 7, 7, 13, 14, and 20 d, and the permanent occlusion group (PO), at 7, 8, 13, and 13 d. The imaging protocol involved simultaneous cardiac MR and PET imaging.

#### **<sup>18</sup>F-FDG PET Imaging**

The suppression of normal myocardial uptake of <sup>18</sup>F-FDG was accomplished by modification of the techniques used to highlight inflamed coronary plaques and for imaging patients with suspected sarcoidosis (4–8). An infusion of 20% lipid (Intralipid; Baxter Healthcare Corp.) at a rate of 0.25 mL/min/kg was administered intravenously over a 50-min period, beginning 20 min before <sup>18</sup>F-FDG injection. In addition, 2,000 units of heparin were given intravenously immediately

before the lipid infusion to further suppress glycolysis, and the animals were fasted between 16 and 21 h before the start of imaging (7,8). All dogs were fed a normal diet. This myocardial suppression protocol was validated before the current study in 3 noninfarcted animals. These were compared with and without lipid emulsion with either a normal diet or a high-fat diet. Measurements of maximum standardized uptake values were taken in the left ventricular wall. Significance was tested using a 1-tailed paired *t* test, with *P* value set at 0.05.

Animals were anesthetized and transferred to the PET/MR imaging scanner (Biograph mMR; Siemens AG). An intravenous injection of <sup>18</sup>F-FDG (250–300 MBq) was given, and data were collected for 70 min in list-mode. Attenuation correction was performed using segmentation of images obtained with a 2-point Dixon MR imaging pulse sequence. In the last 10 min of this <sup>18</sup>F-FDG acquisition,

3-dimensional (3D) LGE MR imaging data were acquired. Retrospective binning of data was done to 6 cardiac phases. PET data were reconstructed using an iterative 3D ordered-subset expectation maximization method with 3 iterations and 21 subsets, a zoom factor of 2.5, 344 × 344 × 127 matrix size, and a 2-mm gaussian smoothing filter. The voxel size with these parameters was 0.83 × 0.83 × 2.03 mm. After the animal was euthanized while in the scanner, and without being moved, a further 10-min PET image was acquired.

#### **Cardiac MR Imaging**

Cardiac MR imaging was initiated at the start of the lipid emulsion infusion. Using Siemens Works-In-Progress software, we acquired a single-slice multiecho T2\* map to estimate the presence of hemorrhage (single breath hold; electrocardiogram-triggered; multiecho gradient-echo sequence; 8 images per slice with echo times of 2.5, 4.8, 7.1,

**TABLE 1**  
Experimental Details and Tissue Characterization

Subject*	Imaging day	PGL	<sup>18</sup> F-FDG (MBq)	<sup>111</sup> In (MBq)	<sup>111</sup> In LE	<sup>99m</sup> Tc-DTPA (MBq)	MO tissue	MO volume images (g) <sup>†</sup>	Hemorrhage tissue <sup>‡</sup>
OR1	7	3.1	259	3.7	13	28	3	5.6	4
OR2	7	3.7	211	8	22	47	2	2.3	4
OR3	13	3.3	226	8	25	37	1	0.0	2
OR4	14	3.1	213	16	51	26	1	0.0	2
OR5	20	2.6	218	3	12	37	1	0.0	1
PO1	7	2.8	257	24	83	48	4	11.6	3
PO2	8	3.0	255	17	62	47	4	4.5	4
PO3	13	3.7	288	14	45	15	1	0.0	1
PO4	13	3.3	221	16	52	41	2	1.0	4

\*Subjects are listed as occlusion type (OR = occlusion reperfusion, PO = permanent occlusion) and day of sacrifice under Imaging day.

<sup>†</sup>Extent of MO in grams was also assessed from LGE in vivo 3D image.

<sup>‡</sup>Sliced excised heart tissue was visually examined to confirm presence or absence of hemorrhage and scored as severe (4) to nonexistent (1).

PGL = arterial plasma glucose concentration in mmol/L; LE = <sup>111</sup>In white blood cell labeling efficiency; MO = microvascular obstruction rated from examination of postmortem LGE images as large (4), medium (3), small (2), or nonexistent (1).

**TABLE 2**  
Regression Equations for Non-MO Tissue

Subject	$^{111}\text{In}/^{18}\text{F}/\text{g}$ vs. $^{18}\text{F}/\text{g}$	$^{111}\text{In}/^{18}\text{F}$ vs. $\lambda_{^{99\text{mTc-DTPA}}^{\text{heart}}}$
OR1	Slope up $+0.0072 \pm 0.0012$ $R^2 = 0.279, P < 0.0001$	No slope $-0.0027 \pm 0.0044$ $R^2 = 0.023, P = 0.54$
OR2	Slope up $+0.0064 \pm 0.0008$ $R^2 = 0.46, P < 0.0001$	Slope up $+0.0043 \pm 0.0007$ $R^2 = 0.31, P < 0.0001$
OR3	Slope up $+0.0029 \pm 0.0007$ $R^2 = 0.1611, P < 0.0001$	No slope $+0.0015 \pm 0.0011$ $R^2 = 0.02, P = 0.19$
OR4	Slope up $+0.0257 \pm 0.0017$ $R^2 = 0.724, P < 0.01$	No slope $-0.017 \pm 0.0022$ $R^2 = 0.006, P > 0.05$
OR5	No slope $-0.0001 \pm 0.0004$ $R^2 = 0.0009, P = 0.78$	Slope up $+0.0026 \pm 0.0011$ $R^2 = 0.06, P = 0.024$
PO1	Slope down $-0.024 \pm 0.011$ $R^2 = 0.046, P = 0.034$	Slope up $+0.047 \pm 0.012$ $R^2 = 0.141, P < 0.0001$
PO2	Slope up $+0.0182 \pm 0.0024$ $R^2 = 0.37, P < 0.0001$	Slope up $+0.0077 \pm 0.0009$ $R^2 = 0.39, P < 0.0001$
PO3	Slope up $+0.0186 \pm 0.0034$ $R^2 = 0.28, P < 0.0001$	No slope $+0.0024 \pm 0.01$ $R^2 = 0.43, P = 0.82$
PO4	Slope up $+0.0819 \pm 0.0097$ $R^2 = 0.44, P < 0.0001$	Slope down $-0.198 \pm 0.0164$ $R^2 = 0.622, P < 0.0001$

Slope up = significant positive slope; no slope =  $P$  value for slope was greater than 0.05; slope down = significant negative slope.

Slope,  $\pm$  slope SE, square of correlation coefficient, and significance of slope obtained from linear regression analysis are given.

9.5, 11.8, 14.2, 16.5, and 18.9 ms; repetition time of 467 ms; 2 averages; 3.5-mm slice thickness;  $192 \times 156$  matrix, resulting in the voxel size of  $1.35 \times 1.35$  mm in plane  $\times$  3.5 mm nominal slice thickness). Because of cardiac motion, only the first 5 echoes were used to estimate the presence of susceptibility, which is consistent with hemorrhage. A bolus of gadolinium contrast (0.2 mmol/kg) (Magnevist; Bayer Inc.) was infused at 0.3 mL/s, followed by a 40-mL saline infusion. Twenty minutes after the bolus infusion, a respiration- and cardiac-gated 3D whole-heart inversion-recovery gradient-echo pulse sequence was acquired as previously described (4,9) using an integrated parallel-acquisition technique and a fat-suppressed isotropic/voxel size of  $1 \times 1 \times 1$  mm (echo time, 1.3 ms;  $20^\circ$  flip angle; and variable inversion times to suppress signal in noninfarcted myocardium). The animal was then euthanized, and the whole-heart inversion-recovery gradient-echo sequence was repeated while the 3D 10-min  $^{18}\text{F}$ -FDG images were acquired.

The timing of the entire experimental protocol is shown in Figure 1.

### Image Processing

The 3D LGE images taken in the live animal after the gadolinium-diethylenetriaminepentaacetic acid (DTPA) injection were processed for the size in grams of the region of MO. Sequential short-axis LGE images underwent signal-threshold-based quantification of total volume of hyperenhancement reported in grams using a signal threshold versus reference mean technique (9).

### In Vitro Tissue Counting

After euthanasia, the hearts were excised and cut into 7 slices (base to apex). The slices were washed and visually examined for evidence of hemorrhage and scored as 4 for severe to 1 for nonexistent. With the postmortem 3D  $^{18}\text{F}$ -FDG and LGE images used as a guide, the myocardial tissue was segmented into an average of 107 (range, 93–123) samples, with an average weight of 0.71 g (SD, 0.18 g).

Blood samples were taken and the white blood cells isolated and then labeled with  $^{111}\text{In}$ -oxine (Cardinal Health) using standard methods (10). Labeled cells were injected between 21 and 28 h before the start of imaging.

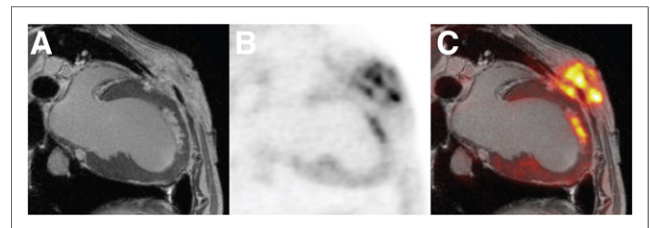
Six minutes after the bolus injection of gadolinium-DTPA, a bolus injection of  $^{99\text{mTc}}$ -DTPA was given to allow for ex vivo measurement of the concentration of a tracer that would simulate the tissue distribution of gadolinium-DTPA (11). On euthanasia, ten 1-mL blood samples were taken so that partition coefficients for  $^{99\text{mTc}}$ -DTPA for heart tissue could be calculated as  $^{99\text{mTc}}$  activity per gram of myocardium divided by  $^{99\text{mTc}}$  activity per milliliter of blood. The partition coefficients ( $\lambda$ ) of  $^{99\text{mTc}}$ -DTPA and gadolinium-DTPA (Magnevist) are related in heart tissue (11):

$$\lambda_{\text{Gd-DTPA}}^{\text{heart}} = 1.094 \times \lambda_{^{99\text{mTc-DTPA}}^{\text{heart}}} \quad \text{Eq. 1}$$

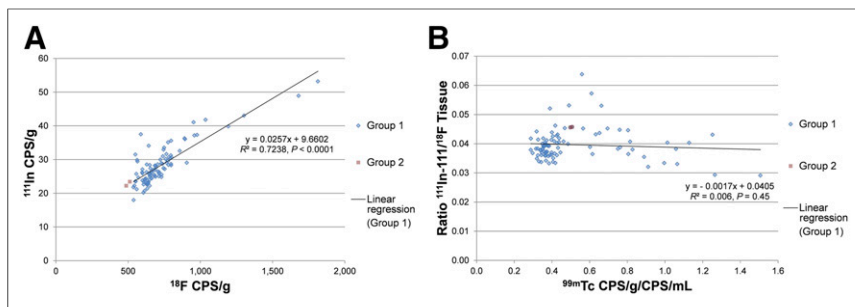
Tissue and blood samples were counted for  $^{18}\text{F}$ ,  $^{99\text{mTc}}$ , and  $^{111}\text{In}$  in a high-purity germanium well counter, with an active volume of 190 mL, an efficiency of greater than 10%, and a spectral resolution of less than 2 keV over the range of energies (140–511 keV) detected. Each sample was counted for 2 min, with counts corrected for detector dead time and radioisotope decay. The tissue samples were separated into 2 groups. Group 2 corresponded to tissue with concentration at least 1 SD below the mean  $^{18}\text{F}$  concentration. These tissue samples generally had been taken from regions in which MO was demonstrated on the gadolinium-DTPA-enhanced images. It was assumed all other samples came from tissue that was not so affected. Results for each sample were tabulated as  $^{111}\text{In}$  (cps/g) versus  $^{18}\text{F}$  (cps/g) and as the ratio of  $^{111}\text{In}$  to  $^{18}\text{F}$  as a function of  $\lambda_{^{99\text{mTc-DTPA}}^{\text{heart}}}$ .

### Statistical Analysis

The results from the tissue counting for group 1 were fit by linear regression (SigmaPlot 13; Systat Software Inc.). For each dog's data, 2 regressions were fit for the non-MO data: 1 to  $^{111}\text{In}$  (cps/g) versus  $^{18}\text{F}$  (cps/g) and 1 to the ratio of  $^{111}\text{In}$  to  $^{18}\text{F}$  versus  $\lambda_{^{99\text{mTc-DTPA}}^{\text{heart}}}$ .



**FIGURE 2.** Postmortem MR (A), PET (B), and PET/MR (C) images of subject OR4 (14 d). MR image demonstrates anteroapical subendocardial infarct. Increased activity of  $^{18}\text{F}$ -FDG in chest wall in PET image is at site of surgery.



**FIGURE 3.** Quantitative analysis of tissue samples for same subject as shown in Figure 2 (subject OR4). (A) Regression of  $^{111}\text{In}$  versus  $^{18}\text{F}$  was strong and slope significantly positive. (B) Regression of  $^{111}\text{In}/^{18}\text{F}$  versus  $\lambda_{99\text{mTc-DTPA}}^{\text{heart}}$  was weak and slope not significant, indicating that  $^{111}\text{In}/^{18}\text{F}$  ratio was independent of  $\lambda_{99\text{mTc-DTPA}}^{\text{heart}}$ .

## RESULTS

In the preliminary experiments evaluating myocardial suppression of  $^{18}\text{F}$ -FDG uptake, there was a significant drop in maximum standardized uptake value when lipid emulsion was administered independent of diet ( $P = 0.02$ ) and when animals were given a normal diet ( $P = 0.0084$ ) but no significant effect of lipid emulsion when a high-fat diet was given ( $P = 0.24$ ). Hence, it was decided to use lipid emulsion infusion and not to use a high-fat diet. For all these experiments, animals were injected with 2,000 IU of heparin (7) as others have demonstrated the combination of heparin and lipid emulsion as effective (8).

The experimental parameters, except for  $^{111}\text{In}$ -labeling efficiency, were consistent for all the animals whereas the variation of extensiveness of myocardial injury varied from mild to severe (Table 1). The linear regression analysis showed an expected relationship between  $^{111}\text{In}/\text{g}$  versus  $^{18}\text{F}/\text{g}$  consistent with  $^{18}\text{F}$ -FDG representing inflammation in 7 of 9 dogs; however, in only 4 of these 7 was the relationship independent of the partition coefficient (Table 2). Subject OR4's (14 d) images showed a subendocardial infarct without evidence of MO and limited evidence of hemorrhage (Fig. 2). The tissue analysis for this animal (Fig. 3) demonstrated that there was a significant positive slope between  $^{111}\text{In}$  and  $^{18}\text{F}$  concentrations. Also, the ratio of  $^{111}\text{In}$  to  $^{18}\text{F}$  was constant (i.e., independent of the  $^{99\text{mTc-DTPA}}$  partition coefficient). Images for PO1 (7 d) show a large area of MO (Fig. 4), with the corresponding tissue analysis (Fig. 5) showing a significant negative slope of  $^{111}\text{In}$  to  $^{18}\text{F}$  and that the ratio of  $^{111}\text{In}$  to  $^{18}\text{F}$  significantly increases as the partition coefficient increases.

Single-slice T2\*-weighted images for OR4 (Fig. 6) show little to no hemorrhage, although visual evaluation of excised heart slices showed mild hemorrhage present (Table 1). T2\*-weighted images for PO1 (Fig. 7) provide evidence of significant hemorrhage supported by visual inspection of the excised heart (Table 1). Results for the other 7 animals fall between what is seen in these 2 animals, which represents the extremes. In all of the other animals but one (OR5; at 20 d), the concentration of  $^{111}\text{In}$  increased as  $^{18}\text{F}$  increased. With respect to the dependence of  $^{111}\text{In}/^{18}\text{F}$  to partition coefficient, 3 were similar to OR4 (Fig. 3), with zero slope; 4 showed a significant positive slope; and 1 showed a significant negative slope.

## DISCUSSION

If  $^{18}\text{F}$ -FDG was strictly a measure of inflammation, it is expected that there would be a strong positive linear regression between the tissue concentration of  $^{111}\text{In}$  and  $^{18}\text{F}$  and that there would be no

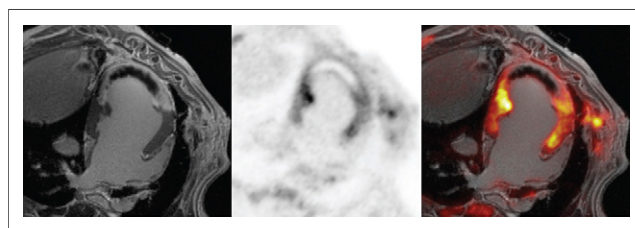
dependence on the ratio of  $^{111}\text{In}/^{18}\text{F}$  to the partition coefficient of DTPA. As seen for subject PO1 in Figures 4 and 5, this clearly cannot be assumed in tissue with MO. At the minimum values of the partition coefficient, this ratio increases 7-fold, suggesting that in the region of MO (reduced apparent partition coefficient), there were  $^{111}\text{In}$ -labeled white blood cells but little  $^{18}\text{F}$  consistent with the postmortem images of  $^{18}\text{F}$  distribution. Furthermore, the pattern shown in Figure 3, which is the desired pattern if  $^{18}\text{F}$ -FDG represents inflammation in the cardiac-suppressed tissue after MI, is not necessarily the pattern seen even in tissue that is reasonably well perfused—that is, that does not

have evidence of MO. Our results are mixed, with clear evidence that the uptake of  $^{18}\text{F}$ -FDG may be at variance with the degree of inflammation as measured with  $^{111}\text{In}$ -labeled white blood cells after myocardial infarction.

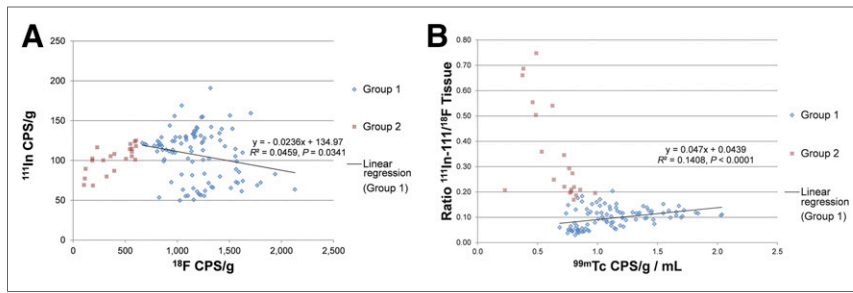
In 7 of 9 animals,  $^{111}\text{In}$  cps/g increased linearly as the  $^{18}\text{F}$  cps/g increased, suggesting that in these animals,  $^{18}\text{F}$ -FDG was increasing as the number of white blood cells was increasing. However, in only 4 of these animals was the  $^{111}\text{In}/^{18}\text{F}$  ratio independent of the partition coefficient in non-MO tissue. In the other 3 animals, the slope was significantly positive in 2 and negative in 1, indicating that although  $^{18}\text{F}$  was increasing as  $^{111}\text{In}$  was increasing the ratio was not independent of  $\lambda_{99\text{mTc-DTPA}}^{\text{heart}}$  (i.e.,  $^{18}\text{F}$ -FDG was underestimating in 2 and overestimating in 1 the number of inflammatory cells).

The potential explanations for this disconnect between indium concentrations, reflective of the presence of inflammatory cells injected back into circulation 1 d earlier, and the  $^{18}\text{F}$ -FDG concentrations in myocardial tissue may be due to the markedly compromised perfusion to the MO zone reducing the availability of  $^{18}\text{F}$ -FDG to inflammatory cells. After a single bolus injection, the concentration of  $^{18}\text{F}$ -FDG rapidly declined. In the setting of markedly reduced flow, there may have been insufficient opportunity for inflammatory cells to take up the tracer, whereas indium-labeled white blood cells had been circulating between 21 and 28 h before animal sacrifice and tissue harvesting. In contrast, in areas of infarction in which perfusion was not severely compromised, our results indicated that although  $^{18}\text{F}$ -FDG accumulation increased as  $^{111}\text{In}$  increased it may not increase proportionately, suggesting that the use of  $^{18}\text{F}$ -FDG imaging in these areas must be interpreted as a nonquantitative indicator of the degree of inflammatory cell activity.

One potential limitation of our study relates to the validity of the measurement of  $\lambda_{99\text{mTc-DTPA}}^{\text{heart}}$  as an indication of the value of  $\lambda_{\text{Gd-DTPA}}^{\text{heart}}$ . This association may have a small uncertainty because



**FIGURE 4.** Postmortem MR (left), PET (middle), and PET/MR (right) images of subject PO1 (7 d). Anteroapical infarct is transmural, with large central MO that is void in both  $^{18}\text{F}$ -FDG and gadolinium-DTPA but is surrounded by zone of increased  $^{18}\text{F}$ -FDG and gadolinium-DTPA.



**FIGURE 5.** Quantitative analysis of tissue samples for same subject as shown in Figure 4 (subject PO1). Regression analysis excluded samples estimated to be from MO tissue (designated as group 2). Regression of  $^{111}\text{In}$  vs.  $^{18}\text{F}$  in A is modest but negative slope significant. Regression of  $^{111}\text{In}/^{18}\text{F}$  vs.  $^{99\text{m}}\text{Tc-DTPA}$  in B is significant and indicates that as partition coefficient increases  $^{111}\text{In}$  increases more quickly than  $^{18}\text{F}$ . This ratio is much higher in MO zone, suggesting that  $^{18}\text{F}$ -FDG significantly underestimated extent of inflammation in this zone.

the  $^{99\text{m}}\text{Tc-DTPA}$  was injected into the live animal on average 6 min (range, 5–7 min) after the gadolinium-DTPA injection. Hence,  $^{99\text{m}}\text{Tc-DTPA}$  had 6 min less time to penetrate areas of MO, possibly resulting in an underestimation of the partition coefficient in some infarcted tissue samples.

A limitation of our work relates to the interpretation of the results given the potential difference between the nature of the white blood cells labeled with  $^{111}\text{In}$  that accumulate in the infarct region and normal myocardium versus those that take up  $^{18}\text{F}$ -FDG. There is evidence in dogs that neutrophil density peaks in reperfused infarcted myocardium within 1 d, with low concentra-

tions by day 7, whereas macrophage concentration peaks around 7 d after infarction (12). Because in blood the number of neutrophils exceeds the number of monocytes by a factor of approximately 10, it is difficult to conclude what fraction of the labeled white blood cells in the infarcted myocardium is  $^{111}\text{In}$ -labeled neutrophils or  $^{111}\text{In}$ -labeled monocytes. This difficulty is further confounded because it is generally assumed that monocytes are transformed to macrophages (2) and then replicate and accumulate in the infarct region. These transformed monocytes, even if still labeled with  $^{111}\text{In}$ , would have the amount of label per cell diluted during replication. In addition, we do not know to what extent neutrophils within the region of MO would be labeled

by  $^{18}\text{F}$ -FDG as well as macrophages. Adding to this is the growing evidence that an appreciable fraction of the macrophages that accumulate in the infarcted region may not be transformed monocytes but rather resident macrophages in the myocardium (13,14).

The study design could have been improved by measuring blood flow using microspheres in the tissue samples. However, the absence of MR imaging contrast after a bolus injection has become an accepted, albeit qualitative, standard. We have characterized tissue blood flow in these 2 animal models using microspheres in the past (15,16).

Although we have followed in detail  $^{111}\text{In}$ -oxine labeling of white blood cells (10), we achieved lower-than-normal labeling efficiencies. Because the greatest discrepancy between the  $^{111}\text{In}$  counts and  $^{18}\text{F}$  counts was in subject PO1, which had the highest labeling efficiency of 83%, we do not believe that the low labeling efficiencies affected the conclusions drawn from our results.

## CONCLUSION

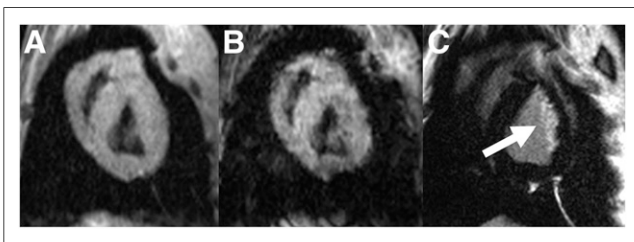
Our results suggest some limitations regarding the use of a bolus injection of  $^{18}\text{F}$ -FDG to study the inflammatory process after myocardial infarction even when active uptake of  $^{18}\text{F}$ -FDG in normal myocardial tissue is suppressed. In the presence of large regions of MO,  $^{18}\text{F}$ -FDG may not reliably and accurately represent the degree of inflammatory cell activity, possibly due to compromised delivery. Further, even in areas of infarction without MO, the degree of inflammation may be underestimated or even overestimated, the reasons for which remain, at this time, unknown.

## DISCLOSURE

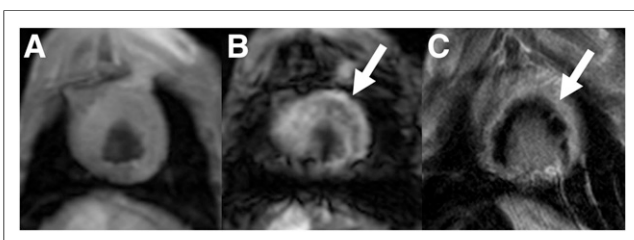
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**FIGURE 6.** Gradient-echo images (i.e.,  $T_2^*$  imaging) of subject OR4 (also featured in Figs. 2 and 3) at echo times of 2.5 (A) and 9.5 ms (B) and matched LGE image (C). No evidence of hemorrhage was identified. White arrow indicates subendocardial infarction. Examination of tissue and assessment of LGE images showed no evidence of MO whereas examination of tissue (Table 1) suggested mild hemorrhage.



**FIGURE 7.** Gradient-echo images (i.e.,  $T_2^*$  imaging) of subject PO1 (also featured in Figs. 3 and 4) at echo times of 2.5 (A) and 9.5 ms (B) and matched LGE image (C). White arrow on B indicates susceptibility indicative of hemorrhage; white arrow on C indicates transmural infarct and associated region of MO. Examination of tissue (Table 2) from this region indicated significant hemorrhage.

## REFERENCES

1. Fishbein MC. The histopathologic evolution of myocardial infarction. *Chest*. 1978;73:843–849.
2. Satomi T, Ogawa M, Mori I, et al. Comparison of contrast agents for atherosclerosis imaging using cultured macrophages: FDG versus ultrasmall superparamagnetic iron oxide. *J Nucl Med*. 2013;54:999–1004.
3. Jones HA, Cadwallader KA, White JF, Uddin M, Peters AM, Chilvers ER. Dissociation between respiratory burst activity and deoxyglucose uptake in human neutrophil granulocytes: implications for interpretation of  $^{18}\text{F}$ -FDG PET images. *J Nucl Med*. 2002;43:652–657.
4. White JA, Rajchl M, Butler J, Thompson RT, Prato FS, Wisenberg GW. First clinical experience of simultaneous PET-MRI imaging for the diagnosis of disease: active cardiac sarcoidosis. *Circulation*. 2013;127:e639–e641.
5. Wykrzykowska J, Lehman S, Williams G, et al. Imaging of inflamed and vulnerable plaque in coronary arteries with  $^{18}\text{F}$ -FDG PET/CT in patients with suppression of myocardial uptake using a low-carbohydrate, highfat preparation. *J Nucl Med*. 2009;50:563–568.
6. Demeure F, Hanin FX, Bol A, et al. A randomized trial on the optimization of  $^{18}\text{F}$ -FDG myocardial uptake suppression: implications for vulnerable coronary plaque imaging. *J Nucl Med*. 2014;55:1629–1635.
7. Ishimaru S, Tsujino I, Takei T, et al. Focal uptake on  $^{18}\text{F}$ -fluoro-2-deoxyglucose positron emission tomography images indicates cardiac involvement of sarcoidosis. *Eur Heart J*. 2005;26:1538–1543.
8. Prinzen FW, van der Vusse GJ, Coumans WA, Kruger R, Verlaan CW, Reneman RS. The effect of elevated arterial free fatty acid concentrations on hemodynamics and myocardial metabolism and blood flow during ischemia. *Basic Res Cardiol*. 1981;76:197–210.
9. White JA, Fine N, Gula LJ, et al. Fused whole-heart coronary and myocardial scar imaging using 3-T CMR: implications for planning of cardiac resynchronization therapy and coronary revascularization. *JACC Cardiovasc Imaging*. 2010;3:921–930.
10. Roca M, de Vries EFJ, Jamar F, Israel O, Signore A. Guidelines for the labelling of leucocytes with  $^{111}\text{In}$ -oxine. *Eur J Nucl Med Mol Imaging*. 2010;37:835–841.
11. Prato FS, Wisenberg G, Marshall TP, Uksik P, Zabel P. Comparison of the biodistribution of gadolinium-153 DTPA and technetium-99m DTPA in rats. *J Nucl Med*. 1988;29:1683–1687.
12. Dewald O, Ren G, Duerr GD, et al. Of mice and dogs: species-specific differences in the inflammatory response following myocardial infarction. *Am J Pathol*. 2004;164:665–677.
13. Epelman S, Lavine KJ, Randolph GJ. Origin and functions of tissue macrophages. *Immunity*. 2014;41:21–35.
14. Yan X, Anzai A, Katsumata Y, et al. Temporal dynamics of cardiac immune cell accumulation following acute myocardial infarction. *J Mol Cell Cardiol*. 2013;62:24–35.
15. Pereira RS, Prato FS, Sykes J, Wisenberg G. Assessment of myocardial viability using MRI during a constant infusion of Gd-DTPA: further studies at early and late periods of reperfusion. *Magn Reson Med*. 1999;42:60–68.
16. Pereira RS, Prato FS, Leks K, Sykes J, Wisenberg G. Contrast-enhanced MRI for the assessment of myocardial viability after permanent coronary artery occlusion. *Magn Reson Med*. 2000;44:309–316.