A Dual Reporter Iodinated Labeling Reagent for Cancer Positron Emission Tomography Imaging and Fluorescence-Guided Surgery

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INTRODUCTION

Cancer is a leading cause of death worldwide, accounting for 8.8 million deaths in 2015. About 14 million people are diagnosed with cancer each year, and this figure is expected to reach 27 million by 2030 according to the World Health Organization.¹ The combination of early diagnosis and complete surgical removal represents the most effective therapeutic modality for malignant tumors.²

Benign diseases such as inflammation or infection are also associated with increased ¹⁸F-FDG uptake, resulting in false-positive diagnosis.³ ImmunoPET employing positron emitter-labeled tumor-specific monoclonal antibodies has shown great promise to stage cancer patients and detect tiny tumor lesions in several clinical trials thanks to their nanomolar binding affinity and specificity for tumor-associated biomarkers.⁴ Zirconium-89 (23% $\beta^+$, $T_{1/2} = 3.3$ days) and iodine-124 (24% $\beta^+$, $T_{1/2} = 4.2$ days) are the most commonly used positron emitters in immunoPET due to their longer half-lives that are compatible with the biological half-lives of the full-size antibodies.⁵ In a feasibility study, $^{89}$Zr-trastuzumab targeting...
both of ccRCC with a hand-held PET probe capable of measuring preoperative mapping with PET and intraoperative localization has been approved by the Food and Drug Administration (FDA).12

ovarian cancer patients.10 The surge in the human epidermal growth factor receptor 2 (HER2) allowed the resolution of the hand-held radiation detection device, it is challenging to distinguish the surgical margins and to identify tissues. In contrast, the interface between the tumor and the surrounding healthy tissues. In contrast, fluorescence imaging offers superior spatial resolution. Ntziachristos et al. demonstrated that a folate receptor-targeting fluorescein molecular probe offered specific and sensitive real-time identification of tumor deposits with clear surgical margins during fluorescence-guided surgery in ovarian cancer patients.10

To harness the synergistic property of PET and fluorescence imaging for cancer diagnosis and precise tumor resection, considerable effort has been made to develop dual PET and fluorescence imaging reagents.11–13 However, none of these has been approved by the Food and Drug Administration (FDA).12

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One limiting factor is the lack of a generic method to prepare these reagents for screening.13 We envisage that a dual PET and fluorescent labeling reagent would act as a generic tool to install both a PET and a fluorescent reporter to any cancer targeting antibody. Thus, it would standardize the preparation protocol, accelerate the screening process, and minimize the translational hurdle.

Recently, we reported a copper-mediated one-pot three-component radiiodination method that allowed the reaction of functionalized alkynes, azides, and radioactive iodine-125 (γ, T1/2 = 60 days) to form trisubstituted iodotriazoles.14,15 To explore the potential of this chemistry in combinational cancer diagnosis and surgery, we have synthesized a new dual modality labeling reagent, 124I-Green. It is equipped with (i) a positron emitter, iodine-124, for PET imaging of cancer, (ii) a clinically approved green fluorescent reporter, fluorescein, for fluorescence-guided surgery, and (iii) a succinimide moiety for antibody coupling (Scheme 1). Here, we exemplify the utility of this approach to allow the one-step installation of a dual PET and fluorescence reporter to a carcinoembryonic antigen (CEA) specific marine antibody ASB7. The dual labeled ASB7 offered excellent tumor-to-background contrast in both PET/CT and ex vivo fluorescence imaging.

RESULTS

Synthetic Chemistry and Radiolabeling. Initially, S(6)-carboxyfluorescein was treated with N,N-diisopropylethylamine in DMF to form the corresponding lactone and then reacted with the carboxylic group with propargylamine using propylphosphonic anhydride as the coupling reagent at 0 °C to form the N-propargyl-S(6)-carboxyfluoresceinamide in 30% yield. The 4-

Scheme 1. Synthesis of the Dual PET and Fluorescent Labeling Reagent, 124I-Green

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<th>entry</th>
<th>ligand (10 mol %)</th>
<th>temperature (°C)</th>
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<td>bathophenanthroline</td>
<td>92 ± 5</td>
<td>(n = 4)</td>
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Table 1. Optimization of the Radioiodination of the Dual Labeling Reagent

“Suspension of N-propargyl-S(6)-carboxyfluoresceinamides (1.0 μmol), CuCl2 (1.0 μmol) TEA (1.5 μmol) and ligand (mol %) was added to a mixture of azidomethyl-N-succinimidyl benzoate (1.0 μmol) and [124I]NaI (6 μL). Radiochemical yield was determined by HPLC. [124I]NaI was used for the reaction. TEA-HCl was used to neutralize the NaOH in the aqueous [124I]NaI.
azidomethyl-N-succinimidyl benzoate was also prepared following a literature method in two steps with an overall yield of 61%.

Subsequently, the nonradioactive reference compound of 124I-Green was synthesized from the reaction of N-propargyl-5(6)-carboxyfluoresceinamide, 4-azidomethyl-N-succinimidyl benzoate, and N-iodosuccinimide in the presence of copper(I) iodide in DMF with a moderate yield of 37% (Scheme 1). Next, we systematically screened different conditions to optimize the radiochemical preparation of 124I-Green. [125I]NaI was used as the surrogate of [124I]NaI for the optimization of the radiolabeling. Under the previously published conditions, low radiochemical yields (RCYs) of 4% and 40% measured by HPLC were observed at room temperature (rt) and 60 °C, respectively (Table 1, entries 1 and 2). Screening of 1,10-phenanthroline copper ligands revealed bathophenanthroline to be optimal for the reaction and provided the 125I-labeled target compound in excellent RCYs of 91 ± 4% measured by HPLC (n = 6) (Table 1, entry 6). When these optimized conditions were applied to the 124I-radiolabeling, excellent RCYs of 92 ± 5% (n = 4) measured by HPLC were achieved in the presence of TEA·HCl (Table 1, entry 10, and Supporting Information, Figure S1). The identity of 124I-Green was proved by co-eluting with its nonradioactive reference compound (Supporting Information, Figure S2).

When prepared under the optimized conditions starting with around 36 MBq of [124I]NaI, 124I-Green was isolated in radioactivity yields of 78 ± 6% (n = 3) with the radiochemical purity ≥95% (Supporting Information, Figure S2) and the molar activity around 3.5 GBq/μmol.

**Dual Labeling Reagent Antibody A5B7 Conjugation and Characterization.** Next, we conjugated 124I-Green with a murine CEA specific antibody A5B7. The A5B7 was incubated with 124I-Green in 10 mM pH 9.0 carbonate buffer at rt for 45 min. Good labeling efficiency of 60 ± 5% (n = 3) was achieved (Supporting Information, Table S1, entry 1) after size exclusion column purification. The recovery efficiency of 124I-Green A5B7 conjugate from this size exclusion column was also determined as 80 ± 4% (n = 3) by passing the purified 124I-Green labeled A5B7 conjugate through a new size exclusion column. To determine the optimal number of fluorophores per antibody, we co-incubated the A5B7 with a mixture of 124I-Green and different amount of the nonradioactive reference compound in 2, 5, 10, and 20 equiv of the antibody in 10 mM pH 9.0 carbonate buffer at rt for 45 min. The radiolabeling efficiencies were approximately 45%, 34%, 27%, and 25% (n = 3), respectively. The average fluorophore per antibody was then determined as 1, 2, 3, or 6, respectively (corrected for the loss of 124I-Green A5B7 conjugate on the size exclusion column) (Supporting Information, Table S1 entries 2–5). Subsequently, the fluorescence intensity of the dual labeled A5B7 (50 μg) with 1, 2, 3, or 6 fluorophores per antibody was measured from 508 to 572 nm with an excitation wavelength at 460 nm. The dual labeled A5B7 with two fluorophores per antibody emitted the strongest fluorescence between 526 and 572 nm (Figure 1).

**Figure 1.** Fluorescence intensity of the dual labeled A5B7 (50 μg) with 1, 2, 3, or 6 fluorophores per antibody. Data are representative of three independent replicates (A). ELISA analysis to determine the EC50 of the dual labeled A5B7 with two fluorophores per antibody and the native A5B7 (mean ± SD, n = 3 independent replicates) (B).

**Figure 2.** Fluorescence images of SW1222 cells stained with the dual labeled A5B7 antibody and bisbenzimide H 33342, without (A) and with blocking using the native A5B7 (B) in vitro. Fluorescence image of an SW1222 tumor xenograft section stained with the dual labeled A5B7 antibody and the bisbenzimide H 33342 in vitro (C).
observed in the tumor, blood, and small intestine compared to other organs. Tumor uptake was 20.21 ± 2.70, 13.31 ± 0.73, and 10.64 ± 1.86 percentage injected dose per gram tissue (% ID/g) at these three time points, respectively (Supporting Information, Table S2). Uptake in the blood and small intestine was 16.42 ± 1.01 and 10.50 ± 1.59%ID/g, respectively, at 24 h, and rapidly decreased to 5.74 ± 2.27 and 2.02 ± 0.73%ID/g, respectively, at 72 h post injection (Supporting Information, Table S2).

Next, the 124I-Green dual labeled ASB7 in PBS was intravenously injected into CD1 nude mice (n = 3) bearing an established SW1222 tumor xenograft in the front-left flank area. Animals were imaged in three sequential 60 min PET/CT scans, performed at 24, 48, and 72 h post injection. The SW1222 tumor xenografts were clearly visualized in the 3D PET/CT images at all three time points, although distribution of the radioactivity signals within the tumors was heterogeneous (Figure 4A,B). Weaker radioactivity signals in the hearts were observed at 24 and 48 h but were cleared in 72 h PET/CT image (Figure 4B). Quantification of radioactivity level in selected organs including tumor, liver, and kidney was largely consistent with the biodistribution data (Supporting Information, Figure S4). As expected, accumulation of 124I in the thyroid glands was also detected and the radioactivity levels were quantified as 32.25 ± 3.24, 33.22 ± 1.09, and 36.92 ± 5.19%ID/mL at 24, 48, and 72 h post injection, respectively (Supporting Information, Figure S4).

All three animals were sacrificed at the end of the last PET/CT scan, and the SW1222 tumor xenografts were exposed. When observed in a whole-body fluorescence imaging system with an excitation wavelength of 460 nm, the xenografts were clearly visualized and distinguished from the surrounding tissue (Figure 5A). As a result, complete resection of the tumor was feasible (Figure 5B). The strength of fluorescence emission from the xenografts was compared with other major thoracoabdominal organs. Apart from the stomach and the large intestine, the tumor xenografts showed the strongest fluorescence (Figure 5C,D).

**Ex Vivo Sequential Autoradiography and Fluorescence Imaging.** Three tumor sections of a SW1222 xenograft resected from a mouse post the PET/CT imaging were exposed to a phosphor screen for 3 days. After scanning on a phosphor imager, autoradiography demonstrated the uneven distribution of the radioactivity signals in each tumor section (Figure 6A). The same tumor sections where then analyzed for fluorescence on the same imager using the fluorescein filter. The distribution and the intensity of fluorescence emission largely co-registered with the radioactivity signals in the autoradiograph of the corresponding tumor sections (Figure 6B).

### DISCUSSION AND CONCLUSION

Biomedical imaging is an essential contributor to cancer patient management that provides vital information required for cancer staging, therapeutic planning, and the monitoring of disease status. Multimodality imaging techniques can overcome the limitations of each imaging modality and supply complementary information to improve cancer care. PET and fluorescence imaging are highly synergistic. The former can provide quantitative measurement without limitation of penetration depth, while the latter offers superior spatial resolution. A cancer-specific dual PET and fluorescence imaging reagent labeled with a longer half-life radioisotope such as 124I (T1/2 = 4.2 days) could enable the cancer surgical treatment to adopt a highly integrated approach. PET/CT imaging of the malignant lesions would enable more accurate preoperative surgical planning. During surgery, the tumors and the metastatic lymph nodes, especially the deep-lying deposits, could then be rapidly localized with a hand-held radiation detector. Subsequently, fluorescence imaging would “light up” these tumors for real-time assessment and more accurate resection.

To prepare our dual PET and fluorescent labeling reagent, we chose the FDA approved fluorescein as the fluorescent reporter because it has successfully been used in fluorescence-guided surgery in clinical trials. The nonradioactive reference
compound of $^{124}$I-Green was synthesized in a moderate yield of 37%. The 5(6)-carboxyfluorescein regioisomer mixture was used for this proof-of-concept study due to its low cost relative to the pure 5 or 6 regioisomers. The mixture has similar chemical and photophysical properties as any of the single regioisomer, and once coupled with an antibody the...
pharmacokinetics of the conjugate will, in any case, be dominated by the macromolecule. Next, we optimized the radiosynthesis of $^{124}$I-Green. Iodine-125 was used as a surrogate of iodine-124 ($\gamma_{\text{max}}$ 35 keV vs 1691 keV for $^{121}$I) for the development work to reduce the cost of the screening and to minimize ionizing radiation exposure. Poor RCYs of the dual labeling reagent were initially obtained when applying the previously published conditions for the formation of radioiodinated 1,2,3-triazoles $^{14,15}$ As a phenanthroline analogue has been reported to accelerate the copper-mediated alkyne “click” reactions, $^{18}$ we screened structurally related ligands and found that addition of bathophenanthroline was beneficial for the radiochemical reaction. Using this ligand, 124I-Green was obtained with excellent RCYs around 92% determined by HPLC.

To explore the application of $^{124}$I-Green to cancer PET imaging and fluorescence-guided surgery, we selected a CEA-specific murine monoclonal antibody ASB7, a CEA-expressing SW1222 human colorectal cell line, and the corresponding tumor xenograft to systematically evaluate this dual modality imaging approach in vitro and in vivo. The specific binding of the directly labeled $^{131}$I-ASB7 to the SW1222 cells and derived xenografts is well documented in the literature. $^{16,19}$ Furthermore, $^{131}$I-ASB7 has been employed in several clinical trials for radioimmunotherapy of metastatic colorectal tumors and advanced gastrointestinal carcinomas. $^{20,21}$ Initially, to optimize the number of fluorophores coupled to the ASB7, we conjugated 1, 2, 3, or 6 of the nonradioactive reference compound per ASB7 molecule under mild basic conditions. As too many fluorophores were coupled to the antibody, this could have a detrimental effect on binding affinity, promote antibody aggregation, and reduce the strength of fluorescence through self-quenching. It became apparent that coupling of two fluorophores per ASB7 antibody emitted the strongest fluorescence. Moreover, this dual labeled ASB7 has similar EC$_{50}$ and HPLC retention time to the native antibody, indicating that its binding affinity was preserved and that no antibody aggregation took place. When incubated with the CEA-expressing SW1222 cells, the cells were stained with bright-green fluorescence. This was largely blocked by the native ASB7 antibody, demonstrating the specific binding of the dual labeled ASB7 to the CEA in live cells. Furthermore, intense green fluorescence was observed in an SW1222 tumor xenograft tissue section stained with the dual labeled ASB7, indicating that it could illuminate the SW1222 tumor xenograft in the ex vivo fluorescence imaging for accurate tumor resection.

Encouraged by these in vitro results, we investigated the biodistribution of the $^{124}$I-Green dual labeled ASB7 in the SW1222 tumor xenograft mouse model at 24, 48, and 72 h post-iv injection. The high tumor uptake and low uptake in thoracoabdominal organs was similar to that previously reported for the directly labeled $^{131}$I-ASB7. $^{16,19}$ The initially high blood pool and small intestine uptake observed at 24 and 48 h, clearing to background levels at 72 h, were in good agreement with the biodistribution pattern of a full-size antibody. To demonstrate the translational potential of the $^{124}$I-Green dual labeled ASB7 for the cancer PET imaging and fluorescence-guided surgery, we carried out three sequential PET/CT scans of SW1222 tumor xenograft-bearing mice at 24, 48, and 72 h post-iv injection followed by ex vivo fluorescence imaging. The tumor xenografts were clearly visualized at all three time points, reflecting the high affinity and specificity of the $^{124}$I-Green dual labeled ASB7 for CEA. Furthermore, uneven distribution of the radioactivity was observed in the tumor xenografts from the PET images, probably due to the heterogeneity of these tumors. In addition, significant thyroid uptake was observed in the PET images, most likely because of the catabolism of the $^{124}$I-Green dual labeled ASB7, generating free iodine-124 which subsequently accumulated in the thyroid. It was noted that the thyroid uptake increased over time, while the tumor, liver, and kidney levels gradually decreased. It expected that this thyroid uptake can be blocked by feeding the animals with 0.1% potassium iodide solution. $^{19}$ Some radioactivity was also observed in the hearts in the 24 h PET images, owing to blood pool activity, but this had largely cleared by the 48 and 72 h PET images. The largest ratio of uptake by tumor compared to blood and other thoracoabdominal organs was observed at 72 h post-iv injection in the biodistribution study. Consequently, we decided to carried out the ex vivo fluorescence imaging at this time point in order to obtain the optimal tumor-to-background contrast. Indeed, tumor xenografts were clearly visualized and exhibited a clear boundary with the surrounding tissue, which facilitated complete resection of the tumor. When compared with other major thoracoabdominal organs, the tumor xenografts emitted the strongest fluorescence apart from the stomach and the large intestine. Much of the strong fluorescence from the stomach and large intestine appeared to arise from food and fecal residue. However, the $^{124}$I-Green dual labeled antibody may also be partially eliminated through the gut, contributing to some of this background fluorescence. In the control experiments by treating two healthy animals with either native or dual labeled ASB7, the ex vivo fluorescence images at 72 h post-iv injection showed identical high fluorescence from the stomach food and large intestine feces (Supporting Information, Figure S5). In addition, tumor uptake was nearly 11 and 6 times greater than stomach and large intestine, respectively, at this time point in the biodistribution study. Thus, it indicates the majority of fluorescence was from the food and feces residue. This strong autofluorescence is not expected to be a problem in the clinical setting as all patients would be fasting and subject to preoperative bowel preparation to ensure that the colon is empty. Future development of the dual labeling reagent with a near-infrared fluorophore such as the indocyanine green would further minimize the background fluorescence. In addition, we also performed sequential autoradiography and fluorescence imaging with the same tumor tissue sections prepared using the xenografts from PET/CT and ex vivo fluorescence imaging. The distribution patterns of the radioactivity and fluorescence in these tumor sections were largely identical, which further confirmed that the observed radioactivity signals and green fluorescence in the tumors were from $^{124}$I-Green. Detection of deeper tumor lesions during fluorescence-guided surgery is challenging due to the limited tissue penetration of fluorescence; $^{17}$ in this context, an added advantage of our $^{124}$I-Green is that it would enable the complementary use of an intraoperative radioactivity detector to map the gross locations of the tumors, followed by real-time fluorescence imaging to define the surgical margins and guide complete tumor resection.

We have developed a generic dual PET and fluorescent labeling reagent, 124I-Green. This approach enables a one-step installation of an iodine-124/fluorescein dual reporter to any antibody. When coupled with the CEA specific antibody ASB7, the affinity of the dual labeled ASB7 was preserved. In vitro
fluorescence imaging of CEA overexpressing SW1222 cells indicated the specific binding of the dual labeled ASB7 to its antigens. The SW1222 xenografts in nude mice were clearly visualized by both PET/CT and ex vivo fluorescence imaging with excellent target-to-background contrast. These data demonstrate the feasibility of 124I-Green as a generic tool to label cancer-specific antibodies for malignant tumor PET imaging and fluorescence-guided surgery, prompting further translational research toward clinical application.

## EXPERIMENTAL SECTION

### Synthetic Chemistry and Radiochemistry. General Information.

H and 13C NMR spectra were recorded at rt on a Bruker Avance 400 instrument operating at the frequency of 400 MHz for H and 100 MHz for 13C. Chemical shifts are reported in ppm relative to DMSO (δ 2.48, m) or CH3CN (δ 9.46, s), and coupling constants (J) are given in hertz. HPLC analysis was performed with an Agilent 1200 HPLC system equipped with a 1200 series diode array detector and a GABI Star radioactivity detector. Reductant-free [125I]NaI was purchased from PerkinElmer (product no. NEZ309) in 0.02 M 5 M NaOH (pH 8) excluding the loading fraction. Formulation: [124I]NaI solution (6.0 MBq) in 0.02 M NaOH solution (6.0 mL). After incubation at rt for 90 min, the reaction mixture was quenched by HPLC using a ZORBAX column (300SB-C18, 9.4 mm, 5 μm) with the following eluent: water (0.1% formic acid) as eluent. The resulting suspension (62.5 μL) was added to a mixture of 4-azidomethyl-N-succinimidyl benzoate (226 μg, 1.0 μmol) in acetonitrile (20 μL) and [124I]NaI (~12 MBq) in 0.02 M NaOH solution (6.0 μL). After washing with water (5 μL), [124I]-Green was released using EtOH. The EtOH was removed with a stream of nitrogen, prompting further use.

For 124I-Labeling. Copper(II) chloride (3.4 mg, 25.3 μmol), triethylamine (4.4 μL, 3.15 μmol), and bipyridinium thiophosphate (850 μg, 2.5 μmol, 10 mol%) were mixed in anhydrous acetonitrile (500 μL). The resulting suspension (62.5 μL) was added to N-propargyl-5(6)-carboxyfluoresceinamide (1.4 mg, 3.1 μmol) in anhydrous DMF (62.5 μL). The resulting suspension was diluted to a mixture of 4-azidomethyl-N-succinimidyl benzoate (226 μg, 1.0 μmol) in acetonitrile (20 μL) and [124I]NaI (~12 MBq) in 1 × 10^{-5} M NaOH solution (6.0 μL).

For 124I-Labeling. Copper(II) chloride (3.4 mg, 25.3 μmol), triethylamine (4.4 μL, 3.15 μmol, 1.38 equiv), and bipyridinium thiophosphate (850 μg, 2.5 μmol, 10 mol%) were mixed in anhydrous acetonitrile (500 μL). The resulting suspension (62.5 μL) was added to N-propargyl-5(6)-carboxyfluoresceinamide (1.4 mg, 3.1 μmol) in anhydrous DMF (62.5 μL). The resulting red suspension (40 μL) was added to a mixture of 4-azidomethyl-N-succinimidyl benzoate (226 μg, 1.0 μmol) and triethylamine hydrochloride (TEA-HCl) (16.5 μg, 0.12 μmol) in acetonitrile (20 μL) and [124I]NaI (~12 MBq) in 0.02 M NaOH solution (6.0 μL).

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respectively) in DMSO (2.5 μL) were added to the antibody ASB7 (100 μL, 2.2 mg/mL) in 10 mM, pH 9.0 carbonate buffer. The resulting solution was incubated in the dark at rt for 45 min. The reaction was diluted to a total volume of 500 μL using pH 7.4 PBS and was purified on a PD-10 minipart G-25 size exclusion column (GE healthcare) eluting with pH 7.4 PBS. The fractions (200–700 μL) containing the dual labeled antibody ASB7 were collected. The purified ASB7 was submitted to HPLC analysis with a Phenomenex size exclusion column (BioSep-SEC-S3000) eluting with PBS-EDTA/ water 1:1 using a flow rate of 5.0 mL/min in 20 min with a UV detector at 280 nm. Fluorescence emission spectra between 508 and 572 nm of each of the dual labeled ASB7 antibodies (50 μg) was measured using the fluorescence detector in an Agilent 1200 HPLC system with an excitation wavelength of 460 nm. The antibody recovery rate from the PD-10 column was determined by loading the dual labeled ASB7 (50 KBq, 500 μL) on a PD-10 column and eluted with pH 7.4 PBS. The radioactivity fractions were collected and measured with a Raytest GABI star radioactivity detector. The antibody recovery rate was 80 ± 4% (n = 3). The average fluorophore(s) per antibody was then calculated based on the following equation:

Average fluorophore(s) per antibody = labeling efficiency × equivalents of added nonradioactive reference compound of dual labeling reagent/recovery efficiency of dual labeled antibody from the PD-10 column.

**ELISA Assay to Measure the EC₅₀ of the Dual Labeled ASB7 with CEA.** A 96-well plate was coated with carcinoembryonic antigen (CEA) at 10 μg/mL in PBS (100 μL) at 4 °C for 24 h. The plate was washed with PBS and blocked with 5% milk/PBST (150 μL) at rt for 1 h. Serial dilutions of the dual labeled or native ASB7 (100 μL, 0.03 nM, 0.14 nM, 0.70 nM, 3.0 nM, 19.2 nM, 96.0 nM, 480.0 nM, 2.4 μM, and 8.3 μM) were added to the plate and incubated at rt for 2 h. The plate was washed with PBS Tween-20 and deionized water, after which the sheep antimouse-horseradish peroxidase-IgG (100 μL) was added and incubated at rt for another 1 h. The samples were washed with PBS Tween-20 and deionized water. O-Phenylenediamine dihydrochloride solution (100 μL) was added to the samples, and color was developed in 5–10 min after addition of HCl (100 μL, 4.0 M). The absorbance was then read at 490 nm. The experiments were repeated three times. The data were analyzed using GraphPad Prism 7 software.

**Cell Line.** The human CEA-producing colorectal adenocarcinoma cell line SW1222 was obtained from the European Collection of Cell Cultures. Cells were cultured in DMEM medium supplemented with 10% FBS, 200 U/L penicillin, 0.1 g/L streptomycin, and 2 mM L-glutamine at 37 °C under humidified atmosphere containing 5% CO₂.

**Tumor Xenograft Model.** The human colorectal adenocarcinoma cell line SW1222 was used to develop a subcutaneous xenograft model by injecting 5 x 10⁶ cells in 100 μL of saline into the front left flank of 4 week old female CD1 nude mice. The CEA expression in the SW1222 tumor xenografts was previously quantified as 22–70 μg/g tumor. Tumor volume was estimated using the formula \( V = \frac{L \times W^2}{2} \) and the xenograft was established within 2 weeks. All animal experiments complied with the Animals (Scientific Procedures) Act (UK 1986) and Home Office (UK) guidelines and were conducted under a Home Office license (P9C94E8A4) with local ethical approval by the KCL College Research Ethics Committee (CREC).

**In Vitro SW1222 Cell Staining and Blocking Study.** SW1222 cells (1 x 10⁶) were plated on a 6-well plate and incubated in the cell culture media for 18 h. The medium was then removed, and the adherent cells were washed twice with PBS (2.0 mL). The dual labeled ASB7 (7 μg/mL) in PBS (1.0 mL) was added, followed by addition of either the native ASB7 solution (10 μL, 3.9 mg/mL) to three wells for blocking study or PBS (10 μL) to the remaining three wells. The cells were incubated for 1 h at 37 °C, 5% CO₂, Bisbenzimide H 33342 (Hoechst 33342) (10 μL, 0.25 mM) was then added 5 min before the end of the incubation period. Subsequently, the solution was removed and the cells were washed twice with ice-cold PBS (2.0 mL) before being viewed under a digital inverted fluorescence microscope EVOS fl (40x objective, AMG, USA) and the dual light cubes GFP and DAPI were selected for fluorescein and Hoechst, respectively.

**Immunofluorescence Imaging.** Frozen SW1222 tumor xenograft sections (10 μm) were fixed in acetone for 10 min at rt, incubated in 3% normal serum/PBS for 30 min, and then incubated with the dual labeled ASB7 (5 μg/mL) for another 30 min. Hoechst 33342 (10 μL, 0.25 mM) was then added 5 min before the end of the incubation period. After rinsing with PBS, sections were mounted in PBS and viewed under a digital inverted fluorescence microscope EVOS fl (4x objective, AMG, USA) and the dual light cubes GFP and DAPI were selected for fluorescein and Hoechst, respectively.

**Dual Labeled ASB7 Biodistribution Studies.** CD1 nude mice bearing established SW1222 tumor xenografts were intravenously injected with approximately 0.8 MBq of the 124I-Green dual labeled ASB7 (~44 μg) in 100 μL of PBS. The mice (n = 3) were culled by cervical dislocation at 24, 48, and 72 h post-injection, respectively. The tumor xenograft, major thoracoabdominal organs, left femur, and thigh muscle were harvested, weighed, and gamma-counted. The radioactivity in each organ was expressed as % ID/g. The total injected dose was defined as the sum of the whole body counts excluding the tail.

**PET/CT and Ex Vivo Fluorescence Imaging.** Preclinical PET/CT images were acquired using a NanoScan PET/CT (Mediso, Budapest, Hungary) scanner with mice under isoflurane (2% in oxygen) anesthesia. The SW1222 tumor xenograft-bearing CD1 nude mice (n = 3) each received approximately 0.8 MBq of the 124I-Green dual labeled ASB7 (~44 μg) in PBS (100 μL) via iv injection. PET scanning was performed for 60 min at 24, 48, and 72 h postinjection followed by a CT scan. All PET/CT data were reconstructed with the Monte Carlo-based full-3D iterative algorithm Tera-Tomo (Mediso Medical Imaging Systems, Budapest, Hungary). Raw PET data were reconstructed into 60 min bins using reconstruction settings (four iterations, six subsets, 0.4 x 0.4 x 0.4 mm³ voxel size) as well as intercrystal scatter correction. All reconstructed data were analyzed with VivoQuant software (v3.0, inviCRO, LLC, Boston, USA). All three animals were culled by cervical dislocation at the end of the last PET/CT scan. The skin surrounding the front left flanks was removed to expose the tumor xenografts. Fluorescence images were taken using a FOBI(tm) BGRN fluorescent imaging system (Neoscience, Suwon, Republic of Korea) both before and after the tumor resection. Subsequently, the tumor xenografts, major thoracoabdominal organs, and thigh muscle were harvested for fluorescence imaging with the FOBI fluorescence imaging system with an excitation wavelength of 460 nm. All fluorescence images were taken using LUCAM capture software.

**Ex Vivo Autoradiography and Fluorescence Imaging.** Three cryosections (5 μm thick) were prepared from the SW1222 tumor xenografts collected 1 week after the PET/CT and ex vivo fluorescence imaging. These were allowed to air-dry and then exposed to a phosphor screen (Molecular Dynamics) for 3 days. The phosphor screen was scanned in a Typhoon 8600 phosphor imager at a resolution of 50 μm. The same three tumor sections from the autoradiography were then used for fluorescence imaging in the same imager with the default emission filter setting for fluorescein (fluorescein 526 SP) with green laser (532 nm) at high sensitivity with a resolution of 50 μm. All images were analyzed using ImageJ software (v1.50, NIH, USA).

**ASSOCIATED CONTENT**

Supporting Information

The Supporting Information is available free of charge on the ACS Publications website at DOI: 10.1021/acs.jmedchem.7b01746.

ASB7 labeling efficiency and the average fluorophores per antibody; HPLC chromatograms including the crude radiodiomination mixture of 121I-Green, 121I-Green coeluting with its nonradioactive reference compound, and dual labeled and native ASB7; 121I-Green dual labeled.
ASB7 biodistribution data; radioactivity accumulation in selected organs extracted from the PET scans; ex vivo fluorescence imaging with healthy CD1 nude mice; $^3$H and $^{13}$C NMR spectra for all compounds prepared (PDF)

Molecular formula strings (CSV)

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Notes
The views expressed are those of the authors and not necessarily those of the NHS, the NIHR, or the DoH. PET scanning equipment was funded by an equipment grant from the Wellcome Trust.

The authors declare no competing financial interest.

■ ACKNOWLEDGMENTS

Dr Ran Yan thanks Bioprobe UK Ltd. for providing the FOBI fluorescence imaging system. Zhi Lu thanks the Scientific Study Project for Institutes of Higher Learning, Ministry of Education, Liaoning Province (L20151-49), China for support of this work. Truc Thuy Pham thanks Rosetrees Trust Ph.D. studentship (M545) for support of this work and Dr. Jayanta Bordoloi’s help with the development of animal model, in vivo PET imaging, and data analysis. The research was funded/ supported by the National Institute for Health Research (NIHR) Biomedical Research Centre at Guy’s and St. Thomas’ NHS Foundation Trust and King’s College London, the Wellcome/EPSCR Centre for Medical Engineering at King’s College London (WT 203148/Z/16/Z), the King’s College London and UCL Comprehensive Cancer Imaging Centre (C1519/A16463) funded by CRUK and EPSRC in association with the MRC and DoH (England), the Experimental Cancer Medicine Centre at King’s College, and the King’s Health Partners/King’s College London Cancer Research UK Cancer Centre. The work undertaken at UCLH/UCL was funded in part by the Department of Health’s NIHR Biomedical Research Centres funding scheme.

■ ABBREVIATIONS USED

Bq, becquerel; cCRC, clear cell renal cell carcinoma; CEA, carcinoembryonic antigen; CT, computed tomography; EC$_{50}$, half-maximal effective concentration; ELISA, enzyme-linked immunosorbent assay; FDA, Food and Drug Administration; $^{18}$F-FDG, 2-deoxy-2-[$^{18}$F]fluoro-D-glucose; HER2, growth factor receptor 2; ICG, indocyanine green; iv, intravenous; % ID/g, percentage injected dose per gram tissue; NIS, N-iodosuccinimide; PET, positron emission tomography; RCYs, radiochemical yields; rt, room temperature; TEA-HCl, triethylamine hydrochloride

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