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Abstract. Rare-earth-doped nanocomposites have appealing optical properties for use as biomedical contrast agents, but few systems exist for imaging these materials. We describe the design and characterization of (i) a preclinical system for whole animal in vivo imaging and (ii) an integrated optical coherence tomography/confocal microscopy system for high-resolution imaging of ex vivo tissues. We demonstrate these systems by administering erbium-doped nanocomposites to a murine model of metastatic breast cancer. Short-wave infrared emissions were detected in vivo and in whole organ imaging ex vivo. Visible upconversion emissions and tissue autofluorescence were imaged in biopsy specimens, alongside optical coherence tomography imaging of tissue microstructure. We anticipate that this work will provide guidance for researchers seeking to image these nanomaterials across a wide range of biological models.

Keywords: short-wave infrared imaging; NIR-II imaging; confocal microscopy; preclinical imaging; multimodal imaging.

1 Introduction

There is growing interest in the development of luminescent optical probes emitting at wavelengths beyond 1000 nm, in the short-wave infrared (SWIR) or NIR-II spectral region. These probes aim to take advantage of low scattering, low autofluorescence, and minima in the absorption spectra of water, lipid, and collagen to enable optical imaging at increased depths in tissue, compared with visible and NIR-I emitting fluorophores. We have been investigating rare-earth-doped nanomaterials, which simultaneously provide both SWIR and visible emissions under continuous wave near-infrared (980 nm) excitation. Although rare-earth-based probe development has gained significant traction in recent years, there is currently a lack of platforms available for imaging these nanomaterials across the whole animal to subcellular scales.

There are several commercial small animal imaging platforms designed for imaging fluorescent and bioluminescent probes, which emit in the visible and NIR-I regions. To the best of our knowledge, there are no commercial small animal whole-body imagers that can work with probes that emit at longer wavelengths, such as the rare-earth-doped nanomaterials used here. For high-resolution imaging, commercial epifluorescence microscopes are typically configured for excitation with UV/visible wavelengths and imaging of longer wavelength emissions. Upconversion light from rare-earth nanocomposites is at shorter wavelengths than the near-infrared excitation. Also, point-scanning confocal or multiphoton microscopes are poorly suited to imaging rare-earth probes because of the unusually long emission lifetime of these materials (milliseconds) compared with conventional fluorescent dyes (nanoseconds). The short pixel dwell time of these scanning microscopes (on the order of 100 ns) results in very little of the rare-earth emission being collected due to its long emission lifetime.

To overcome these limitations, we describe the design, assembly, and characterization of two platforms for rare-earth-based nanoprobe imaging. The first is a preclinical in vivo imaging system using SWIR emissions for deep tissue imaging. The second is an integrated optical coherence tomography (OCT) and multichannel line-scanning confocal microscopy system for high-resolution imaging of ex vivo tissues. This system uses backscattered SWIR light for OCT and visible upconversion emissions for confocal imaging. We demonstrate the capabilities of these systems by imaging rare-earth-doped nanoprobe administered to a murine model of metastatic breast cancer.

2 Methods

2.1 Wide-Field Short-Wave Infrared Imaging System

A fully automated system was developed for in vivo SWIR imaging in small animals [Figs. 1(a) and 1(b)]. Compared with our previous generation system, the spatial resolution of the system reported here is 4× greater, animal imaging is performed by automated scanning of the illumination beam, the emission filter combination has over 3× higher transmission, and real-time overlay of rare-earth signal has been implemented. The distal tip of a multimode fiber-coupled 980-nm diode laser (Opto Engine LLC) was mounted on a dual-axis motorized translation stage (Zaber, X-LSM-E) and the output beam collimated to a 9.6 mm (1/e²) diameter (Thorlabs, F810SMA-780). An InGaAs camera with 512 × 640 pixels, 25-μm square, (Sensors Unlimited, 640HSX) was fitted with a 25-mm fixed focal length lens (StingRay Optics, SR0907, f/1.4), providing a 195 mm × 155 mm field-of-view at a 300-mm working distance [Fig. 1(e)]. The translation stage raster scans the illumination beam across the sample platform while the camera continuously transfers images to a CameraLink frame grabber (NI PCIe-1427) at 30 frames per second. A custom LabVIEW VI creates an image from SWIR emissions by retaining the maximum intensity value at each pixel as images are acquired. After the scan is complete, a 1550-nm LED (Thorlabs, LED1550L) is turned on and a single full-field reflectance image is acquired for coregistration.
and overlay of SWIR emissions with anatomical features. Two gold mirrors are mounted at 45 deg on either side of the sample platform. An 805-nm short-pass dichroic mirror (Thorlabs, DMSP805, λ = 1325 nm) is positioned as close as possible to the galvo-mounted mirror. A 150-mm focal length achromatic cylindrical lens (Thorlabs, L2) forms a focused illumination line at the back aperture of the objective lens, which is important when imaging rare-earth-doped nanoparticles due to their long emission lifetimes.9 The OCT source (Thorlabs, SLD1325), λ = 1325 nm, ΔFWHM = 100 nm) is collimated to a 3.4-mm diameter beam (1/e²) using a 19-mm focal length achromatic doublet lens (Thorlabs, L3). Confocal illumination light from a 980-nm single-mode fiber-coupled laser (QPhotonics, QFBGLD-980-500) is collimated to an 8.0-mm diameter by a 36.6-mm focal length achromatic cylindrical lens (Thorlabs, L2) forms a focused illumination line at the back aperture of the objective lens, which is positioned as close as possible to the galvo-mounted mirror. A dichroic mirror (Semrock, FF495-Di03, D1), with a reflection band at the OCT source wavelength combines the illumination beams along a coaxial path. Both beams are then reflected by an 805-nm short-pass dichroic mirror (Thorlabs, DMSP805, D2) used to separate the nanoparticle excitation and (upconversion) emission wavelengths. To additionally allow for tissue autofluorescence excitation, a single-mode fiber-coupled laser with 488-nm wavelength (Blue Sky Research, FiberTec II) was collimated to an 8.0-mm diameter using a 35-mm focal length aspheric achromatic lens (Edmund Optics, L5) and then reshaped by a 150-mm focal length achromatic cylindrical lens (Thorlabs, L6). This beam is reflected by a 495-nm long-pass dichroic mirror (Semrock, D3), which transmits autofluorescence and rare-earth upconversion emissions. All three illumination beams are reflected at the galvo mirror. A 20×/0.4 objective lens (Olympus) was used for confocal imaging, whereas a 40×m focal length achromatic doublet lens (Thorlabs) was used for OCT. Erbium-doped nanoparticle emissions were transmitted through a 519- to 700-nm band-pass filter (Semrock) and focused by a 200-mm focal length achromatic doublet lens (Thorlabs, L4) onto a line-scan camera (Aviiva, EM1) with a single row of 1024 pixels, each 14-μm wide and 28-μm tall. The height of the pixels was ∼0.76 Airy units; therefore, no additional confocal slit was used. Backscattered OCT light returned through the sample arm fiber to the detection spectrometer, which was configured with a 1200 l/mm transmissive diffraction grating (Wasatch Photonics) and high-speed InGaAs line scan camera (Sensors Unlimited, LDH-2) as described previously.10

A 1951 USAF resolution target was used to measure a confocal lateral resolution of 2.8 μm. The full width at the half maximum (FWHM) of the mean pixel intensity over the central region of a mirror scanned through the focus indicated an optical sectioning thickness of 11.3 μm with the 20×/0.40 objective. The axial resolution of the OCT system was measured to be 14 μm in air over a 2.0-mm imaging depth, by setting a mirror at the focus of the OCT imaging beam and adjusting the reference path length over the imaging range. The OCT system resolved group 5, element 3 on a 1951 USAF resolution target with 34.7% contrast, corresponding to a lateral resolution of 27.9 μm. These values can be compared with theoretical figures of 7.7-μm axial resolution, 3.0-mm imaging depth, and lateral resolution of 19.8 mm.

2.2 Optical Coherence Tomography/Confocal Microscopy Imaging System

For high-resolution imaging, the sample arm of a spectral-domain OCT system was integrated within a dual-channel line-scanning confocal microscope (Fig. 2). Line scanning allows imaging with pixel exposure times far longer than point-scanning systems, which is important when imaging rare-earth-doped nanoparticles due to their long emission lifetimes.9 The OCT source (Thorlabs, SLD1325, λ = 1325 nm, ΔFWHM = 100 nm) is collimated to a 3.4-mm diameter beam (1/e²) using a 19-mm focal length achromatic doublet lens (Thorlabs, L3). Confocal illumination light from a 980-nm single-mode fiber-coupled laser (QPhotonics, QFBGLD-980-500) is collimated to an 8.0-mm diameter by a 36.6-mm focal length achromatic cylindrical lens (Thorlabs, L2) forms a focused illumination line at the back aperture of the objective lens, which is positioned as close as possible to the galvo-mounted mirror. A dichroic mirror (Semrock, FF495-Di03, D1), with a reflection band at the OCT source wavelength combines the illumination beams along a coaxial path. Both beams are then reflected by an 805-nm short-pass dichroic mirror (Thorlabs, DMSP805, D2) used to separate the nanoparticle excitation and (upconversion) emission wavelengths. To additionally allow for tissue autofluorescence excitation, a single-mode fiber-coupled laser with 488-nm wavelength (Blue Sky Research, FiberTec II) was collimated to an 8.0-mm diameter using a 35-mm focal length aspheric achromatic lens (Edmund Optics, L5) and then reshaped by a 150-mm focal length achromatic cylindrical lens (Thorlabs, L6). This beam is reflected by a 495-nm long-pass dichroic mirror (Semrock, D3), which transmits autofluorescence and rare-earth upconversion emissions. All three illumination beams are reflected at the galvo mirror. A 20×/0.4 objective lens (Olympus) was used for confocal imaging, whereas a 40×m focal length achromatic doublet lens (Thorlabs) was used for OCT. Erbium-doped nanoparticle emissions were transmitted through a 519- to 700-nm band-pass filter (Semrock) and focused by a 200-mm focal length achromatic doublet lens (Thorlabs, L4) onto a line-scan camera (Aviiva, EM1) with a single row of 1024 pixels, each 14-μm wide and 28-μm tall. The height of the pixels was ∼0.76 Airy units; therefore, no additional confocal slit was used. Backscattered OCT light returned through the sample arm fiber to the detection spectrometer, which was configured with a 1200 l/mm transmissive diffraction grating (Wasatch Photonics) and high-speed InGaAs line scan camera (Sensors Unlimited, LDH-2) as described previously.10

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2.3 Rare-Earth-Doped Nanocomposites

Rare-earth-doped nanocomposites were synthesized using a solvothermal decomposition method, forming a NaYF₄ core doped with ytterbium (Yb) and erbium (Er), surrounded by an undoped...
NaYF₄ shell.¹¹ Under continuous wave excitation at 980 nm, these particles emit fluorescence around 1550 nm, as well as upconversion light around 515, 540, and 650 nm. These nanocomposites (~10-µm diameter) were rendered biocompatible through encapsulation in human serum albumin¹² and functionalized with the chemokine receptor antagonist AMD3100, forming functionalized rare-earth-doped albumin nanocomposites (fREANCs), each ~100 nm in diameter and containing ~30 individual nanoparticles.¹² AMD3100 has been shown to preferentially bind to cells with high expression levels of CXCR4, a chemokine receptor overexpressed on many types of primary and metastatic cancer.¹³

2.4 Animal Model

All animal procedures were carried out under a protocol approved by the Rutgers University Institutional Animal Care and Use Committee. SCP28 cells (provided by Dr. Yibin Kang, Princeton University), derived from the MDA-MB-231 line and transfected with luciferase, were cultured in Dulbecco’s modified Eagle’s medium supplemented with 10% fetal bovine serum and 1% penicillin-streptomycin. Two 5-week-old female athymic nude mice (Taconic Laboratories) were inoculated with 300,000 SCP28 cells via tail vein injection, with two additional animals serving as healthy controls. Tumor uptake and progression were evaluated through weekly bioluminescence imaging (BLI) (Bruker, MS FX PRO). Each animal received an intra-peritoneal injection of D-Luciferin (Perkin Elmer, 150 mg/kg). Animals were then anesthetized under inhaled isoflurane at a concentration of 4%. Ten minutes later, animals were transferred to the BLI imaging chamber and imaged under general anesthesia with 1 to 2% isoflurane. BLI was performed using a 10-min exposure and 8 × 8 pixel binning. For each animal, once tumor uptake was confirmed on BLI, SWIR imaging was performed within 24 h.

2.5 Small Animal Short-Wave Infrared Imaging Procedures

For in vivo imaging, animals were anesthetized under 4% inhaled isoflurane maintained at a concentration of 1% to 2%. The collimated 980-nm laser beam [9.6 mm diameter (1/e²), 1.7 W power] was continuously scanned over a 100 mm × 50 mm field at 15 mm/s. Under these conditions, each individual point on the animal is illuminated for 640 ms resulting in a delivered fluence of 1.5 J/cm², less than half of the maximum permissible exposure under these conditions (3.6 J/cm²) defined by the American National Standards Institute (ANSI z136.1). The beam has a Gaussian intensity profile rather than a uniform distribution, so some overlap is required between successive sweeps of the raster scan to achieve uniform illumination. A lateral scan step of 2-mm results in 7.6-mm overlap between each sweep and a variation in illumination intensity of 26%. A smaller lateral step would reduce this variation but extend the scan time. Using the above parameters, each scan took ~3 min to complete. For ex vivo imaging, the lungs and liver from each animal were excised immediately following euthanization. Ex vivo imaging was performed using the same instrument parameters as for in vivo imaging.

Whole animal SWIR images were processed by manually outlining regions-of-interest (ROIs) corresponding to the liver and the lungs in the full-field reflectance images captured pre and post-nanoparticle injection. The mean SWIR pixel value in the pre-injection ROI was subtracted from each of the pixels in the post-injection ROI. These background-corrected pixel values were then displayed as a partially transparent overlay on the post-injection reflectance image with a user-defined colormap. With no pre-injection image of the organs available in the ex vivo cases, a region within the field-of-view that was not illuminated was used to estimate the background level.

2.6 Optical Coherence Tomography/Confocal Imaging of Biopsy Specimens

For each of the excised lungs, tissue samples were collected using a 2-mm-diameter punch biopsy, from regions with suspected tumor, and from the lungs of the healthy control animal. After OCT imaging of these biopsies, specimens were frozen at ~80°C in optimal cutting temperature compound for cryosectioning. About 60-µm-thick tissue sections were collected for confocal microscopy and 20-µm sections were collected and stained with hematoxylin and eosin (H&E). The line-scanning confocal microscope was used to examine the specimens and identify areas exhibiting the fREANC upconversion signal. Autofluorescence images of each field were obtained under 488-nm excitation.

3 Results

3.1 Wide-Field Short-Wave Infrared Imaging, In Vivo and Ex Vivo

Both tumor-bearing animals were imaged in vivo immediately following i.v. administration of fREANCs [Figs. 3(a) and 3(b), “0 h” panel]. One animal was euthanized immediately after imaging [Fig. 3(a)] and the other was reimaged 24 h later [Fig. 3(b), “24 h” panel], then euthanized. SWIR signal was evident in the abdominal region of both animals immediately after fREANC administration. This SWIR signal likely originated from both tumor-bound and free nanocomposites. At 24 h, this abdominal signal was reduced almost to the background level.

![Image](https://example.com/image.png)
level [Fig. 3(b), “24 h” panel], indicating that a substantial proportion of administered nanoparticles cleared the body within this time frame. SWIR images of the excised lungs (top organ) and liver (bottom organ) are shown in the right-hand panels of Figs. 3(a) and 3(b). The liver showed a higher mean signal than the lungs for the animal euthanized immediately following fREANC administration [Fig. 3(a)]. At 24 h, the in vivo liver [Fig. 3(b)] had lower SWIR intensity than the liver excised immediately after fREANC administration, whereas the lungs at 24 h exhibited a similar mean SWIR intensity to the “0 h” case.

The healthy control animal that did not receive fREANCs showed no significant SWIR autofluorescence signal in vivo or in organs excised immediately after imaging [Fig. 3(c)]. The control animal that received fREANCs showed abdominal signal on in vivo imaging immediately after fREANC administration [Fig. 3(d), “0 h”], with significantly reduced signal at 24 h. Ex vivo imaging of the lungs and liver of this control animal showed higher mean intensity in the lungs compared with the liver [Fig. 3(d)].

3.2 High-Resolution Tissue Imaging, Ex Vivo

Figure 4 shows representative confocal images (a,b) and coregistered OCT images (c,d) of tissue specimens collected at the study end point from the lungs of healthy control (a,c) and tumor bearing (b,d) animals. The rare-earth nanocomposite up-conversion signal is overlaid (red) on the tissue autofluorescence signal (green). Tissue from the healthy control animals contained confocal fields with a few small discrete regions of fREANC up-conversion signal [Fig. 4(a)]. Some tissue sections from the tumor bearing animal had high levels of discrete, localized fREANC up-conversion [e.g., Fig. 4(b)], with other sections appearing similar to healthy tissue, with few discrete regions of fREANC up-conversion. The OCT image from the corresponding normal tissue [Fig. 4(c)] has punctate regions of increased backscattering near the tissue surface, whereas the OCT image of the tumor tissue has a more uniform backscattering signal [Fig. 4(d)]. On H&E sections prepared from these specimens, alveolar structure is readily apparent in the healthy tissue [Figs. 4(e) and 4(f)] whereas sections from the tumor exhibit denser tissue with few, small alveoli [Figs. 4(g) and 4(h)].

4 Conclusions

This article describes the design and characterization of two imaging systems for evaluating rare-earth-doped nanocomposites in biological systems. Whole animal in vivo imaging and whole organ ex vivo imaging were implemented using scanned excitation light and an InGaAs camera. Ex vivo OCT imaging was integrated with dual-channel confocal microscopy for imaging tissue microstructure, autofluorescence, and nanoparticle distribution. The capabilities of these systems are illustrated by imaging albumin-encapsulated erbium-doped nanocomposites in a mouse model of metastatic breast cancer. We anticipate that this work will guide other researchers studying these emerging nanomaterials in the design and assembly of imaging platforms to suit their specific needs.

Disclosures

The authors have no relevant financial interests in this article and no potential conflicts of interest to disclose.

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