Non-Invasive Imaging of Cysteine Cathepsin Activity in Solid Tumors Using a $^{64}$Cu-Labeled Activity-Based Probe

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Abstract

The papain family of cysteine cathepsins are actively involved in multiple stages of tumorigenesis. Because elevated cathepsin activity can be found in many types of human cancers, they are promising biomarkers that can be used to target radiological contrast agents for tumor detection. However, currently there are no radiological imaging agents available for these important molecular targets. We report here the development of positron emission tomography (PET) radionuclide-labeled probes that target the cysteine cathepsins by formation of an enzyme activity-dependent bond with the active site cysteine. These probes contain an acyloxymethyl ketone (AOMK) functional group that irreversibly labels the active site cysteine of papain family proteases attached to a 1,4,7,10-tetraazacyclododecane-1,4,7,10-tetraacetic acid (DOTA) tag for labeling with $^{64}$Cu for PET imaging studies. We performed biodistribution and microPET imaging studies in nude mice bearing subcutaneous tumors expressing various levels of cysteine cathepsin activity and found that the extent of probe uptake by tumors correlated with overall protease activity as measured by biochemical methods. Furthermore, probe signals could be reduced by pre-treatment with a general cathepsin inhibitor. We also found that inclusion of a Cy5 tag on the probe increased tumor uptake relative to probes lacking this fluorogenic dye. Overall, these results demonstrate that small molecule activity-based probes carrying radio-tracers can be used to image protease activity in living subjects.

Introduction

Proteases play important roles in the regulation of both normal and disease processes. In particular, the papain family cysteine cathepsins are frequently over-expressed in a number of human cancers [1-7]. In addition, expression of a number of cysteine cathepsin including cathepsins B and L is increased in pre-neoplastic lesions and changes in both localization and sub-cellular distribution of these proteases are often observed in tumors [1,3,7]. This combination makes them potentially valuable cancer biomarkers [1–5,8,9]. Novel imaging methods and molecularly targeted tracers can now be used not only to locate a tumor, but also to visualize the expression and activity of specific molecular targets and biological processes in a tumor [10,11]. These imaging methods have the potential to facilitate both early disease detection and to aid in the process of drug development by allowing non-invasive monitoring of target inhibition. Directed targeting of enzymatic proteins such as proteases using imaging agents also has the potential to provide greater detail about the basic biological framework of a tumor and provide better resolution of the disease phenotype [10,12–16].

Since most proteases are initially synthesized as inactive zymogens that are activated by a complex set of post-translational mechanisms, tools that report on enzyme activity rather than protein abundance will be required to fully understand their function in complex biological processes. For this reason, a number of recent studies have focused on the development of imaging agents that act as substrates for a target protease [10,13,15,16]. In addition, fluorescent molecules that only penetrate a cellular membrane when processed by a target protease have been developed [17–19]. While all of these methods have provided valuable new tools, none of these methods have been translated for use in radiological imaging.

We report here the development and application of radio-labeled small molecule activity-based probes (ABPs) that can be used for positron emission tomography (PET) imaging of cysteine...
Generation of a PET Probe Based On the GB123 Structure

ABPs provide an indirect readout of protease activity in complex proteomes. A number of ABPs targeting various cysteine cathepsins have recently been reported [20,21,23,24]. In particular, the peptide acylxymethyl ketones have been validated in optical imaging applications [8,21,22,25]. The covalent modification of a cysteine protease by an AOMK results in alkylation of the active site thiol and loss of acyloxy group (Figure 1A). One of the initially reported AOMK derivatives, Z-FR-AOMK, exhibited high potency and specificity for cathepsin B and L [20]. Moreover, conversion of Z-FR-AOMK to the related Z-FK-AOMK and modification of the lysine side chain with bulky organic fluorescent dye resulted in a probe, GB123, that could be used to non-invasively image cathepsins B and L activity [8,21]. Therefore, we synthesized a probe in which the fluorescent reporter of GB123 was replaced by the 1, 4, 7, 10-tetraazacyclododecane-1, 4, 7, 10-tetraacetic acid (DOTA) for coordination of the $^{64}$Cu tracer (Z-FK(DOTA)-AOMK; Figure 1B). We also synthesized a probe in which the original carboxylbenzoyl (Z) capping group was replaced by the DOTA group and an additional phenylalanine amino acid was added to mimic the properties of the Z group that was lost (GB170; Figure 1B). Finally, we generated a control analog that lacked the reactive AOMK electrophile (GB173; Figure 1B).

We radiolabeled the Z-FK(DOTA)-AOMK, GB-173 and GB-170 conjugates with $^{64}$Cu ($t_{1/2}$ = 12.7 h, $E_{\text{max}}^* = 656 \text{ keV}$, 19%) at 50 $^\circ$C for 1 h and purified the resulting probes to greater than 95% radiochemical purity using analytical RP-HPLC (Figure S1). The specific radioactivity of $^{64}$Cu-Z-FK(DOTA)-AOMK was 0.48 Ci/μmol (17.8 MBq/μmol, 500 μCi/μg) at the end of synthesis (EOS). The specific radioactivity of GB170 and GB173 were determined to be around 20 μCi/μg EOS. These labeled compounds were then used for in vitro and in vivo labeling experiments.

Labeling properties of $^{64}$Cu labeled AOMK analogs

We tested probes in the human breast cancer cell line MBA-MB-435 and a mouse myoblastoma cell line that had been transformed by over-expression of the ras oncogene (C2C12/Ras) [26]. Both of these cell lines were originally used for in vivo imaging studies with the NIRF-labeled cysteine cathepsin probes [21]. We labeled both cell lines with GB123 and found that cathepsin B and L activities of the C2C12/Ras cell line were higher than the activities observed for the same targets in the MBA-MB-435 cells (Figure 2A). Labeling of the same cells with $^{64}$Cu-Z-FK(DOTA)-AOMK confirmed that the DOTA probes produced a similar labeling pattern to that observed for GB123 (Figure 2B). In addition, the specificity of the Cu-labeled probe for the cysteine cathepsins was confirmed by the complete loss of labeling of the protease targets when cells were pretreated with the general cysteine cathepsin inhibitor JPM-OEt [25]. Finally, we compared the labeling properties of the dual fluorescent and PET tagged probe GB170 and GB173 to the original PET-only probe Z-FK(DOTA)-AOMK. Both GB170 and GB173 have fluorescent tags and can be used to directly label intact NIH 3T3 cells. For comparison with the non-fluorescently labeled probe, we labeled residual activity using a BODIPY-TMR labeled version of GB123 (Figure 2C). This analysis confirmed that the GB170 probe was able to effectively label active cathepsins B and L. Interestingly, the dual labeled probe GB170 showed slightly reduced potency in the competition assay compared to Z-FK(DOTA)-AOMK and GB123 even though it showed the most effective labeling of cathepsins as measured by direct labeling. GB173, which lacks the AOMK reactive group, failed to label active cathepsins and did not compete for labeling by the BODIPY-TMR GB123, making it an ideal negative control probe.

Biodistribution of $^{64}$Cu-labeled probe in subcutaneous C2C12/Ras and MDA-MB-435 tumors

We next examined the overall biodistribution of the various $^{64}$Cu-labeled probes using athymic nude mice carrying subcutaneously grafted C2C12/Ras and MDA-MB-435 tumors. We measured uptake of Z-FK(DOTA)-AOMK in both the C2C12/Ras and MDA-MB-435 tumor models (Figure S2). This probe displayed rapid blood clearance, resulting in low blood and muscle uptake even at the early time points in both models. However, we did observe some accumulation of radioactivity in the C2C12/Ras tumors expressing high levels of cysteine cathepsins (0.35 ± 0.13%ID/g at 0.5 h p.i.). This activity remained in the tumors resulting in an uptake of 0.27 ± 0.05%ID/g at 24 h p.i. For the MDA-MB-435 tumors with lower cysteine cathepsin activity, tumor uptake was significantly lower at each time point relative to the C2C12/Ras tumors ($P<0.05$). Furthermore, while moderate tumor-to-background ratios (tumor/blood 1.25 and tumor/muscle 5.64 at 2 h p.i.) were observed in the C2C12/Ras tumors, only low tumor-to-background ratios (tumor/blood 0.61 and tumor/muscle 3.03 at 2 h p.i.) were found for the MDA-MB-435 tumors. For both tumor models, $^{64}$Cu-Z-FK(DOTA)-AOMK displayed very low accumulation in most non-tumor tissues. The highest radioactivity was found in the liver, consistent with high endogenous expression of various cysteine cathepsins in this organ. Moderate renal accumulation was also observed at all times in both tumor models. These high levels of non-tumor signals may be partially due to loss of the copper radiotracer from the probe. This is an issue when using DOTA as a chelation agent. Subsequent studies will be aimed at using more optimal chelators and possibly other radio-tracers that can be secured to the probe through covalent modification. Regardless, the relatively low tumor accumulation of $^{64}$Cu-Z-FK(DOTA)-AOMK suggested that this probe is not optimal for in vivo applications.

For comparison, we measured the biodistribution of GB170 as well as its corresponding negative control GB173 in the C2C12/Ras tumor model (Figure S2). GB170 rapidly accumulated in the C2C12/Ras tumors ($4.28 \pm 0.91%ID/g$, with higher levels of probe accumulation throughout all tissues compared to Z-FK(DOTA)-AOMK. The control probe GB173 showed some tumor accumulation ($1.96 \pm 0.73%ID/g$) but this level was significantly ($P<0.05$) lower than the levels observed for GB170. Furthermore, we observed moderate to high tumor-to-background activity. These probes are based on the peptide acylxymethyl ketones (AOMKs) that have been reported to be highly selective labels of a number of classes of cysteine cathepsins [20]. This class of reagents has also recently been used for optical imaging of cysteine cathepsin activity in live cells [8] and in near-infrared fluorescent (NIRF) labeled form for non-invasive optical imaging of cysteine cathepsin activity in living subjects [21,22].
ratios (tumor/blood 3.1 and tumor/muscle 13.0 at 24 h p.i.) for GB170 in the C2C12/Ras tumors, while only low tumor-to-background ratios (tumor/blood 1.84 and tumor/muscle 4.4 at 24 h p.i.) for GB173.

MicroPET imaging of cathepsin activity in tumor models

We performed microPET imaging of mice bearing the MDA-MB-435 and C2C12/Ras tumors using the 64Cu-Z-FK(DOTA)-AOMK probe (Figure 3). Although the probe signal in the liver and kidney is relatively high, the C2C12/Ras tumors could be clearly visualized with good tumor to contralateral background contrast of 2 at 24 h p.i. (Figure 3B). In contrast, we observed low tumor uptake and poor tumor to normal tissue contrast in the MDA-MB-435 tumors (Figure 3A). Quantification of the accumulation of the probe in the tumor and contralateral muscle tissue also showed that 64Cu-Z-FK(DOTA)-AOMK had higher tumor uptake and a higher tumor-to-muscle ratio in C2C12/Ras than in MDA-MB-435 (Figure 3B).

We next performed microPET imaging of mice bearing C2C12/Ras tumors using 64Cu-GB170 and the corresponding negative control, 64Cu-GB173 (Figure 4A). As expected, GB170 produced substantially higher signals that the 64Cu-Z-FK(DOTA)-
AOMK probe with tumor to contralateral background contrast of 4 at 24 h p.i. In addition, $^{64}$Cu-GB173 showed only weak tumor uptake and poor tumor to normal tissue contrast. Specific probe uptake into tumors was confirmed by quantification of the tumor and muscle signals (Figure 4B) and by comparison of tumor to muscle and tumor to blood ratios (Figure 4C).

While these data were suggestive that probe accumulation correlates with levels of active cathepsins in the tumor tissues, we needed to demonstrate this correlation by direct analysis of cathepsin activity in samples after imaging. Therefore, we performed a study in which three different cell lines were used to generate xenograft tumors on nude mice. We chose the original MDA-MB-435 cells with low cathepsin activity and used the C2C12/Ras cells with high cathepsin levels. We also included a third tumor cell line (4T1) that also has high expression of cathepsins. We then performed PET imaging studies at various time points using the GB170 probe (Figure 5A). After quantification of tumor and muscle signals and subsequent biodistribution studies of collected tissues (Figure S3), we lysed the tumor tissues and directly labeled target cathepsins by probes was obtained by SDS-PAGE followed by scanning of the gel using a flatbed laser scanner (Top gel). Inhibition of the same targets was observed by scanning of the gel for the BODIPY labeled probe using a Typhoon 9410 scanner (Bottom gel).

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Figure 3. In vivo imaging of tumor-derived cathepsin activity using first generation activity-based probes. (A) Decay corrected coronal (top) and transaxial (bottom) microPET images of a nude mouse bearing MDA-MB-435 tumor at 1, 4 and 28 hours after tail vein injection of (5.55 MBq, 150 μCi) $^{64}$Cu-Z-FK(DOTA)-AOMK. The location of the tumor is indicated by arrows. Average values (n = 3) for tumor uptake (%ID/g ± SD) relative to muscle signals at each time point are plotted (at right) (B) Decay corrected coronal (top) and transaxial (bottom) microPET images of a nude mouse bearing C2C12/Ras tumors at 2, and 24 hr after administration of (3.7 MBq, 100 μCi) $^{64}$Cu-Z-FK(DOTA)-AOMK. Arrows indicated location of tumors. Average values (n = 3) for tumor uptake (%ID/g ± SD) relative to muscle signals at each time point are plotted (at right). For all images (T: tumor; L: liver; B: bladder).

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Figure 4. In vivo imaging of tumor-derived cathepsin activity using dual optical/PET imaging probes. (A) Coronal and transaxial decay corrected microPET images of an athymic nude mouse bearing C2C12/Ras tumor at different time points after tail vein injection of (1.85 MBq, 50 μCi) $^{64}$Cu-GB170 (bottom) or $^{64}$Cu-GB173 (top). The location of the tumor is indicated by arrows. (B) Tumor-to-normal tissues ratios of $^{64}$Cu-GB170 or $^{64}$Cu-GB173 in C2C12/Ras cancer bearing mice at different time points (n = 3 each). (C) Plots of average ratios (n = 3) of tumor to blood and tumor to muscle signals at the indicated time points. For all images (T: tumor; L: liver; B: bladder).

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measured by PET imaging, indicated a strong correlation between the two set of values (Figure 5D). Specifically, the two high expressors of cathepsins (4T1 and C2C12/Ras) showed indistinguishably (p = 0.7) high levels of cathepsin activity as measured by gel, while the tumors with low cathepsin activity (MDA-MB-435) showed significantly (p<0.05) reduced levels of cathepsins relative to either of the high cathepsin tumors. These levels matched the relative levels of probe observed in tumors by PET imaging.

Figure 5. Probe signals in tumors correlate with levels of active cathepsins. (A) Representative coronal and transaxial decay corrected microPET images of athymic nude mice bearing 4T1, MDA-MB-435 and C2C12/Ras tumors at different time points after tail vein injection of (1.85 MBq, 50 μCi) 64Cu-GB170. Locations of tumors (T) and liver (L) are indicated. (B) SDS-PAGE and fluorescence scanning of tumor lysates from mice in (A) after the final 24 hr time point. The positions of fluorescently labeled cathepsins are indicated. (C) Determination of residual cathepsin activity in tumors from mice in (A) as measured by labeling of lysates with GB123 followed by SDS-PAGE and scanning of the gel with a flatbed laser scanner. (D) Comparison of the levels of probe levels in tumor relative to muscle as measured by PET (right bar graph) to the levels of active cathepsins as measured by quantification of gels in (C; left bar graph). The p-values are indicated and defined as follows: n.s. = not significant, *<0.05, **<0.01, ***<0.001.

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suggesting that probe signal is the result of retention at sites of high cysteine cathepsin activity.

Finally, to confirm that the probe accumulation was due to modification of active cathepsins, we pre-treated C2C12/Ras tumor bearing mice with a broad-spectrum inhibitor of the cysteine cathepsins (Figure 6). This compound, K11777 was used in previous optical imaging studies to pre-block cathepsin activity [21]. MicroPET images show a marked drop in probe accumulation in tumors of mice treated with the inhibitor compared to those treated with the vehicle control (Figure 6A). Further quantification of tumor signals in vehicle and drug-treated animals compared to muscle signals confirmed the drop in specific signal as a result of inhibition of cysteine cathepsin activity (Figure 6B). Thus, virtually all of the probe signal observed above muscle background could be blocked by inhibition of the target proteases.

Discussion

The cysteine cathepsins are potentially valuable molecular targets for tumor imaging because of their significant function in tumor progression, invasion and metastasis [1-7]. Although NIRF labeled probes that target this class of protease have been successfully applied to optical imaging applications [10,13,15,16,21,22], radiolabeled probes have not been developed. Part of the reason for the lack of probes for radiological imaging applications is the difficulty in designing radioactive probes based on protease substrates. Therefore, in order to prepare new PET agents, we chose to make use of highly selective activity-based probes that covalently target proteases. The main disadvantage of using an inhibitor-based approach is the lack of amplification of signal that is obtained with substrate-based approaches. However, the high expression of cysteine cathepsins in tumor tissue [27], and the prior validation of these probes in optical imaging applications suggested that they would be potentially valuable for use in PET imaging.

We chose to use the validated peptide AOMK scaffold for development of our first generation PET probe. Simple conversion of the Z-FK-AOMK probe to the corresponding DOTA labeled analog Z-FK(DOTA)-AOMK (Figure 1B) allowed labeling with $^{64}$Cu. We selected $^{64}$Cu as a PET radiolabel because it can be readily produced using a medical cyclotron and the intermediate half-life of $^{64}$Cu makes it suitable for small molecule and peptide radiolabeling [28-30]. Overall labeling and biodistribution studies with $^{64}$Cu-Z-FK(DOTA)-AOMK indicated that the probe showed rapid clearance in blood but accumulated in tumor tissues with reasonable signal levels (Figure 3). Unfortunately, the probe showed overall low tumor uptake (only 0.33±0.13%ID/g at 0.5 h p.i. in C2C12/Ras tumor model), possibly due to its lipophilicity (Figure S1) and loss of the fluorogenic Cy5 group found on the corresponding optical imaging probe. We have found that the addition of bulky aromatic groups on probes enhances their labeling of lysosomal proteases, possibly due to increased endocytosis. We therefore synthesized a PET probe in which the Cy5 fluorophore was retained on the P1 lysine sidechain and the DOTA tag for PET labeling was moved to the N-terminus of the peptide. Analysis of the GB170 probe in microPET imaging studies confirmed that the in vivo tumor uptake was clearly superior to $^{18}$F-Z-FK(DOTA)-AOMK in C2C12/Ras tumor bearing mice. We were also able to confirm that the signals observed for the GB170 probe were due to retention at the site of active cysteine cathepsins by showing correlation between imaging signals and protease activity as measured biochemically (Figure 5) as well as by showing a reduction in probe retention in mice pretreated with a broad-spectrum inhibitor of the cathepsins (Figure 6). These combined data confirm the selectivity of the tumor signals and also demonstrates the utility of PET probes for monitoring in vivo efficacy of small molecule drugs.

Although the probes reported here support the use of ABPs for PET imaging, significant work will be required to improve the probes. For example, release of free $^{64}$Cu from the radiolabeled compound or transchelation to other proteins may also be partially responsible for high background signal. Other chelators such as cross-bridged cyclam ligands may also be used in place of DOTA to improve the metal-chelate stability and subsequently improve the biodistribution of the probes [30,31]. Regardless of the issues of probe background, the use of positron emitting labels allows application of tomographic methods to resolve signals in specific locations that are free from high protease background. In addition, unlike the NIRF versions of these ABPs the PET probes have the potential to be used to image tissues at much greater depths and with higher overall resolution [21]. The work presented here with $^{64}$Cu-labeled probes serves as the foundation for future studies using the more clinically suitable $^{18}$F tracer. We believe that cathepsin-targeted PET imaging agents should find broad applications once these agents become available to the nuclear medicine community.

Figure 6. Inhibition of tumor-derived cathepsin activity can be visualized non-invasively using PET probes. Coronal and transaxial decay corrected microPET images of an athymic nude mouse bearing MDA-MB-231MFP tumors treated with the general cathepsin inhibitor K11777 (treated) or vehicle (control) followed by $^{64}$Cu-GB170 (1.85 MBq, 50 μCi). Images were taken at the indicated times after injection of the imaging probe. The location of the tumor is indicated by arrows. (B) Quantification of average levels (n = 3) of probe uptake in tumors at the various time points for control treated and inhibitor treated mice relative to uptake into muscle tissue. For all images (T: tumor; L: liver; B: bladder). doi:10.1371/journal.pone.0028029.g006
Materials and Methods

Ethics Statement

All animal experiments were conducted according to relevant national and international guidelines and approved by the Stanford Institutional Animal Care and Use Committee (IACUC; approval number A3213-01). All experiments strictly followed the panel’s specific guidelines regarding the care, treatment and euthanasia of animals used in the study.

General

1,4,7,10-tetraazacyclododecane-1,7,10-tetraacetic acid (DOTA) was obtained from Macrocyclics Inc. (Richardson, TX). All other chemicals were obtained from Sigma-Aldrich Chemical Co. (St. Louis, MO). AOMK analogs, were prepared based on a reported synthetic method [20]. 64Cu was provided by the Department of Medical Physics, University of Wisconsin at Madison (Madison, WI). A CRC-15R PET dose calibrator (Capintec Inc., Ramsey, NJ) was used for all radioactivity measurements. Reverse phase high performance liquid chromatography (RP-HPLC) was performed on a Dionex Summit HPLC system (Dionex Corporation, Sunnyvale, CA) and analytical (Dionex, Sunnyvale, CA. Acclaim120 C18, 4.6 mm×250 mm) RP-HPLC columns were used. The mobile phase was solvent A, 0.1% trifluoroacetic acid (TFA)/H2O, and solvent B, 0.1%TFA/acetonitrile. Matrix-assisted laser desorption/ionization time of flight mass spectrometry (MALDI-TOF-MS) were performed on a Perseptive Voyager-DE RP Biospectrometry instrument (Framingham, MA) by the Stanford Protein and Nucleic Acid Biotechnology Facility. Alpha cyano-4-hydroxy-cinnamic acid (α-CHCA, prepared as 10 g/L in 33.3% CH3CN : 33.3% EtOH : 33.3% H2O : 0.1% TFA) were used as the solid matrix. NIH3T3 mouse fibroblast cells was a generous gift from Dr. P. Jackson, Stanford University, CA. The tumorigenic murine skeletal myoblast cell line C2C12/Ras (retrovirally transduced with the ras oncogene) was a generous gift from Dr. Helen Blau, Stanford University, CA, human breast cancer MDA-MB-435 was a generous gift from Dr. X. Chen, Stanford University, CA. NIH3T3 cells were treated similarly as described above but were labeled with GB111-TMR-X (1 μM final concentration) after probe treatment and then lysed. Cells were scanned for fluorescence at 532/580 nm with a Typhoon scanner.

Subcutaneous Tumor Model

All animal studies were carried out in compliance with Federal and local institutional rules for the conduct of animal experimentation. Female athymic nude mice (nu/nu) were obtained from Charles River Laboratories (Boston, MA) at 7–8 weeks old and kept under sterile conditions. The nude mice were inoculated subcutaneously in the right shoulder with 1×106 cultured C2C12/Ras cells or 5×106 MDA-MB-435 cells or 3×106 cultured 4T1 cells. For inhibition studies, 3×106 MDA-MB-231 MFP cells were used. When the tumors reached 0.5–0.8 cm in diameter, the tumor bearing mice were used for in vivo PET imaging studies (see below).

MicroPET Imaging

PET imaging of tumor-bearing mice will be performed on a microPET R4 rodent model scanner (Siemens Medical Solutions USA, Inc., Knoxville, TN). The mice bearing C2C12/Ras and MDA-MB-435 tumors were injected with 740 KBq (20 μCi) 64Cu AOMK probe via the tail vein. At different times after injection, 5 or 10-min static scans were obtained and the images were reconstructed by a two-dimensional ordered subsets expectation maximum (OSEM) algorithm. Regions of interest (ROIs) were then drawn over the tumor on decay-corrected whole-body coronal images. The mean counts per pixel per minute were obtained from the ROI and converted to counts per milliliter per minute by using a calibration constant. By assuming a tissue density of 1 g/mL, the ROIs were converted to counts/g/min. An image ROI-derived %ID/g of tissue was then determined by dividing counts per gram per minute with injected dose (ID). No attenuation correction was performed.

Biodistribution Studies

Mice bearing xenografts were injected with 740 KBq (20 μCi) of 64Cu labeled tracer through the tail vein and sacrificed at different time points after injection. Tumor and normal tissues were removed and weighed, and radioactivity was measured by gamma-counter. The radioactivity uptake in the tumor and normal tissues was expressed as a percentage of the injected radioactive dose per gram of tissue (%ID/g). To inhibit the in vivo
cathepsin activity, K1177 was injected intraperitoneally at the dose of 100 mg/kg/day in 40% DMSO/sterile PBS in a final volume of 100 μL for 5 days before the imaging sessions [21].

Biochemical analysis of cathepsin activity in tumors

Tumors were removed after the 24 hr time point and homogenized in citrate buffer (50mM Citrate pH 5.5, 5 mM DD1, 0.5% CHAPS, 0.1% Triton X1). Total protein (40 μg) was labeled with GB123 (1 μM final concentration) for 1 hr at 37°C. The proteins were resolved on SDS-PAGE (15%). Labeled proteases were visualized by scanning the gel with a Typhoon 9410 imager (excitation/emission 633/630 nm) [GE Healthcare, Piscataway, NJ]. Labeling intensities were quantified using Image J software.

Statistical Method

Statistical analysis was performed using the Student’s t-test for unpaired data. A 95% confidence level was chosen to determine the significance between groups, with P < 0.05 being significantly different.

Supporting Information

Figure S1 Characterization and purification PET probes. HPLC radiochromatogram of purified (A) 64Cu-Z-FK(DOTA)-AOMK, (B) 64Cu-GB170, and (C) 64Cu-GB173. (DOC)

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Figure S2 Biodistribution of probes in vivo. Biodistribution results of 64Cu-Z-FK(DOTA)-AOMK in (A) C2C12/Ras (top) or MDA-MB-435 (bottom) cancer bearing mouse models as well as for (B) 64Cu-GB170 (top) and 64Cu-GB173 (bottom) in C2C12/ras tumor bearing mice. Data are expressed as the percentage administered activity (injected dose) per gram of tissue (%ID/g) after intravenous injection of 740 kBq (20 μCi) of 64Cu-Z-FK(DOTA)-AOMK at 0.5, 2, and 24 h pi (n = 3). Significant lower tumor uptake and tumor/blood, tumor/muscle ratio in MDA-MB-435 breast cancer (P < 0.05) were observed. (DOC)

Figure S3 Biodistribution of 64Cu-GB170 in tumor bearing mice. N = 4 for each. (DOC)

Methods S1 Chemistry and Radiochemistry. (DOC)

Author Contributions

Conceived and designed the experiments: GR GB MV HL LEE SSG MB ZC. Performed the experiments: GR GB MV HL SS LEE OG ZM HJ. Analyzed the data: GR GB MV HL LEE SSG MB ZC. Wrote the paper: GR GB MV HL MB ZC.
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