Enhancing magnetic resonance imaging tumor detection with fluorescence intensity and lifetime imaging

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Abstract. Early detection is important for many solid cancers but the images provided by ultrasound, magnetic resonance imaging (MRI), and computed tomography applied alone or together, are often not sufficient for decisive early screening/diagnosis. We demonstrate that MRI augmented with fluorescence intensity (FI) substantially improves detection. Early stage murine pancreatic tumors that could not be identified by blinded, skilled observers using MRI alone, were easily identified with MRI along with FI images acquired with photomultiplier tube detection and offset laser scanning. Moreover, we show that fluorescence lifetime (FLT) imaging enables positive identification of the labeling fluorophore and discriminates it from surrounding tissue autofluorescence. Our data suggest combined-modality imaging with MRI, FI, and FLT can be used to screen and diagnose early tumors. © 2010 Society of Photo-Optical Instrumentation Engineers. [DOI: 10.1117/1.3509111]

Keywords: cancer detection; pancreatic cancer; orthopedic models; magnetic resonance imaging; red fluorescent protein; fluorescence intensity.

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1 Introduction

Early detection is critical to increase the cure rate of cancer. Unfortunately, the images provided by ultrasound, magnetic resonance imaging (MRI), and computed tomography (CT) applied alone or in combination are often inconclusive1–10 for the identification of tumors less than 10 mm. Endoscopic ultrasonography (EUS) offers high resolution and high sensitivity but often cannot discriminate tumor tissue from inflammation, as in the case of pancreatic cancer.5

The present report addresses the hypothesis that MRI augmented by fluorescence intensity (FI) imaging is more likely to detect a small tumor than MRI alone.11–15
modality approach may be further augmented by fluorescence lifetime (FLT) imaging. The use of FLT imaging has recently been extended from fluorescence lifetime imaging microscopy (FLIM) to whole-body imaging of rodents in vivo. Fluorescence lifetime depends on the unique decay rate of an excited fluorophore relaxing to its ground state. Generally, the rate of decay is understood to be a sum of the rates that depopulate the excited-state of the fluorophore. The emitted fluorescence decay is understood to be a sum of the rates that depopulate the excited-state of the fluorophore. Briefly, the mutual information criteria. Briefly, the journal of biomed imaging may be further augmented by fluorescence lifetime (FLT) imaging. The use of FLT imaging has recently been extended from fluorescence lifetime imaging microscopy (FLIM) to whole-body imaging of rodents in vivo. Fluorescence lifetime depends on the unique decay rate of an excited fluorophore relaxing to its ground state. Generally, the rate of decay is understood to be a sum of the rates that depopulate the excited-state of the fluorophore. The emitted fluorescence decay is understood to be a sum of the rates that depopulate the excited-state of the fluorophore. Briefly, the mutual information criteria. Briefly, the

2.1 Imaging

2.1.1 MRI

Four mice, at each of three time points, which were 5, 7, and 12 days after implantation, were anesthetized with 1.5% isoflurane. They received intravenous gadolinium contrast agent (Magnevist), and were placed within a bird-cage-type of mouse body coil. A respiratory bellows was used to monitor respiration and a nose cone delivered the inhalation anesthetic. The mouse was positioned at the isocenter of the scanner magnet, and the bore was warmed to 36°C. A fast 3-D localized image acquisition was performed, followed by T1-weighted 3-D structural scans. The 3-D scan acquisition time was 1357 s, the acquisition matrix was 256 × 128, repetition time (TR) = 885.7, the time to echo (TE) = 10 ms, averages = 6, and the slice thickness was 0.6 mm. The 3-D data sets were reconstructed and saved in standard DICOM (digital imaging and communications in medicine) format.

2.1.2 FL and FLT imaging

The Advanced Research Technologies (ART) Optix-MX2 imaging system was used to acquire both the intensity and the time-domain-based fluorescence images. This apparatus includes four pulsed lasers (picosecond pulse, 80-MHz duty cycle), eight filters, and a TCSPC PMT for TD detection. A brightfield image was acquired first, followed by TD fluorescence detection from which FI and FLT images were obtained. The FL and FLT image acquisition times ranged between 2 and 7 min, depending on the size of region of interest. The Optix-MX2 images were acquired with euthanized mice that were placed supine. The image data were saved directly on a PC for later processing.

2.2.3 Confirmation of tumors in each subject

To determine whether each mouse had developed a tumor, and to provide a record of tumor location, the mice were euthanized and then laparotomized. The pancreatic tumor was exposed by blunt dissection. Direct brightfield and fluorescence imaging of the exposed tumor provided a record of the tumor for each mouse. FI images were acquired with an Olympus OV100 Small Animal Imaging System (Olympus Corp., Tokyo, Japan). This instrument was used because of its ability to magnify over a large range. The OV100 included a DP70 CCD camera (Olympus Corp., Tokyo, Japan), with a range of lens magnifications [0.14 × (63 × 47 mm imaging area) to 16 × (0.6 × 0.45 mm)].

2.3 Image Data Processing

2.3.1 Image coregistration and fusion

Coronal MRI volume image slices were coregistered with the Optix FI/FLT images using a custom program prepared with IDL software (ITT Visual Solutions, Boulder, Colorado). Registration was based on the mutual information criteria. Briefly, the

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3-D MR data volume of the mouse torso was coregistered to the 2-D fluorescence image by a computer algorithm that iteratively rotated and translated the MR volume in three rotational and three translational parameters. Initially, regions of interest were drawn to isolate the anatomic regions to be matched and then the mutual information criteria between the two images were maximized. During each iteration, the 3-D MR data volume was reprojected into a 2-D image prior to determining the mutual information criteria. The mutual information criteria incorporated in the software has a maximum at the best registration for a given set of images. This maximum value varies depending on the content of the two images. The final coregistered MR tomographic set was used for tumor identification. The MR volume slice that most closely corresponded in terms of $z$ to the FI/lifetime image used for coregistration was selected and alpha blended with the fluorescence image to create a fused image. The opacity of the FI image was adjusted to permit simultaneous viewing of its corresponding MR. The FI image was processed prior to alpha blending, so that only the lifetime pixels corresponding to the fluorophore were visible.

2.3.2 Assessing Tumor Detection with MRI, FI, and FLT

The images from the MRI volumes were assembled into panels for each mouse with an identification code. Subsequently, two trained observers, one an oncologist and the second a radiologist, skilled in oncology screening, were asked to determine which mice had identifiable tumors in the pancreas, and then tabulate their assessments. These two evaluators were not informed as to the stage or age of the tumor implantation in each mouse.

3 Results

3.1 Tumor Detection by MRI Alone

MRI enabled detection of tumors for all four of the most advanced tumors. These were imaged at 12 days from implantation (Table 1) and were over 1.0 cm in diameter. For the 7-day group, however, two of the four tumors were not considered detectable by one of the expert readers, and both readers had difficulty detecting tumors at the 5-day time point (smallest tumors). With both readers, only one positive identification at 5 days was tabulated, and this was considered questionable. Figure 1(a) left panel is a T1-weighted gadolinium-enhanced MR image slice of a mouse bearing a 5-day pancreatic tumor, which was not detectable. In contrast, the tumors at 7 and 12 days, middle and right panels, respectively, both were visible by MRI. Figure 1(b) shows a fluorescence image of the same subject in the Fig. 1(a) left panel, after laparotomy and dissection, and the tumor is clearly highlighted confirming its presence.

3.2 Tumor Detection by MRI and FI Together

FI acquired by photon counting with scanning laser illumination and a PMT detector on the Opitx-MX2® system, supported tumor detection (Table 1). For all 12 mice, MRI unambiguously detected 6/12 tumors, for reader 2 the worst case, while FI alone detected 10/12, and combined modality detection was 12/12. MRI versus FI did not differ significantly, but MRI alone versus combined modality (MRI + FI) differed according to Fisher’s exact probability ($p = 0.014$). FLT did not contribute significantly to detection.

3.3 Tumor Detection by FLT and FI Together

Figure 2, left panel, shows the same mouse as in Fig. 1(a); the tumor presence is indicated by the fluorescence hotspots near the left flank, but other intensely fluorescent sites are present. These likely arose from vegetable food matter in the gut. Figure 2, middle panel, shows a cleanly delineated area of the gated FLT map near the left flank in the same approximate areas as a candidate FI hotspot. In this image, only pixels from the FLT associated with the RFP fluorophore were displayed. The tumor is separated from the surrounding tissue. Figure 2, right panel, is a fluorescence image acquired after laparotomy...
Table 1  Tumor detection with different imaging modalities applied alone and in combination.

<table>
<thead>
<tr>
<th>Imaging Modality</th>
<th>Time point [days]</th>
<th>MRI Reader 1</th>
<th>MRI reader 2</th>
<th>FI (PMT-Laser Scanning)</th>
<th>Combined MRI + FI</th>
<th>Confirmed by FLT</th>
</tr>
</thead>
<tbody>
<tr>
<td>5 subject 1</td>
<td>ND</td>
<td>ND</td>
<td>Det</td>
<td>FI</td>
<td>Det</td>
<td>Yes</td>
</tr>
<tr>
<td>5 subject 2</td>
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<td>ND</td>
<td>Det</td>
<td>FI</td>
<td>Det</td>
<td>Yes</td>
</tr>
<tr>
<td>5 subject 3</td>
<td>ND</td>
<td>ND</td>
<td>Det</td>
<td>FI</td>
<td>Det</td>
<td>No</td>
</tr>
<tr>
<td>5 subject 4</td>
<td>Det-AMB</td>
<td>Det</td>
<td>Det</td>
<td>FI</td>
<td>Det</td>
<td>Yes</td>
</tr>
<tr>
<td>7 subject 1</td>
<td>Det</td>
<td>Det</td>
<td>Det</td>
<td>FI</td>
<td>Det</td>
<td>Yes</td>
</tr>
<tr>
<td>7 subject 2</td>
<td>Det</td>
<td>Det</td>
<td>ND</td>
<td>FI</td>
<td>No</td>
<td></td>
</tr>
<tr>
<td>7 subject 3</td>
<td>Det</td>
<td>ND</td>
<td>Det</td>
<td>FI</td>
<td>No</td>
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<tr>
<td>7 subject 4</td>
<td>Det</td>
<td>ND</td>
<td>Det</td>
<td>FI</td>
<td>No</td>
<td></td>
</tr>
<tr>
<td>12 subject 1</td>
<td>Det</td>
<td>Det</td>
<td>ND</td>
<td>FI</td>
<td>Det</td>
<td>No</td>
</tr>
<tr>
<td>12 subject 2</td>
<td>Det</td>
<td>Det</td>
<td>Det</td>
<td>FI</td>
<td>Det</td>
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<tr>
<td>12 subject 3</td>
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<td>Det</td>
<td>Det</td>
<td>FI</td>
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<tr>
<td>12 subject 4</td>
<td>Det</td>
<td>Det</td>
<td>Det</td>
<td>FI</td>
<td>Yes</td>
<td></td>
</tr>
</tbody>
</table>

Det, detected; Det-AMB, detected but not confirmed; ND, not detected.

and dissection, confirming the presence of the tumor. The use of FLT with FI confirmed the presence of the RFP fluorophore in three of four of the smallest (5 day) tumors.

The top image on the left of Fig. 3 shows tumor RFP FI and abdominal autofluorescence FI. The lower image acquired from a different subject indicates tumor FI and two intestinal FI sites. The FLT graph was prepared using data for all sites in the two images. Note that the RFP FLT decay curves for tumors from two different subjects are virtually identical (upper curves), and that the intestinal autofluorescence FLT decay curves (lower) are also the same but clearly different from the RFP curves.

3.4 Coregistration of MRI, FI, and FLT
Coregistration of the MRI image slices with gray-scale optical, FI, and FLT images, enabled the tumor to be highlighted within its anatomical context, as shown in Fig. 4 (left and right panels). The FLT decay image in particular (Fig. 4, right panel), where nontumor FLT decay pixels were set to zero, clearly delineated the tumor within its anatomical environment, which was described by the fused MRI and gray-scale (optical) images of the mouse torso.

4 Discussion
The combination of MRI along with laser-scanning, offset-illumination FI imaging markedly improved the detection of small tumors. The addition of FLT mapping further augmented this enhancement in a significant proportion of cases, allowing discrimination of tumor tissue from surrounding host tissue. The fluorescence detection process can be viewed as comprising two steps: (1) detection of fluorophores associated with the target and (2) discrimination of the target fluorescence from surrounding autofluorescence. Fluorescence emission lifetime decay, owing to its independence from intensity, addresses the second part of the process.
In this study, each mouse served as its own control whereby the presence of a tumor was confirmed by dissection and direct observation after it had been imaged. We found no “false positives,” i.e., incorrectly calling a tumor to be present on MRI or combined modality when in fact one was not present. Since these tumors were entirely confined to the pancreas, especially in the early time points when the tumors were extremely small, normal tissue autofluorescence was not different between tumor-bearing and non-tumor-bearing mice. The intestines, rather than the pancreas, were the main source of autofluorescence due to their food content.

In many instances, the tumor FLT was discriminated cleanly from surrounding tissues, but in other instances, the weaker intensity of the fluorescence signal did affect separation on the basis of lifetime decay. This highlights one difficulty inherent in FLT imaging, which is that a very large number of decay curves must be tediously analyzed to separate the fluorophore of interest.21

The best solution to minimize interference from autofluorescence when the fluorophore signal is expected to be weak, for example, due to sparse labeling of a tumor, is to use a labeling fluorophore with the longest lifetime possible. Unfortunately, lifetime has an inverse relationship with another desirable fluorophore attribute, namely, long emission wavelength to reduce scatter and improve tissue penetrance. Most organic fluorophores that have longer wavelength emissions also have short fluorescence emission lifetimes.17 An exception are quantum dots, which emit long wavelengths and at the same time have a relatively long fluorescence decay.15 An example is Qdot800,
which emits at 800 nm, a very long wavelength that enables deeper imaging, and it has a very long FLIM decay of 20 ns, which is ideal. However, Qd680 is difficult to link to biological molecules such as targeted antibodies, and one of its components is cadmium metal, which is toxic. Therefore, new labeling fluorophores need to be developed that are optimized in terms of emission wavelength, fluorescence lifetime, and toxicity profile.

In summary, this paper demonstrated that combined modality imaging of the mouse pancreatic tumor model using MRI along with FI, augmented tumor detection over MRI alone. Future work will address the clinical potential of this approach in particular with fluorescence endoscopy.

Future studies will involve comparison of MRI to an injected fluorophore, to account for tumor delivery issues related to the targeted fluorophore. This early study used cells expressing a fluorophore to avoid complexities such as the distribution of the fluorophore, the kinetics of tumor accumulation of the fluorophore, extravasation into normal tissues, and concentration issues. We then sought to demonstrate that with optical methods it is feasible to augment MRI with the best-case scenario. A much more comprehensive study comparing MRI to an injected fluorophore will be carried out in the future. This study, however, indicates that MRI can be augmented by FI, which has implications for clinical application for early detection of cancer.

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