
Placental Growth Factor (PIGF)–Specific Uptake in Tumor Microenvironment of ⁸⁹Zr-Labeled PIGF Antibody RO5323441

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Placental growth factor (PIGF) is a member of the proangiogenic vascular endothelial growth factor family, which is upregulated in many tumors. RO5323441, a humanized monoclonal antibody against PIGF, showed antitumor activity in human tumor xenografts. We therefore aimed to radiolabel RO5323441 and preclinically validate this tracer to study drug tumor uptake and organ distribution by PET imaging. ⁸⁹Zr-RO5323441 was tested for stability and immunoreactivity in vitro. **Methods:** The tumor uptake and organ distribution for 10, 50, and 500 μg of ⁸⁹Zr-RO5323441 was assessed in mice bearing human PIGF-expressing hepatocellular cancer (Huh7) xenografts or human renal cell carcinoma (ACHN) xenografts without detectable human PIGF expression. The effect of pretreatment with RO5323441 (20 mg/kg) on ⁸⁹Zr-RO5323441 tumor uptake was analyzed in Huh7 xenografts. ¹¹¹In-IgG served as a control for nonspecific tumor uptake and organ distribution. Cy5-RO5323441 was injected to study the intratumor distribution of RO5323441 with fluorescence microscopy. **Results:** ⁸⁹Zr-RO5323441 showed a time- and dose-dependent tumor accumulation. Uptake in Huh7 xenografts at 10 μg of ⁸⁹Zr-RO5323441 was 8.2% ± 1.7% injected dose (ID)/cm³ at 144 h after injection, and in ACHN xenografts it was 5.5 ± 0.3 %ID/cm³ (*P* = 0.03). RO5323441 pretreatment of Huh7 xenograft-bearing mice reduced ⁸⁹Zr-RO5323441 tumor uptake to the level of nonspecific ¹¹¹In-IgG uptake. Cy5-RO5323441 was present in the tumors mainly in the microenvironment. **Conclusion:** The findings show that RO5323441 tumor uptake is PIGF-specific and time- and dose-dependent.

Key Words: placental growth factor; RO5323441; angiogenesis; positron emission tomography; ⁸⁹Zr

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Angiogenesis is a key feature of tumors (1). Inhibition of angiogenesis by targeting vascular endothelial growth factor (VEGF) and VEGF receptors (VEGFRs) has emerged as treatment for various tumor types (2). However, sustained clinical benefit of angiogenesis inhibitors can be hampered by compensatory mechanisms such as the upregulation of proangiogenic factors such as placental growth factor (PIGF) (2–5). Targeting PIGF could, thus, be a new strategy for tumor angiogenesis inhibition, complementary to VEGFR inhibition, and might circumvent resistance observed during current antiangiogenic therapies.

PIGF, a VEGF homolog, can be expressed by tumor cells and is expressed by several other cells (e.g., endothelial and smooth muscle cells, macrophages, fibroblasts, and leukocytes) in the tumor microenvironment (4). PIGF is present in low levels in normal tissue. It contributes to the angiogenic switch in pregnancy, wound healing, ischemic conditions, and tumor growth (6). PIGF inhibition preclinically slows down growth and metastasis of various tumors, including those resistant to VEGFR inhibitors, and enhances the efficacy of chemotherapy and VEGFR inhibitors. Additionally, PIGF inhibition reduces angiogenesis, lymphangiogenesis, and tumor cell motility. Distinct from VEGFR inhibitors, PIGF inhibition preclinically prevents infiltration of angiogenic macrophages and severe tumor hypoxia and, thus, does not initiate the angiogenic rescue program responsible for resistance to VEGFR inhibitors (7,8).

These findings led to the clinical development of RO5323441 (TB-403) (9,10). RO5323441 is a humanized monoclonal antibody against PIGF-1 and PIGF-2. In a phase I trial with RO5323441 in cancer patients, stable disease was observed in 6 of 23 patients at different dose levels. No dose-limiting toxicities were observed with doses of up to 30 mg/kg administered once every 3 wk. Because no maximum-tolerated dose could be defined (11), determination of the optimal dose for phase II studies was hampered. Rational dosing might be obtained when tumor and normal

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tissue uptake of the antibody is defined with ^{89}Zr -RO5323441 PET. The feasibility of this approach for visualization of soluble angiogenic factors was already shown for the VEGF-A antibody bevacizumab (12–14). The aim of the current study was therefore to study ^{89}Zr -RO5323441 tumor uptake and organ distribution in human tumor xenograft models with different PIGF expression.

MATERIALS AND METHODS

Cell Lines

Human hepatocellular carcinoma Huh7 (Health Science Research Resources Bank) with high human PIGF (hPIGF) expression (hPIGF messenger RNA expression 62-fold higher than HEK293 native) and the human renal cell carcinoma ACHN (American Type Culture Collection) without detectable hPIGF expression (hPIGF messenger RNA expression equal to HEK293 native) were cultured in Dulbecco modified Eagle medium containing glucose (1 g/L) supplemented with 10% heat-inactivated fetal calf serum (Bodinco BV) at 37°C in humidified atmosphere containing 5% CO_2 .

The Abelson murine leukemia virus–induced tumor BALB-c–derived RAW264.7 murine macrophage cell line (American Type Culture Collection) was cultured in Dulbecco modified Eagle medium containing glucose (4.5 g/L) supplemented with 10% fetal calf serum and 1% L-glutamine at 37°C in humidified atmosphere containing 5% CO_2 .

Conjugation, ^{89}Zr Labeling, and Quality Control of ^{89}Zr -RO5323441

RO5323441 was conjugated and labeled as described by Verel et al. (15). In short, RO5323441 (25 mg/mL; Roche) was first conjugated with a 5-fold molar excess of the chelator *N*-succinyl-desferrioxamine-tetrafluorophenol (*N*-sucDf-TFP; provided by Dr. Guus van Dongen, VU University Medical Center). *N*-sucDf-RO5323441 was purified by ultracentrifugation using a 30-kDa Vivaspin-2 filter (Sartorius), diluted in water for injection (2.5 mg/mL), and stored at –20°C. Labeling was performed with ^{89}Zr (IBA) within 24 h before use. Radiochemical purity was evaluated by size-exclusion high-performance liquid chromatography using a Superdex 200 10/300 GL column (GE Healthcare) and showed no aggregates, fragments, free ^{89}Zr , or other impurities (Supplemental Fig. 1A; supplemental materials are available online only at <http://jnm.snmjournals.org>). ^{89}Zr -RO5323441 stability was evaluated at 4°C in solvent (0.9% NaCl) and at 37°C in phosphate-buffered saline (140 mM NaCl, 9 mM Na_2HPO_4 , and 1.3 mM NaH_2PO_4 ; pH 7.4) and in human serum by trichloroacetic acid precipitation. ^{89}Zr -RO5323441 was highly stable in 0.9% NaCl at 4°C and in phosphate-buffered saline or human serum at 37°C with less than 5% ^{89}Zr release over 168 h in all tested conditions. The preservation of immunoreactivity was tested in a competition assay with unlabeled RO5323441 with recombinant hPIGF-1 (Peprotech) as the target antigen, according to the assay previously described (16). The RO5323441 concentration required for 50% reduction in hPIGF-1 binding of 5 nM ^{89}Zr -RO5323441 was 5.1 ± 1.5 nM RO5323441, showing a fully preserved immunoreactivity (Supplemental Fig. 1B). ^{111}In -IgG was produced as described previously (16).

Generation of Cy5-Labeled RO5323441

Purified RO5323441 was incubated in 100 mM potassium phosphate, pH 8.5, with a 5-fold molar excess of Cy5 *N*-hydroxysuccinimide ester (GE Healthcare Life Sciences) dissolved in dimethyl sulfoxide.

The reaction was stopped after 60 min by the addition of 10 mM L-lysine, and the surplus of the labeling reagent was removed by dialysis against 20 mM histidine, 200 mM sodium chloride, 5% saccharose, pH 6.0. The Cy5 labeling degree was determined at an absorbance of 280 and 650 nm and was 2.7:1. The specificity of Cy5-labeled RO5323441 was analyzed by confocal fluorescence microscopy using recombinant Hek293 cells expressing membrane-tagged PIGF. Cy5-labeled RO5323441 was bound to the surface of the cells expressing membrane-anchored PIGF. The specificity was confirmed by competition with unlabeled RO5323441.

Animal Studies

All invasive procedures and imaging in animal experiments were performed with isoflurane inhalation anesthesia (induction, 5%; maintenance, 2%). Tumor cells were harvested by trypsinization, resuspended in culture medium and Matrigel (BD Biosciences), and inoculated subcutaneously (5×10^6 per mouse) in 6- to 8-wk-old male athymic nude mice (BALB-c/Ola HSD-fox nude; Harlan). Each subgroup consisted of 4 mice. Tumor growth was assessed 2 times per week with caliper measurement. When tumors measured 6–8 mm in diameter (~ 0.3 cm³), in vivo imaging studies were started using a small-animal PET Focus 220 rodent scanner (CTI Siemens).

Huh7 xenograft-bearing mice were coinjected in the penile vein with 10, 50, or 500 μg of ^{89}Zr -RO5323441 (5 MBq) and corresponding protein doses of ^{111}In -IgG (1 MBq). Coinjection with ^{111}In -IgG provides insight into behavior of an intraanimal nonspecific control for RO5323441 tumor uptake and organ distribution, improves the statistical power, and reduces animal numbers. ACHN xenograft-bearing mice and non-tumor-bearing mice were coinjected with ^{89}Zr -RO5323441 (10 μg ; 5 MBq) and ^{111}In -IgG (10 μg ; 1 MBq). In the pretreated group, mice were injected intraperitoneally with RO5323441 (20 mg/kg) 2 d before ^{89}Zr -RO5323441 injection (10 μg ; 5 MBq). Animals were imaged at 24, 72, and 144 h after injection, followed by ex vivo biodistribution analysis as described previously (16).

For microscopic analysis of tumor localization of RO5323441, Huh7 and ACHN xenograft-bearing mice were injected intravenously with 50 μg of Cy5-RO5323441, followed by tumor excision at 24 h after injection. In vivo stability of Cy5-RO5323441 was confirmed by near-infrared fluorescence imaging, which indicated accumulation in Colo205 tumors over 48 h without specific accumulation in the liver or kidneys.

All animal experiments were approved by the animal experiments committee of the University of Groningen.

Tumor Tissue Analyses

Tumors were kept on ice during biodistribution analysis and subsequently processed for histology and fluorescence microscopy. Hematoxylin and eosin staining was performed on formalin-fixed, paraffin-embedded tumors to assess tumor morphology. For fluorescence microscopy, tumor slides were stained for cell nuclei with Hoechst 33258 (Invitrogen). Macrophages were visualized by a 2-step staining with goat-antimouse CD68 (Santa Cruz) and donkey-antigoatIgG-Alexa488 (Invitrogen). Slides were analyzed with a Leica DM6000B microscope, and images were captured with a DFC360FX camera (Leica) and processed with LAS-AF2 software (Leica).

Fresh-frozen tumors were lysed and analyzed for hPIGF and murine PIGF (mPIGF) protein levels using an enzyme-linked immunosorbent assay (ELISA; R&D Systems) according to the manufacturer's protocol.

⁸⁹Zr-RO5323441 Cell-Binding Studies

Binding of ⁸⁹Zr-RO5323441 to the human Huh7 and ACHN cell lines and the murine RAW264.7 macrophage tumor cell line was studied in the presence or absence of hPIGF-1 (Peprotech) or mPIGF-2 (R&D Systems). ⁸⁹Zr-RO5323441 (0.22 nM; 10 kBq) was incubated for 24 h at 37°C with 1 × 10⁶ cells per well. During incubation, PIGF (human or murine) was absent or added in a 10:1, 1:1, and 1:10 molar ratio with RO5323441. An Fc receptor blocking reagent (20 μL; no. 130-059-901 [MiltenyiBiotec]) was used to block the Fc_γ receptors (Fc_γR) in RAW264.7. After incubation, cells were washed before harvesting, and the cell-bound ⁸⁹Zr was measured in a well-type γ-counter (LKB Wallac). All conditions were tested in 3 independent experiments.

Statistical Analysis

Data are presented as mean ± SD. For statistical analysis, GraphPad Prism (version 4.00 for Windows; GraphPad Software) was used. Statistical analysis was performed using the 2-tailed unpaired *t* test. A *P* value of less than 0.05 was considered significant.

RESULTS

⁸⁹Zr-RO5323441 Uptake in hPIGF-Expressing Huh7 Xenografts

In Huh7 tumors, ⁸⁹Zr-RO5323441 uptake was seen at all dose cohorts, and uptake increased over time, indicating specific tumor uptake. The 144-h scan for the 10-μg ⁸⁹Zr-RO5323441 dose cohort showed the best tumor visualization, with a higher tumor-to-background ratio than the 50- and 500-μg ⁸⁹Zr-RO5323441 dose cohorts (Fig. 1A).

Quantification of the ⁸⁹Zr-RO5323441 presence in the small-animal PET scans showed that ⁸⁹Zr-RO5323441 blood-pool levels were similar for the 3 dose cohorts, with a decrease between 24 and 144 h (Fig. 1B). ⁸⁹Zr-RO5323441 tumor uptake increased over time at all protein dose cohorts; the highest percentage injected dose per centimeter cubed (%ID/cm³) in tumors occurred in the 10-μg cohort at 144 h.

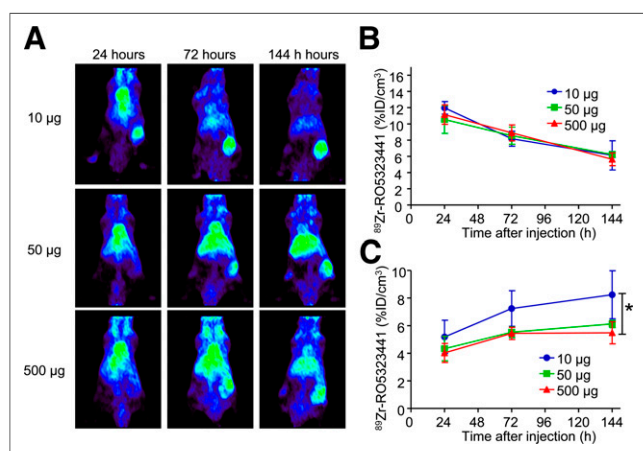


FIGURE 1. (A) Representative small-animal PET examples of Huh7 xenografts injected with 10, 50, or 500 μg of ⁸⁹Zr-RO5323441 and scanned at 24, 72, and 144 h after tracer injection. (B and C) Small-animal PET quantification of ⁸⁹Zr-RO5323441 blood-pool activity (B) and tumor uptake (C).

For the 50- and 500-μg cohorts, maximal tumor uptake was observed at 72 h and remained the same at 144 h (Fig. 1C). The absolute ⁸⁹Zr-RO5323441 tumor uptake levels were 0.8 ± 0.2, 3.1 ± 0.2, and 27.4 ± 3.98 μg/cm³ for, respectively, the 10-, 50-, and 500-μg injected dose cohorts. For ⁸⁹Zr-RO5323441, the quantified small-animal PET data show a non-dose-dependent blood clearance and a time- and dose-dependent tumor uptake.

Ex vivo biodistribution analysis in Huh7 xenograft-bearing mice confirmed a normal IgG distribution of ⁸⁹Zr-RO5323441 in nontumor organs. This distribution was hardly RO5323441 protein dose-dependent. In contrast, relative ⁸⁹Zr-RO5323441 tumor uptake showed a clear RO5323441 dose-dependent decrease in %ID per gram (%ID/g) of tumor (Fig. 2A). In the 10-μg cohort, ⁸⁹Zr-RO5323441 tumor uptake was, respectively, 72% (*P* = 0.016) and 109% (*P* = 0.023) higher than ¹¹¹In-IgG at 72 and 144 h after tracer injection (Fig. 2B). At 144 h, tumor uptake of ⁸⁹Zr-RO5323441 in the 50-μg cohort was 91% (*P* = 0.0041) and in the 500-μg 79% (*P* = 0.0004) higher than for ¹¹¹In-IgG (Figs. 2C and 2D). These biodistribution data confirm the time- and dose-dependent ⁸⁹Zr-RO5323441 tumor uptake in Huh7 xenografts observed with the small-animal PET quantification.

On the basis of the high tumor-to-nontumor contrast on small-animal PET images, the tumor accumulation over time shown by small-animal PET scan quantification and biodistribution data, and the high ratio with ¹¹¹In-IgG observed in our biodistribution analysis, 10 μg was chosen as the ⁸⁹Zr-RO5323441 protein dose for further experiments.

⁸⁹Zr-RO5323441 Biodistribution in Non-Tumor-Bearing Mice

For non-tumor-bearing mice, organ distribution of 10 μg of ⁸⁹Zr-RO5323441 was similar to that of ¹¹¹In-IgG (Supplemental Fig. 2). There was no specific accumulation of

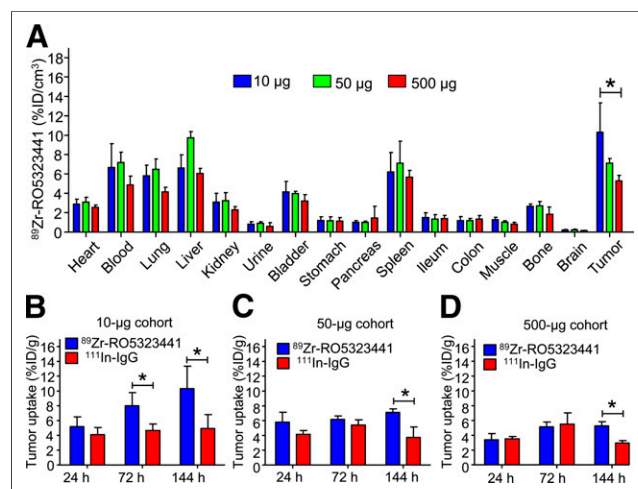


FIGURE 2. (A) ⁸⁹Zr-RO5323441 biodistribution in Huh7 xenografts at 144 h after tracer injection in all 3 dose cohorts. (B–D) ⁸⁹Zr-RO5323441 and ¹¹¹In-IgG tumor uptake of 10- (B), 50- (C), and 500-μg (D) cohorts.

^{89}Zr -RO5323441 over time in most nontumor organs, except for the liver (+26% from 24 to 144 h after injection; $P = 0.027$), and a nonsignificant trend in bone. Liver and bone uptake were possibly the result from binding to scavenging receptors in the liver or could indicate *in vivo* tracer metabolism to ^{89}Zr -labeled antibody fragments and release of free ^{89}Zr , which could accumulate in bone. Organ distribution of ^{89}Zr -RO5323441 was similar in non-tumor- and tumor-bearing mice.

RO5323441 Pretreatment Reduces ^{89}Zr -RO5323441 Uptake in Huh7 Tumors

Small-animal PET quantification showed that a 20 mg/kg pretreatment of RO5323441 reduced the tumor uptake of 10 μg of ^{89}Zr -RO5323441 by 33% ($P = 0.026$) (Fig. 3A). ^{89}Zr -RO5323441 tumor uptake in RO5323441-pretreated mice was comparable to the tumor uptake in the 500- μg ^{89}Zr -RO5323441 dose cohort. Biodistribution data showed that pretreatment with RO5323441 in Huh7 xenografts reduced ^{89}Zr -RO5323441 tumor uptake to background levels—that is, not higher than ^{111}In -IgG (Fig. 3B)—confirming the small-animal PET quantification data. Biodistribution of ^{89}Zr -RO5323441 in nontumor organs was not influenced by RO5323441 pretreatment (data not shown).

^{89}Zr -RO5323441 Uptake in ACHN Tumors That Do Not Express hPIGF

Small-animal PET scans showed that ^{89}Zr -RO5323441 tumor uptake in ACHN tumors was 33% lower ($P = 0.021$) than in Huh7 tumors at 144 h after injection (Fig. 4A). Biodistribution results revealed that ^{89}Zr -RO5323441 uptake in ACHN tumors was comparable to ^{111}In -IgG uptake. The ratio of ^{89}Zr -RO5323441 to ^{111}In -IgG in Huh7 tumors was higher than the ratio in ACHN tumors at all time points: 46% at 24 h ($P = 0.0003$), 61% at 72 h ($P = 0.023$), and 89% at 144 h ($P = 0.0005$) after tracer injection (Fig. 4B). These results indicate PIGF tumor level specific uptake in Huh7 tumors.

Ex Vivo Analysis of Huh7 and ACHN Tumors

Tumor histology of Huh7 and ACHN xenografts obtained from the 10- μg ^{89}Zr -RO5323441 cohorts at 24, 72, and 144 h after injection showed areas of tumor cells, stromal cells,

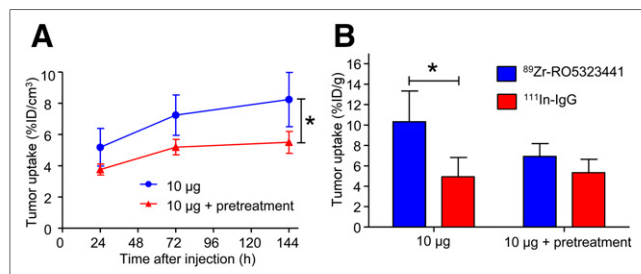


FIGURE 3. Influence of RO5323441 pretreatment on ^{89}Zr -RO5323441 tumor uptake in Huh7 xenografts quantified from small-animal PET data (A) and biodistribution data at 144 h after tracer injection (B).

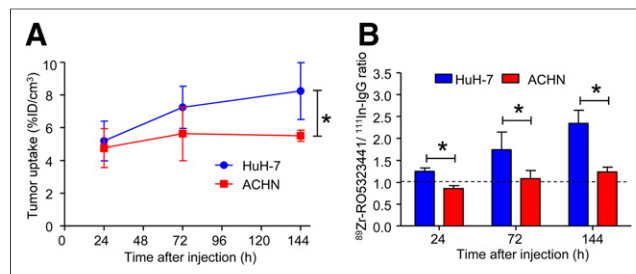


FIGURE 4. ^{89}Zr -RO5323441 uptake in ACHN and Huh7 tumors as assessed by small-animal PET (A) and biodistribution (B).

vasculature, and some necrotic areas (Fig. 5A shows representative hematoxylin and eosin staining).

Fluorescence microscopy revealed that in Huh7 tumors, Cy5-RO5323441 accumulated in areas of tumor cells and in necrotic regions, vasculature, and the surrounding connective tissue (Fig. 5A). In ACHN tumors, Cy5-RO5323441 accumulated only in the surrounding connective tissue, necrotic regions, and vasculature but not in the areas of tumor cells. Costaining of the Huh7 tumor slides with an anti-CD68 antibody to visualize macrophages indicated that Cy5-RO5323441 was present in the CD68-positive regions and in the vasculature (Fig. 5B).

hPIGF protein levels measured with ELISA were 153 ± 62 pg/mg of protein in Huh7 xenografts and nondetectable in ACHN xenograft tissue. Murine PIGF levels were 253 ± 177 pg/mg of protein in Huh7 and $1,063 \pm 207$ pg/mg of protein in ACHN xenograft tissue.

^{89}Zr -RO5323441 Binding to RAW264.7 Macrophages

To further study the mechanism by which cells, such as macrophages, that are in a microenvironment might be involved in RO5323441 tumor uptake, we studied the

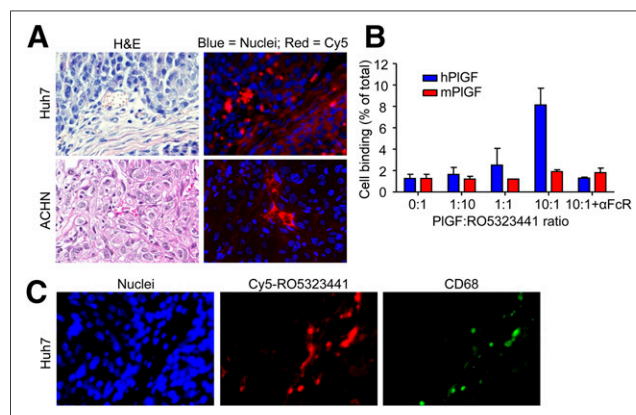


FIGURE 5. (A) Representative examples of Huh7 and ACHN histology, as assessed by hematoxylin and eosin (H&E) staining, and Cy5-RO5323441 microscopic tumor distribution, as assessed by fluorescence microscopy. (B) ^{89}Zr -RO5323441 binding to RAW264.7 macrophages after 24 h incubation with hPIGF or mPIGF and an Fc receptor blocking reagent (αFcR). (C) Hoechst staining visualizes cell nuclei. Costaining of Huh7 tumor with anti-CD68 antibody to visualize macrophages.

uptake of ^{89}Zr -RO5323441 by RAW264.7 murine tumor macrophages in vitro. The binding of ^{89}Zr -RO5323441 to RAW264.7 concentration dependently increased in the presence of hPIGF, with a 6.4-fold increase ($P = 0.0018$) at a 10-fold molar excess of hPIGF (Fig. 5C). The hPIGF-induced macrophage binding of ^{89}Zr -RO5323441 was completely blocked by coincubation with a Fc receptor blocking reagent. mPIGF was not able to induce binding of ^{89}Zr -RO5323441 to RAW264.7 macrophages, indicating the higher affinity of RO5323441 for hPIGF.

The binding of ^{89}Zr -RO5323441 to Huh7 cells was 2-fold higher than binding to ACHN cells ($P = 0.0030$, Supplemental Fig. 3), possibly due to the autocrine production of heparin-binding hPLGF by Huh7. ^{89}Zr -RO5323441 binding to Huh7 and ACHN was not affected by the non-heparin-binding hPIGF-1 isoform. The addition of mPIGF, which also binds heparin, induced a slight increase in both cell lines (1.6- and 1.9-fold, respectively).

DISCUSSION

In this study, we determined ^{89}Zr -RO5323441 tumor uptake and organ distribution in human tumor-bearing mice. We used 3 different tracer doses and 3 different imaging and biodistribution time points, 2 tumor models with different target expression levels, an unspecific IgG control, and a RO5323441 pretreatment dose. All together, these data showed that ^{89}Zr -RO5323441 tumor uptake was time-, dose-, and PIGF-dependent, supporting the feasibility of implementing the use ^{89}Zr -RO5323441 with PET scanning for clinical studies.

Because this is the first study, to our knowledge, on in vivo visualization of PIGF, the best possible comparison is with results obtained with ^{89}Zr -labeled bevacizumab. For optimal comparison, the antibody tracer dose should be considered. ^{89}Zr -bevacizumab showed a SKOV-3 tumor uptake of 6.8 ± 1.8 %ID/g with 100 μg of antibody (12), which is similar for ^{89}Zr -RO5323441 tumor uptake. Higher levels of ^{89}Zr -RO5323441 tumor uptake, expressed as %ID/g, might have been obtained when lower antibody doses would have been used. ^{111}In -bevacizumab biodistribution studies showed uptake levels of up to 25 %ID/g for a 1- μg bevacizumab dose (17). In the present study, we used 10 μg of ^{89}Zr -RO5323441 because this was the lowest dose that allowed sufficient counts for PET quantification at 6 d after injection. It would have been possible to perform a biodistribution-only experiment with a lower dose (e.g., 1 μg), and this would likely have resulted in a higher tumor uptake when expressed as %ID/g. However, ex vivo biodistribution studies are nonsequential and not translational, and we aimed to validate ^{89}Zr -RO5323441 as an imaging tool. Overall, ^{89}Zr -RO5323441 PIGF PET preclinical performance is comparable to that of ^{89}Zr -bevacizumab VEGF PET.

Fluorescence-labeled RO5323441 was present in the tumor microenvironment at 24 h after injection. Our mouse models were engrafted with human Huh7 and ACHN tumors, and therefore the microenvironment of these human tumor

xenografts consisted of human tumor cells and murine stroma. This microenvironment, as shown with ELISA, contained both hPIGF and mPIGF. RO5323441 binds both hPIGF and mPIGF, with a Biacore k_{off} rate of 80 and 35 min, respectively, indicating a more tight binding to hPIGF than mPIGF (data not shown). Additionally, RO5323441 binding experiments with RAW264.7 cells support the higher affinity for binding to hPIGF. This higher affinity explains the PIGF-specific uptake of ^{89}Zr -RO5323441 in the hPIGF-overexpressing Huh7 tumors. As expected, binding of ^{89}Zr -RO5323441 to Huh7 and ACHN tumor cells in vitro was low. mPIGF induced a small increase in ^{89}Zr -RO5323441 binding to Huh7 and ACHN cells. One explanation could be that we used hPIGF isoform 1 as hPIGF and mPIGF isoform 2 as mPIGF, because mice only express this isoform. One of the differences between PIGF1 and PIGF2 is that PIGF1 lacks a heparin-binding domain, which is necessary for binding to the neuropilin-1 coreceptor (18). mPIGF2 therefore could have been bound to Huh7 and ACHN cells via heparin, allowing binding of ^{89}Zr -RO5323441.

The in vitro experiments with RAW264.7 macrophages suggest that ^{89}Zr -RO5323441 forms immune complexes preferably with hPIGF. The ^{89}Zr -RO5323441 and hPIGF complex in turn binds to the Fc_γ receptors on macrophages, followed by phagocytosis of the immune complex; this is especially supported by the finding that ^{89}Zr -RO5323441 itself shows little interaction with macrophages. Only in the presence of hPIGF is there hPIGF concentration-dependent ^{89}Zr -RO5323441 binding. Because hPIGF is the target of RO5323441, competition of RO5323441 binding would intuitively be expected when hPIGF is added and thus decreased cellular binding. However, our results and those of others (19) provide evidence for the Fc_γ receptor-mediated macrophage uptake of IgG immune complexes, such as the complex of RO5323441 with hPIGF. This evidence can be explained by the fact that macrophages express Fc_γ receptor subtypes II and III ($\text{Fc}_\gamma\text{RII}$ and $\text{Fc}_\gamma\text{RIII}$), which have a low affinity for monomeric IgG. However, $\text{Fc}_\gamma\text{RII}$ and $\text{Fc}_\gamma\text{RIII}$ interact with high avidity with IgG immune complexes, with consequential selective phagocytosis of IgG immune complexes (19). The relevance of Fc_γ receptor interactions of antibody-antigen immune complexes in the tumor accumulation of soluble antigen-targeted antibodies such as RO5323441 is not exactly clear yet. Our Cy5-RO5323441 ex vivo data and ^{89}Zr -RO5323441 in vitro data, however, indicate that PIGF-specific tumor uptake of RO5323441 might be modulated by macrophages. The Fc_γ receptor-mediated uptake of antibody-antigen complexes is rather generic. It is therefore unlikely that this mechanism is specific for the RO5323441-PIGF complex. Tumor uptake of other soluble antigen-targeted antibodies such as the VEGF-A antibody bevacizumab might therefore also be dependent on immune complex formation. Bevacizumab-VEGF-A₁₆₅ immune complexes can interact with platelets via $\text{Fc}_\gamma\text{RIIA}$, potentially playing a role in the thrombotic events often seen during bevacizumab therapy (20).

Conflicting opinions exist on the significance of PIGF as a target for treatment in oncology (21). In preclinical models, PIGF plays a role in resistance to antiangiogenic therapies. PIGF inhibition showed antitumor activity in tumor models resistant to VEGFR inhibitors (7). Additionally, elevated serum levels of PIGF were reported in patients after antiangiogenic therapy with bevacizumab or sunitinib (22–24). In a preclinical study with the VEGF- and PIGF-neutralizing decoy receptor sFLT01, increased serum PIGF levels reflected a systemic host response instead of a tumor response. In addition, circulating PIGF was upregulated in mice with responding as well as progressing tumors (25). These findings suggest limited potential and utility of serum PIGF as a biomarker for antiangiogenic therapies. ^{89}Zr -RO5323441 tumor uptake was found to be PIGF-dependent. Therefore, ^{89}Zr -RO5323441 PET imaging might well be superior to measuring circulating PIGF levels to monitor PIGF tumor expression during antiangiogenic treatment. When the angiogenic rescue program is activated in the tumor, this will coincide with a local PIGF upregulation in the tumor microenvironment and an increased ^{89}Zr -RO5323441 uptake and possibly support the rationale for combining RO5323441 with VEGFR inhibitors. The vasculature normalization effect of bevacizumab can hamper the tumor penetration of simultaneously injected large molecules (e.g., antibodies such as RO5323441) (26). PIGF PET with ^{89}Zr -RO5323441 might therefore also be of value in the optimization of scheduling combined antiangiogenic treatments.

The therapeutic potential of PIGF inhibition recently has drawn additional attention. In contrast to the initially published antitumor effects of an anti-PIGF antibody, other PIGF-neutralizing antibodies had no significant effect on tumor growth and angiogenesis in several preclinical tumor models (27). There is no unifying explanation for these conflicting results; however, differences in physicochemical properties, posttranslational modifications, or other alterations may have influenced the efficacy of the different PIGF antibodies (8). Moreover, it was found that a functional VEGFR-1 in tumor cells is required for an antitumor effect of anti-PIGF antibodies (28). It is as of yet unclear how the antibody dose affects the efficacy (9). In the clinical phase I dose-escalation studies with RO5323441, no dose-limiting toxicities were found with doses of up to 30 mg/kg (10,11). It is possible that the maximum effective dose was already reached with optimal target saturation of its target. Insight in biodistribution of RO5323441 by quantification of ^{89}Zr -RO5323441 tumor uptake may offer an interesting support for this dosing dilemma by serving as readout for target saturation by different doses of RO5323441. Our results showed that a 20 mg/kg dose of RO5323441 reduced the human tumor uptake of the ^{89}Zr -RO5323441 tracer dose in mice to background ^{111}In -IgG levels. These results likely reflect PIGF saturation by the treatment dose but could also be a consequence of RO5323441-induced vessel normalization. Vessel normalization, which is now widely

acknowledged for bevacizumab, also occurs after PIGF inhibition (8,29). Dynamic contrast-enhanced MR imaging to monitor changes in tumor perfusion in the clinical evaluation of ^{89}Zr -RO5323441 and RO5323441 could potentially be used to distinguish PIGF saturation effects and vessel normalization.

CONCLUSION

The performance of ^{89}Zr -RO5323441 PET for PIGF detection is of interest because ^{89}Zr -bevacizumab and ^{111}In -bevacizumab have already proven to be valuable VEGF-A imaging tracers in clinical studies. The ^{89}Zr -bevacizumab uptake in tumor lesions in renal cell cancer and melanoma patients is beyond even the expectations based on preclinical results (13,14). Given the extensive similarity between VEGF-A and PIGF, an enhanced ^{89}Zr -RO5323441 tumor-to-background ratio versus findings in the xenograft model is expected also in the human setting. Because in the human setting both tumor cell- and stromal cell-derived PIGF are of human origin and will thus be recognized by ^{89}Zr -RO5323441, it is possible that our preclinical findings with ^{89}Zr -RO5323441 PET underestimate the tumor uptake that will be seen in cancer patients. In addition, the antibody tracer dose of 10 μg in a mouse of 25 g versus a typical dose of 5 mg in a human of 70 kg favors the antibody-antigen ratio in humans. To determine RO5323441 human tumor uptake and how this is affected by bevacizumab, we will quantify the tumor uptake of RO5323441 by serial ^{89}Zr -RO5323441 PET scans in patients with recurrent glioblastoma (EudraCT no. 2011-004974-27).

DISCLOSURE

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