An imaging agent to detect androgen receptor and its active splice variants in prostate cancer

Yusuke Imamura,1 Amy H. Tien,1 Jinhe Pan,2 Jacky K. Leung,1 Carmen A. Banuelos,1 Kunzhong Jian,1 Jun Wang,1 Nasrin R. Mawji,1 Javier Garcia Fernandez,1 Kuo-Shyan Lin,2 Raymond J. Andersen,3 and Marianne D. Sadar1

1Genome Sciences Centre and 2Department of Molecular Oncology, British Columbia Cancer Agency, Vancouver, Canada.
3Chemistry and Earth, Ocean, and Atmospheric Sciences, University of British Columbia, Vancouver, Canada.

Introduction

Most castration-resistant prostate cancer (CRPC) continues to be dependent on transcriptionally active androgen receptor (AR). Recently, powerful agents in the arsenal of hormone therapies for advanced prostate cancer, such as abiraterone and enzalutamide, have been approved based upon 5- to 6-month improvements in overall survival. Approximately 20%–40% of patients with metastatic CRPC have primary resistance to enzalutamide or abiraterone (1–4), and patients that were originally responsive will all eventually fail. One mechanism of resistance to current hormone therapies is the presence of constitutively active splice variants of AR (AR-Vs) (5–7).

Full-length AR is a ligand-activated transcription factor with distinct functional domains that include the C-terminal ligand-binding domain (LBD), to which androgens and antiandrogens bind; the hinge region, which contains a nuclear translocation sequence; the DNA-binding domain (DBD), which binds to androgen response elements (AREs) in the enhancers/promoters of target genes; and the N-terminal domain (NTD), which contains activation function-1 (AF-1) that is essential for AR transcriptional activity (8–10).

Constitutively active splice variants of androgen receptor (AR-Vs) lacking ligand-binding domain (LBD) are a mechanism of resistance to androgen receptor LBD–targeted (AR LBD–targeted) therapies for metastatic castration-resistant prostate cancer (CRPC). There is a strong unmet clinical need to identify prostate cancer patients with AR-V–positive lesions to determine whether they will benefit from further AR LBD–targeting therapies or should receive taxanes or investigational drugs like EPI-506 or galectonere. Both EPI-506 (NCT02606123) and galectonere (NCT02438007) are in clinical trials and are proposed to have efficacy against lesions that are positive for AR-Vs. AR activation function-1 (AF-1) is common to the N-terminal domains of full-length AR and AR-Vs. Here, we provide proof of concept for developing imaging compounds that directly bind AR AF-1 to detect both AR-Vs and full-length AR. 123I-EPI-002 had specific binding to AR-AF-1, which enabled direct visualization of CRPC xenografts that express full-length AR and AR-Vs. Our findings highlight the potential of 123I-EPI-002 as an imaging agent for the detection of full-length AR and AR-Vs in CRPC.

Conflict of interest: Y. Imamura, A.H. Tien, C.A. Banuelos, J. Wang, N.R. Mawji, K. Jian, J.G. Fernandez, R.J. Andersen, and M.D. Sadar are inventors and have licensed the technology to ESSA Pharma Inc. C.A. Banuelos, J. Wang, N.R. Mawji, J.G. Fernandez, R.J. Andersen, and M.D. Sadar all have shares in ESSA Pharma Inc., and R.J. Andersen and M.D. Sadar are directors and officers and receive consulting fees.

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Unfortunately, approaches to determine if a patient harbors metastases that express AR-Vs are not developed or validated, with each approach having significant shortcomings. Measurement of AR-Vs in CTCs is dependent on abilities to isolate/detect CTCs and the sensitivity of the assay to measure levels of AR-Vs. Current assays for AR-Vs in CTCs have limited reproducibility, sensitivity, and specificity, with the predictive value of the assay limited to only patients with detectable CTCs. Detection of CTCs is dependent on tumor volume and the presence of a surface antigen, such as HER2 (human epidermal growth factor receptor 2), EpCAM (epithelial cell adhesion molecule), or CD45; detection may not capture the entire CTC population. Circulating tumor-derived cell-free DNA (ctDNA) is also undergoing evaluation alongside CTCs for its utility in clinical practice (19, 20). Conceptually, ctDNA and/or ctRNA may be derived from primary tumors, metastatic lesions, CTCs, or from the plethora of benign tissues that are known to express AR-Vs (21). ctDNA analyses are restricted to measurable changes at the DNA level, and to date the clinical relevance of genomic rearrangement to generation of AR-Vs is lacking. Thus, there is an urgent need to develop clinical methods to determine expression of AR-Vs in metastatic CRPC. One possible approach is application of molecular imaging agents to detect AR-Vs. EPI small molecules provide an opportunity to develop such a prognostic imaging tool because they bind to AF-1 (22–24) that is common to both full-length AR and AR-Vs. EPI-001 and its stereoisomers (e.g., EPI-002) and analogs (e.g., EPI-506) were developed as AF-1 inhibitors to block transactivation of both full-length AR and AR-Vs. EPI-506 received investigational new drug approval from the FDA and Health Canada (i.e., acceptable safety profile at therapeutic doses) and is currently in clinical trials in the US and Canada for CRPC patients that have failed abiraterone and/or enzalutamide. Here, we provide proof of concept for developing an analog of EPI for imaging full-length AR and AR-Vs in CRPC patients. We provide evidence that single-photon emission computed tomography–computed tomography (SPECT/CT) successfully and specifically imaged prostate cancer xenografts that express AR-Vs after tail vein administration of microdoses of radioiodine 123-EPI-002 (123I-EPI-002). To specifically reveal AR-Vs, a discordant distribution and/or discordant level of uptake between 16β-[18F]-fluoro-5α dihydrotestosterone (18F-FDHT), which binds only full-length AR, and an imaging agent, such as 123I-EPI-002, which binds AF-1 that is common to both full-length AR and AR-Vs, would indicate the expression of solely AR-Vs. Currently, this may be technically challenging. However, if such an imaging approach were developed, it could reveal patients with lesions that are positive for AR-Vs who may not benefit from further therapies that target AR LBD. Our findings suggest that an imaging probe targeted to AR AF-1 may aid in monitoring treatment responses to significantly affect the clinical management of the disease as well as provide insight into the role of all AR species in resistance mechanisms.
Figure 2. I-EPI-002 has better potency than EPI-002 for AR and maintains specificity. (A) Dose-dependent inhibition of androgen induced transcriptional activity of endogenous full-length AR in LNCaP cells by I-EPI-002. n ≥ 3. Representative data are shown. (B) Effects of I-EPI-002 on androgen-dependent proliferation of LNCaP cells treated with R1881 compared with PC3 and DU145 cell viability. n ≥ 3. Data represent mean ± SD. (C) Effect of I-EPI-002 (2 μM) vs. EPI-002 (25 μM) on AR transcriptional activity in LNCaP cells that were transiently transfected with PSA(6.1 kb) luciferase reporter and induced with R1881 (1 nM). n = 5. Data represent mean ± SEM. (D) Effect of I-EPI-002 (2 μM) vs. EPI-002 (25 μM) on 4-pregnene-3,20-dione (Progesterone, 10 nM) induced PR transcriptional activity in LNCaP cells that were transiently transfected with PRE luciferase reporter and expression vector for PRβ. n = 5. Data represent mean ± SEM. (E) A representative competition binding curve showing displacement of 1 nM fluorescently labeled ligand from recombinant AR LBD (25 nM) by enzalutamide and R1881 but not with I-EPI-002 or EPI-002. The experiment was repeated 3 times. (F) Androgen-induced AR-transcriptional activity measured in LNCaP cells transfected with PSA(6.1 kb) luciferase reporter and treated with vehicle (DMSO), enzalutamide (ENZ, 5 μM), I-EPI-002 (5 μM), or EPI-002 (25 μM) and increasing concentrations of R1881. n = 4. Values shown are the mean ± SD. I-EPI, I-EPI-002; EPI, EPI-002; ENZ, enzalutamide. (A-F) Each independent experiment was performed in triplicate. **P < 0.01. One-way ANOVA Dunnett’s multiple comparison test was utilized.
Results
I-EPI-002. EPI-001 and its analogs bind AF-1 (22) that is common to both full-length AR and AR-Vs. For an imaging agent to be successful, it must have excellent binding affinity for its target. The most potent stereoisomer of the EPI-001 mixture is EPI-002 (23), which was employed as the starting scaffold for synthesis of cold 15-IodoEPI002 (called I-EPI-002) (Figure 1 and described in the Supplemental Experimental Procedures; supplemental material available online with this article; doi:10.1172/jci.insight.87850DS1).

Iodination improves the potency of EPI-002 in functional assays. Iodination of EPI-002 (I-EPI-002) could potentially alter many of the properties of the parent compound EPI-002, such as potency, biological activity, and off-target effects. Since excellent binding affinity is critical for an imaging agent to be successful, we first tested whether iodination of EPI-002 improved or diminished its potency using well-established cell-based functional assays. Dose response curves using the AR-driven prostate-specific antigen (PSA) (6.1 kb) luciferase reporter were used to calculate IC<sub>50</sub> values for inhibition of endogenous AR transcriptional activity by I-EPI-002 compared with EPI-002 in LNCaP cells. I-EPI-002 had an IC<sub>50</sub> of 1.17 ± 0.22 μM for inhibition of AR transcriptional activity (Figure 2A). This is comparable to that reported for the antiandrogen enzalutamide, which is considered to have strong affinity for the AR. At 1 μM, enzalutamide caused 67% inhibition of ARE(4x) luciferase activity (25). I-EPI-002 had an IC<sub>50</sub> of approximately 1 μM to block AR-driven proliferation of LNCaP cells in response to androgen (Figure 2B). I-EPI-002 had no effect on proliferation or viability of PC3 and DU145 human prostate cancer cells that do not express functional AR at up to 10 times the IC<sub>50</sub> required to reduce AR-dependent proliferation, thereby supporting its specificity for AR. I-EPI-002 is 10 times more potent than EPI-002, which has IC<sub>50</sub> in the range of 10 μM for these cell-based assays (22, 23). Together, these data support that I-EPI-002 has improved potency compared with EPI-002 and are consistent with saturation transfer difference–NMR data, showing strong interaction of the bisphenyl rings of EPI with Tau-5 of AR AF-1 (24).

I-EPI-002 maintains specificity for AR. Imaging agents are administered at microdoses and thus must be highly specific for their targets. Therefore, we next examined whether I-EPI-002 would potentially interact with the proteins most highly related to AR in the human proteome, which are the other steroid receptors, progesterone receptor (PR), glucocorticoid receptor (GR), and estrogen receptor (ER). In addition to high sequence identity in the LBD and DBD of this class of receptors, they also interact with many of the same proteins, such as CBP (CREB-binding protein) and SRC1–3. A sensitive method to determine if I-EPI-002 interacts with any of these other receptors is to measure their transcriptional activities. Therefore, reporter gene assays were used to determine if I-EPI-002 would inhibit PR, GR, or ER transcriptional activities. Cells were cotransfected with expression plasmids for full-length human PRβ, GR, ERα, and their respective reporters and then treated with ethanol vehicle, 4-pregnene-3,20 dione (progesterone) (PRβ), dexamethasone (GR), or estradiol (E2) (ERα). At a concentration of ~2 μM, I-EPI-002 inhibited AR transcriptional activity to similar levels (75%) to that achieved with 25 μM EPI-002 (Figure 2C). Importantly, I-EPI-002 did not inhibit the transcriptional activities of PR, GR, or ER in response to their cognate ligands (Figure 2D and Supplemental Figure 1, A and B). These data provide evidence that I-EPI-002 does not have general effects on transcription or translation because it did not inhibit induction of PRE (progesterone response element), GRE (glucocorticoid response element), orERE (estrogen response element) luciferase reporters in response to ligand. Thus, I-EPI-002 appears to maintain specificity for AR.

I-EPI-002 does not interfere with ligand binding. EPI analogs bind specifically to AR AF-1 without being affected by, or interfering with, ligand binding to AR LBD (22, 23). This is an important attribute for an imaging agent for CRPC, because AR LBD may be occupied by antiandrogen or steroid/androgen. To test whether I-EPI-002 maintained this property and does not affect ligand binding to AR LBD, we employed the fluorescence polarization assay. This assay measures competition of a test compound for binding of fluoromone to recombinant AR LBD or other related steroid hormone receptor LBDs. R1881 and the antiandrogen enzalutamide both bound to the AR LBD to compete with the fluoromone as expected (Figure 2E). I-EPI-002 behaved consistently to EPI-002 and did not prevent ligand-binding to the LBDs of AR or any steroid hormone receptors tested (Supplemental Figure 1, C–F). Consistent with these data, I-EPI-002 and EPI-002 maintained effective inhibition of AR transcriptional activity with increasing concentrations of androgen (Figure 2F), unlike enzalutamide, which markedly lost potency, as expected for a competitive antiandrogen. These data support that imaging with a molecular probe targeted to AR AF-1 should not be affected by an occupied AR LBD.

I-EPI-002 targets AR-Vs. Constitutively active AR-Vs that lack LBD are detected in clinical samples of CRPC, and levels of expression are correlated with poor prognosis and resistance to abiraterone.
In LNCaP95 cells, proteins in LNCaP95 cells are shown by Western blot analyses. LNCaP95 cells were serum starved for 24 hours and then treated with DMSO, enzalutamide (5 μM), or EPI-002 (25 μM) treatment for 48 hours. Bivariate plots show cell cycle distribution of a representative experiment. The stacked graph (below) represents the average results from 3 independent experiments. The percentage of cells in each phase of cell cycle was calculated. Effects of enzalutamide, EPI-002, and I-EPI-002 on cell cycle–regulated proteins in LNCaP95 cells are shown by Western blot analyses. LNCaP95 cells were serum starved for 24 hours and then treated with DMSO, enzalutamide (5 μM), EPI-002 (25 μM), or I-EPI-002 (5 μM), or EPI-002 (25 μM) for 2 days. Data represent mean ± SEM. Effects of enzalutamide, EPI-002, and I-EPI-002 on cell cycle–regulated proteins in LNCaP95 cells are shown by Western blot analyses. LNCaP95 cells were serum starved for 24 hours and then treated with DMSO, enzalutamide (5 μM), EPI-002 (25 μM), or I-EPI-002 (5 μM) for 48 hours (Figure 3A, left). Consistent with previous studies, enzalutamide had no effect in blocking AR transcriptional activity, either in the presence or absence of androgen, when V567es or V7 was expressed (Figure 3A, middle and right). I-EPI-002 showed good inhibition against full-length AR as well as mixed populations of full-length AR with variant V567es or V7 at concentrations 10 times lower than those of EPI-002. This suggests that I-EPI-002 interacts with the truncated AR variants, consistent with EPI-002, but with better potency. Western blot analysis using an antibody against the AR NTD confirmed the approximate 1:1 ratio of endogenous full-length AR to ectopic V567es or V7 in whole cell lysates of cells treated with I-EPI-002, EPI-002, and enzalutamide (Figure 3B). I-EPI-002 was next tested to examine if it had effects on AR-V–driven proliferation. LNCaP95 cells are enzalutamide resistant (12, 28) and increase proliferation in the absence of androgen by a mechanism driven by AR-Vs. I-EPI-002 (2 μM) decreased proliferation of LNCaP95 cells as well as EPI-002 (25 μM), whereas enzalutamide (5 μM) had no effect (Figure 3C). I-EPI-002 had an IC50 of 6.89 ± 2.85 μM for inhibition of LNCaP95 cell proliferation (Figure 3D). Consistent with these data, cell cycle analysis revealed that EPI analogs caused G0/G1 arrest. Approximately 25%–30% of cells treated with enzalutamide or DMSO vehicle were in S phase (Figure 3E). Whereas I-EPI-002 or EPI-002 decreased S phase cells by approximately 2-fold or more with a concomitant increase of cells in G1 phase. Western blot analysis of cell cycle G1/S-related proteins revealed that EPI analogs decreased the expression of UBE2C (ubiquitin-conjugating enzyme E2C), an AR-V7–regulated protein; cyclinD3; and cyclin-dependent kinases (CDKs). EPI analogs increased protein expression of the CDK inhibitor p27kip1. Importantly, enzalutamide had no effect on the levels of expression of these proteins, which was consistent with these cells being resistant to enzalutamide (Figure 3F). Together, these data support that I-EPI-002 targets full-length AR and AR-Vs.

Synthesis of 123I-EPI-002 imaging probe. The preparation of [123I]15-iodoEPI002 is shown in Figure 4A. The radiosynthesis involves three reactions. First, the [123I]iodoacetone was prepared in 90% ± 10% radiochemical yield via aromatic electrophilic iodination reaction using chloramines-T as the oxidizing reagent and the acetonitrile 3 as the precursor. The isolated [123I]iodoacetone was subsequently coupled with (2R)-(−)-glycidyl tosylate to form [123I]iodoepoxide 8 in 37% ± 17% radiochemical yield. At the final step, [123I]15-iodoEPI002 was obtained in 70% ± 11% radiochemical yield by hydrolysis of the [123I]iodoepoxide 8 with CeCl3·7H2O. Due to multiple radiolabeling and HPLC purification steps, [123I]15-iodoEPI002 (123I-EPI-002) was isolated in an overall 14% ± 4% (n = 12) decay-corrected yield in a 4.5-hour synthesis time with >99% radiochemical purity and 3.4 ± 1.0 Ci/μmol specific activity at the end of synthesis.

123I-EPI-002 binds to AF-1 in the AR NTD in cells. EPI-002 and analogs bind to AR AF-1 (22, 23) and specifically to Tau-5 in the AF-1 region (24). Here, we show that 123I-EPI-002 binds to recombinant full-length AR (Figure 4B). To determine if 123I-EPI-002 binds the same site as EPI-002, AF-1 protein was preincubated with excess cold EPI-002 prior to addition of radioactive 123I-EPI-002 probe. Excess cold EPI-002 displaced binding of 123I-EPI-002 to AF-1, thereby suggesting they bind to the same site (Figure 4C). Ultimately, to ensure specific binding of 123I-EPI-002 to endogenous AR in cells, LNCaP95 cells were treated with 123I-EPI-002 for 16 hours, followed by SDS-PAGE and Western blot analysis. 123I-EPI-002 binds covalently with Tau-5 in the AF-1 region (24).
Figure 4. Synthesis of $^{123}$I-EPI-002 and its binding to AR AF-1. (A) Synthetic route to $[^{123}]$15-iodoEPI-002 starting from (R)-tosylglycerolactonide[1] and bisphenol A[1] and using Na$^{123}$I as a source of radiolabeled iodine. (B) Recombinant full-length AR protein was incubated with $^{123}$I-EPI-002 for 16 hours at room temperature. Binding of $^{123}$I-EPI-002 to AR was detected by SDS-PAGE followed by phosphorimaging. Coomassie blue staining was used to provide an indication of equal loading. (C) Recombinant AF-1 protein was incubated with vehicle (DMSO) or EPI-002 (60 $\mu$M) for 6 hours at room temperature prior to addition of $^{123}$I-EPI-002 (200 $\mu$Ci) and then incubated for 16 hours more prior to SDS-PAGE and phosphorimaging. Less binding of $^{123}$I-EPI-002 was observed when AF-1 was preincubated with excess cold EPI-002. Coomassie blue staining was used to provide an indication of equal loading (refer to full and uncut gel images in Supplemental Materials). (D) Binding of $^{123}$I-EPI-002 to endogenous AR in LNCaP95 cells that were incubated with or without 25 $\mu$M EPI-002 overnight at 37°C prior to harvesting cells. Whole cell lysates were used for SDS-PAGE to reveal $^{123}$I-EPI-002 covalently bound to full-length AR in the LNCaP95 cells, as detected by phosphorimaging. Western blot analysis detection of bands corresponding to AR for radiolabeled phosphorimaged bands. $n \geq 3$ for B, C, and D binding experiments.
and specifically to AR in the LNCaP95 cells (Figure 4D). There was no indication that 123I-EPI-002 was a random alkylator, consistent with previous reports that EPI-001 and EPI-002 do not react with glutathione at physiological pH (23, 29). These data support the specificity of 123I-EPI-002 binding AR AF-1.

In vivo biodistribution of 123I-EPI-002

Biodistribution studies of 123I-EPI-002 were conducted in male castrated NOD/SCID mice that each carried both a LNCaP95 and a PC3 xenograft. The LNCaP95 cell line was selected as the target CRPC xenograft due to its high level of expression of endogenous full-length AR and AR-V7. PC3 was selected as the nontarget xenograft because it has very low/negligible protein levels of AR. Table 1 outlines the in vivo tissue biodistribution results of 123I-EPI-002. The cLogP of 123I-EPI-002 is 4.2, which means it is highly lipophilic, a property that predicts it would be eliminated by the hepatobiliary system. Consistent with this projection, 123I-EPI-002 accumulation was highest in the gallbladder, liver, and intestines (Table 1). At 4 hours after injection, 66 percent of the injected dose per gram (66% ID/g) of 123I-EPI-002 was detected in the large intestine. Generally, there was low uptake and fast washout from nonspecific tissues.

Accumulation of 123I-EPI-002 in LNCaP95 tumors did not support that it was binding irreversibly in this or any other tissues (Table 1). To date, covalent binding of EPI to AR NTD has only been shown in vitro and in closed systems over a long time (23, 24). Binding of EPI-002 is proposed as a 2-step process: a quick reversible binding, followed by a very slow covalent binding (23, 24). It may be plausible that the short t½ of EPI in vivo (~3 hours) may result in negligible covalent binding because it is not around long enough for covalent binding to occur.

123I-EPI-002 reached a maximum 2.2% ± 0.5% ID/g uptake in LNCaP95 tumors at 1 hour after injection. This was greater than the maximum achieved for 18F-FDHT (0.432% ± 0.183% ID/g ± SEM) in the target tissue of ventral prostate in rodents at 1 hour after injection (30). Both 123I-EPI-002 and 18F-FDHT target the nuclear AR. Based on similar studies with 18F-FDHT, the accumulation of 123I-EPI-002 in AR-rich tissues is expected for an AR-mediated process (31). At this time point, 123I-EPI-002 uptake in AR-deficient PC3 tumors was only 0.7% ± 0.4% ID/g, providing a ratio of 3.2

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Tumor and normal organ uptakes are expressed as percent of the injected dose per gram (%ID/g). Values are shown as mean ± SD. ND, not determined.
for AR-rich tissue (LNCaP95) compared with AR-deficient tissues (PC3). LNCaP95 tumor to blood ratio was 3.58 ± 0.66 (Figure 5A), which was similar to the 3.5 ratio achieved with 18F-FDHT, which is used clinically (31). LNCaP95 tumor-blood and tumor-muscle ratios were greater than that of PC3 for all time points examined. Blocking with excess cold EPI-002 (50 mg/kg body weight) caused a 74% decrease (P < 0.001) in 123I-EPI-002 accumulation in the LNCaP95 xenograft (Figure 5B), suggesting that the uptake of 123I-EPI-002 was AR mediated. This 74% drop in LNCaP95 tumors by blocking with excess cold EPI-002 was similar to the 70% drop achieved with testosterone to block 18F-FDHT (31). Blocking with excess cold EPI-002 had no significant effect on levels of 123I-EPI-002 uptake in blood, muscle, and PC3 xenograft (Figure 5B). These data suggest that the accumulation of 123I-EPI-002 in LNCaP95 xenografts was specifically AR mediated.

Micro-SPECT/CT imaging of 123I-EPI-002. On the basis of the results from the biodistribution study and ratios of target tissues to nontarget tissues that were at similar to those achieved with clinically employed 18F-FDHT, a whole-body micro-SPECT/CT study was performed to evaluate the specific tumor-targeting ability of 123I-EPI-002. To do this, male castrated NOD/SCID mice bearing both LNCaP95 and PC3 xenografts of approximately equal volumes were scanned using SPECT. Scanning at 2 hours after tail vein injection of 123I-EPI-002, when the tumor-blood and tumor-muscle ratios were sufficiently high, provided SPECT images shown in Figure 5C. 123I-EPI-002 distinguished LNCaP95 tumors from PC3 tumors, whereas the liver region had relatively higher uptake (Figure 5C; Unblock). The amount of radiopharmaceutical uptake seen in the liver, which was concurrent with the low amount of gastrointestinal uptake, was likely due to the lipophilic nature of 123I-EPI-002 and was consistent with the in vivo biodistribution data. LNCaP95 tumor uptake was similar to the tumor uptake observed in the biodistribution study (1.09% ± 0.41% ID/g at 2 hours after injection, n = 3). In addition, the 123I-EPI-002 tracer accumulation was absent from SPECT images when blocked with excess cold EPI-002 (Figure 5C; Block). Importantly, the thyroid uptake was approximately 2.31% ± 0.59% ID/g and was not altered by blocking with EPI-002 (2.32% ± 0.33% ID/g). These data suggest that 123I-EPI-002 may not be metabolically stable and that the 123I may fall off the compound to accumulate in the thyroid.

Histological analysis showed that LNCaP95 xenografts have more necrotic centers than PC3 xenografts (Figure 5D). This suggests that poor vascularization and consequent central necrosis may prevent optimal access of 123I-EPI-002 to AR in LNCaP95 tumors. Western blot analysis using an antibody against the AR NTD and V7 confirmed the expression of full-length AR and V7 in LNCaP95 xenografts, which were undetectable in PC3 xenografts (Figure 5E). These data support that 123I-EPI-002 accumulation in the LNCaP95 tumor was specific to full-length AR and AR-Vs. Together, all of these data support the feasibility of developing molecular imaging agents directed to AR AF-1 to detect the AR and AR-Vs to improve the clinical management of prostate cancer.

Discussion

Bone scans are routinely used for metastatic CRPC but they do not provide insight about the cancer itself or whether the lesions are AR positive to aid in treatment selection. Although bone metastases are the predominate site of dissemination, approximately 20%–30% of metastatic CRPC patients have visceral metastases (32, 33). If visceral metastases are positive for full-length AR, they could benefit from hormone therapies that target AR LBD, such as enzalutamide (34). Even so, approximately one-third of CRPC patients will have intrinsic resistance, and eventually all patients will acquire resistance to enzalutamide or abiraterone (1–4). The major mechanisms of resistance to these therapies include gain-of-function mutations in AR LBD and/or expression of constitutively active AR-Vs that lack LBD (5–7). An important clinical example is drawn from a report using CTCs from CRPC patients, which revealed that, upon initiating enzalutamide treatment, those patients with CTCs that express V7 have a lower PSA response rate, shorter PSA-progression-free survival, shorter clinical or radiographic progression-free survival, and reduced overall survival compared with patients that had CTCs that were V7 negative (5). Compared with taxanes, hormone therapies are the preferred therapeutic due to low toxicity. However, taxanes are more efficacious than AR LBD-targeting therapies in V7-positive CRPC patients (18). Therefore, the ability to distinguish those patients that express AR-Vs that are resistant to AR LBD therapies would be a significant advance in the clinical management of CRPC.

To date, assays to detect AR-Vs are still in early developmental stages, with most focused upon detection of AR-Vs in CTCs. This is because biopsy of the multiple lesions that predominate in the bone of CRPC patients is not viable, making the comprehensive assessment of expression of AR-Vs in metastatic tissues...
unfeasible. Clinical application of measurement of expression of AR-Vs using CTCs has several caveats that include abilities to isolate or detect CTCs and the sensitivity of the assay to measure levels of AR-Vs. Thus, a noninvasive imaging approach that could robustly reveal the presence of full-length AR and AR-Vs in all metastatic lesions is of utmost importance to identify patients that may or may not benefit from further antiandrogen or androgen ablation therapies.

Figure 5. 123I-EPI-002 shows uptake in xenografts that endogenously express ARs. (A) Tumor-blood and tumor-muscle ratios of LNCaP95 xenografts. \( n = 3–8 \) at each time point. Data are shown as mean ± SEM. (B) Effects of EPI-002 cotreatment on 123I-EPI-002 accumulation in blood, muscle, LNCaP95, and PC3 xenografts (1-hour treatment). The accumulation of 123I-EPI-002 in LNCaP95 xenografts was significantly decreased by blocking with EPI-002. \( n = 3–7 \). Data represent mean ± SD. *** \( P < 0.001 \). One-way ANOVA Dunnett’s multiple comparison test was utilized. (C) Axial and coronal micro-SPECT/CT images of castrated NOD/SCID mice bearing LNCaP95 (arrow) and PC3 (arrowhead) xenografts 2 hours after coinjection of 123I-EPI-002 with (Block) or without excess dose of EPI-002 (Unblock). The images are scaled to the same threshold. Color scale indicates the level of radioactivity. \( n = 2 \). (D) H&E staining and immunohistochemistry (AR) of representative sections of formalin-fixed, paraffin-embedded LNCaP95 and PC3 xenografts with lower magnification. Xenografts were extracted approximately 6 weeks after subcutaneous injection of LNCaP95 and PC3 cell lines. The histological analysis demonstrated that LNCaP95 cells that expressed the AR formed a largely concentrated tumor, which resulted in a necrotic center. Scale bar: 40 \( \mu \)m. (E) Protein expression obtained from LNCaP95 and PC3 xenografts. Four xenografts from each cell line are shown. Western blot analysis of LNCaP95 xenografts showing both full-length AR and AR-V7 expression, both of which were negative in PC3 xenografts. \( \beta \)-Actin is shown as a protein loading control. Refer to full and uncut gel images in Supplemental Materials.
The optimal qualities for a potentially useful imaging probe to provide a noninvasive approach to localize metastatic lesions that express full-length AR and AR-Vs are a high binding affinity to AR NTD and low nonspecific binding affinity. Here, we propose developing an analog of EPI-002 as an AR NTD-targeted molecular imaging probe to detect all AR species. The fact that AR protein is commonly overexpressed in CRPC tissues supports the application of AR NTD–based imaging agents. We show here that I-EPI-002 (a) has 10 times greater potency than EPI-002; (b) maintained specificity for AR; (c) bound to the target, AR AF-1, to yield loss of functional transcriptional activities as expected; and (d) most importantly, that in vivo 123I-EPI-002 enabled specific visualization of xenografts that express full-length AR and AR-Vs. I-EPI-002 is potentially the first imaging agent that can directly detect both full-length AR and AR-Vs. Importantly, these data also support the specificity of EPI-002 for AR.

The addition of a substituent in EPI-002 chemical structure could alter the binding of the probe to AR directly or by conformational effects. Here, the addition of iodine to EPI-002 had favorable effects on blocking the transcriptional activity of AR NTD without losing its specificity. The fact that 123I-EPI-002 binds the AR NTD and does not affect binding of ligand to AR LBD is an important attribute for an imaging agent for this indication and molecular target. This is because imaging agents are applied at microdoses and hence are highly susceptible to competing agents. For most applications, it is predicted that patients failing enzalutamide or other antiandrogens may have an AR LBD occupied by the antiandrogen. Washout of enzalutamide takes approximately 5 half-lives ($t_{1/2}$ = 6–9 days) or more than 1 month. In addition, if there is steroid hormone breakthrough in castrated or abiraterone-treated patients, or a mutation in AR LBD making the receptor promiscuous to binding other steroid, AR LBD could be occupied by steroid. Under these clinical conditions, an imaging agent that binds to AR NTD would theoretically not be affected, contrary to 18F-FDHT, which competes for binding to LBD and has been shown to have diminished uptake at the tumor site in the presence of testosterone (35). The quality of competing for the AR LBD can, however, be beneficial, as in application of 18F-FDHT as a pharmacodynamic marker to demonstrate the effects of agents targeting the full-length AR (1). Most importantly, in order to distinguish full-length AR from AR-Vs, an imaging agent that binds to the AR NTD, such as 123I-EPI-002, which detects full-length AR and AR-Vs, would be required to be used in combination with an imaging agent that only detects full-length AR, such as 18F-FDHT. By using sequential imaging, a discordant distribution or discordant level of uptake between 18F-FDHT and a radiolabeled AR NTD imaging agent would indicate the presence of AR-Vs. This experiment was beyond the scope of these proof-of-concept studies; but, in the future, this experiment can be carried out with improved analogs. Such improvements that are currently underway encompass synthesizing a metabolically more stable imaging agent to use in place of 123I-EPI-002 and moving away from an iodocompound. However, 124I-EPI-002 might be useful for PET imaging and an 125I-EPI-002 might be useful as a Auger emitter for therapeutic applications. In the event that an AR NTD–targeted molecular imaging probe has appropriate uptake and biodistribution, it could be further developed as a targeted radionuclide to deliver α- or β-emitting radionuclides to lesions that express full-length AR and/or AR-Vs in patients with metastatic CRPC.

In conclusion, these studies suggest that an AR NTD–targeted molecular imaging probe has significant potential for monitoring treatment response; providing insight into the role of all AR species, including AR-Vs, in resistance mechanisms of CRPC; and may provide a means of targeted radiotherapy.

Methods

Cells, plasmids, transfections, and reporter assays. The synthetic androgen R1881 was purchased from PerkinElmer. EPI-002 was provided by NAEJA. Enzalutamide was purchased from Omega Chem. Progesterone (4-pregnene-3,20-dione) was obtained from Steraloids Inc. Dexamethasone and HEPES were obtained from Sigma-Aldrich. LNCaP, PC3, and DU145 cell lines as well as PSA(6.1 kb) luciferase, probasin luciferase, PRE luciferase, GRE luciferase, ERE luciferase, AR-V7 and AR-V567es plasmids, and transfection of cells have been described previously (22, 23, 26, 29). LNCaP cells were obtained from Leland Chung (Cedars-Sinai Medical Center, Los Angeles, California, USA) in September 1993. DU145 cells were from Victor Ling (British Colombia Cancer Agency, Integrative Oncology) in October 1998. LNCaP95, an androgen-independent cell line that expresses full-length AR and AR-V7, was provided by Stephen R. Plymate (University of Washington, Seattle, Washington, USA) in February of 2012 and reported in previous studies (12, 14, 36, 37). PC3 cells were purchased from ATCC. LNCaP95 cells were not authenticated in our laboratory but were regularly tested to ensure that they were mycoplasma free (VenorTMGeM Mycoplasma Detection kit,
Sigma-Aldrich). LNCaP, PC3, and DU145 cells were authenticated by short tandem repeat analysis and tested to ensure that they were mycoplasma free by DDC Medical in September 2013. All cells used in the experiments were passaged in our laboratory for fewer than 3 months after resurrection.

General procedures for synthesis of I-EPI-002 and [123I]-EPI-002. [123I] NaI was purchased from Nordion. (S)-1,2-isopropylidenediglycerol, 4-(dimethylamino)pyridine, pyridine, NN-dimethylformamide, p-toluenesulfonyl chloride, bisphenol A, sodium iodide, sodium perchlorate, sodium hydroxide, HPLC grade methanol, sodium hydroxide, (2R)-glycidyl tosylate, cerium chloride heptahydrate, HPLC grade acetonitrile, phosphate buffer, and chloramine-T were purchased from Sigma Aldrich. Purification and quality control of [123I]-labeled compounds were performed on an Agilent HPLC System equipped with a model 1200 quaternary pump, a model 1200 UV absorbance detector (set at 220 nm), and a Bioscan NaI scintillation detector. The radiodetector was connected to a Bioscan B-FC-1000 Flow-count System, and the output from the Bioscan Flow-count system was fed into an Agilent 35900E Interface, which converted the analog signal to digital signal. The operation of the Agilent HPLC system was controlled using the Agilent ChemStation software. The HPLC columns used were a semipreparative column (Phenomenex Luna C18, 5 μ, 250 x 10 mm) and an analytical column (Phenomenex Luna C18, 5 μ, 250 x 4.6 mm). Radioactivity of [123I]-labeled compounds was measured using a Capintec CRC-25R/W dose calibrator. Details are available in the Supplemental Experimental Procedures.

Fluorescence polarization. Androgen, PR, ER, and GR PolarScreen Competitor Assay kits (Invitrogen) were used according to the manufacturer’s protocol. Serial dilution was done for each small molecule, and solvent was compensated to ensure equal volume of DMSO and ethanol in each sample. Fluorescence polarization at excitation wavelength 470 nm and emission at 530 nm was measured in Greiner 384 black clear-bottomed plates using Infinite M1000 (TECAN).

Viability and proliferation assays. PC3 cells (2,000 cells/well), DU145 cells (2,000 cells/well), and LNCaP cells (5,000 cells/well) were plated in 96-well plates in respective media plus 0.5% FBS. The next day, PC3 and DU145 cells were treated with DMSO (vehicle control) and I-EPI-002 for 3 days, and LNCaP cells were pretreated with vehicle and I-EPI-002 for 1 hour before treatment with 0.1 nM R1881 for 4 days. Proliferation and cell viability was measured using alamarBlue Cell Viability Assay (Invitrogen) following the manufacturer’s protocol.

For BrdU incorporation experiments, LNCaP95 cells (8,000 cells/well) were seeded in 96-well plates for 48 hours in RPMI with 10% charcoal stripped serum, which was changed to serum-free media 24 hours before treatment with 5 μM enzalutamide, 25 μM I-EPI-002, 2 μM I-EPI-002, or serial concentrations of I-EPI-002. BrdU incorporation was measured after 2 days using the BrdU ELISA kit (Roche Diagnostics) according to the manufacturer’s protocol.

FACS analysis. LNCaP95 cells (75,000 cells/dish) were plated in 10-cm dishes and were treated with inhibitors under serum-free and phenol red-free conditions for 48 hours. Cells were pulse labeled with 10 μM BrdU for 2 hours and fixed in 70% ethanol. BrdU-labeled cells were probed with anti-BrdU-FITC antibody (BD Biosciences), and DNA was stained with 7-aminoactinomycin D (Sigma-Aldrich). Data were acquired using a FACS Calibur (BD Biosciences). Bivariate analysis was performed using FlowJo 7 software (Ashland).

Binding assays. The cell-free binding assay was performed with commercially available full-length AR recombinant protein or AR AF-1 recombinant protein that was expressed and purified as previously described (22, 38), with additional purification by size exclusion chromatography. Binding reactions between EPI-002 or DMSO (vehicle control) and the recombinant AR AF-1 (amino acid residues 142–485) or full-length AR were carried out by mixing 60 μM EPI-002 or DMSO with 10 μM AR AF-1 and 1 μM EPI-002 or DMSO with 0.35 μM full-length AR in equivalent amounts. Binding reaction was incubated at room temperature for 6 hours. 200 μCi of [123I]-EPI-002 was then mixed in each sample at room temperature for 16 hours prior to SDS-PAGE. Proteins were separated from free probes on 8% SDS-PAGE, and radioactivity was visualized using Fujifilm FLA-7000 image analyzer (GE Healthcare). The same gel was stained with Coomassie blue R-250. To examine binding to the endogenous AR, LNCaP95 cells (60,000 cells/well) were seeded in 12-well plates and serum starved for 24 hours before treating cells for 16 hours with DMSO or 25 μM EPI-002 and with 400 μCi of [123I]-EPI-002. Proteins were extracted from treated cells with RIPA buffer containing 50 mM tris (pH 8.0), 150 mM NaCl, 0.5% Na Deoxycholate, 1% NP-40, 0.1% SDS, and EDTA-free protease inhibitors. [123I]-EPI-proteins were separated by SDS-PAGE, and radioactivity was visualized using Fujifilm FLA-7000 image analyzer, and subjected to Western blot analysis using anti-AR antibody.
Immunohistochemistry and Western blot analysis. Immunohistochemistry was performed as previously described (22, 23). For analysis of AR protein, concentrations of lysates of homogenized LNCaP95 and PC3 xenografts were measured by bicinchoninic acid assay after albumin depletion. Proteins (20 μg) were resolved on a NuPAGE 4%–15% Bis Tris gradient gel, transferred to nitrocellulose membrane, and probed for AR species using antibodies to the AR NTD and AR-V7. LNCaP95 cells (250,000 cells/well) were seeded in a 6-well plate for 48 hours and serum-starved for 24 hours, followed by treatment with DMSO, enzalutamide (5 μM), EPI-002 (25 μM), or I-EPI-002 (5 μM) for 48 hours. Cells were harvested, and whole-cell lysate (10 to 15 μg) was subjected to SDS-PAGE. Antibodies used were as follows: AR N-20 (sc-816, Santa Cruz Biotech Inc.); AR-V7 (AG10008, Precision Antibody); UBE2C (A-650, BostonBiochem); and cyclinD1, cyclinD3, p27kip1, CDK2, CDK4, and CDK6 from the Cell Cycle Regulation Sampler Kit (9932 from Cell Signaling Technology). β-Actin (ab6276, Abcam) was used as a loading control.

Xenografts and animal biodistribution studies. Six- to eight-week-old male NOD/SCID mice were cas-trated 2 weeks before inoculating LNCaP95 cells (10 million cells/tumor) and PC3 cells (2 million cells/tumor) subcutaneously. The mice bearing LNCaP95 and PC3 xenografts were injected with 370 kBq to 7.4 MBq (10–200 μCi) of 123I-EPI-002 in 200 μl of 30% PEG solution via the tail vein. At 0, 1, 2, 4, 6, and 12 hours after injection, blood, liver, kidney, spleen, lung, large intestine, small intestine, stomach, heart, muscle, bone, bladder, brain, gallbladder, and tumor samples were collected and weighed and their radioactivity content was determined using an automated γ-spectrometer. The results were expressed as %ID/g. More than 3 animals per time point were used.

Micro-SPECT/CT imaging. SPECT/CT studies were performed using the MILabs U-SPECT II (Advanced Molecular Vision). LNCaP95 and PC-3 tumor-bearing mice (20-25 g) were anesthetized with isoflurane inhalation at 2% and placed on a heating pad to maintain body temperature. The mice were then injected with 123I-EPI-002 (37–74 MBq; 1–2 mCi, in 200 μl of 30% PEG solution) via the tail vein. A competition experiment was performed by coinjection of the radiotracer with 50 mg/kg of EPI-002 in a total volume of 200 μl of 30% PEG solution. SPECT scans were acquired over 30 minutes and 2 frames at 15 minutes each using 1.0-mm multi-pinhole collimator, after which CT scans were performed for anatomic reference (parameters; 60 kV, 600 μA). SPECT imaging data were reconstructed using MI Labs reconstruction software (2 subsets, 30 iterations). PMOD software was used to analyze and view the images, and a Gaussian filter was applied after reconstruction.

Statistics. Statistical analysis was performed using GraphPad Prism (version 6.01 GraphPad Software Inc.). Values are presented as mean ± SD or mean ± SEM. Except where specified, comparisons between groups were performed with 1-way ANOVA Dunnett’s multiple comparisons test, and differences were consider statistically significant at P values less than 0.05.

Study approval. All experiments involving animals conformed to the relevant regulatory and ethical standards established by the University of British Columbia Animal Care Committee. The University of British Columbia Animal Care Committee approved the experiments.

Author contributions
YI acquired and analyzed the data and wrote the manuscript. AHT acquired and analyzed the data. JP synthesized the compound. JKL performed BrdU assay. CAB performed FACS analysis. KJ administrated technical or material support. JW performed animal studies. NRM analyzed the data and administrated material support. JGF administrated technical or material support. K-SL supervised and administrated material support. RJA developed the chemical compound. MDS conceived of the concept; developed, designed, and supervised the work; and wrote the manuscript. All the authors approved the paper.

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Address correspondence to: Marianne D. Sadar, Department of Genome Sciences Centre, British Columbia Cancer Agency, 675 West 10th Avenue, Vancouver, British Columbia, Canada V5Z1L3. Phone: 604.675.8157; E-mail: msadar@bcgsc.ca.


