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In vitro cytotoxicity of Auger electron-emitting [\(^{67}\text{Ga}\)]Ga-trastuzumab

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Introduction: Molecular radiotherapy exploiting short-range Auger electron-emitting radionuclides has potential for targeted cancer treatment and, in particular, is an attractive option for managing micrometastatic disease. Here, an approach using chelator-trastuzumab conjugates to target radioactivity to breast cancer cells was evaluated as a proof-of-concept to assess the suitability of \(^{67}\text{Ga}\) as a therapeutic radionuclide.

Methods: THP-trastuzumab and DOTA-trastuzumab were synthesised and radionlabelled with Auger electron-emitters \(^{67}\text{Ga}\) and \(^{111}\text{In}\), respectively. Radiopharmaceuticals were tested for HER2-specific binding and internalisation, and their effects on viability (dye exclusion) and clonogenicity of HER2-positive HCC1954 and HER2-negative MDA-MB-231 cell lines was measured. Labelled cell populations were studied by microautoradiography.

Results: Labelling efficiencies for \(^{67}\text{Ga}\)-Ga-THP-trastuzumab and \(^{111}\text{In}\)-DOTA-trastuzumab were 90% and 98%, respectively, giving specific activities 0.52 \(\pm\) 0.16 and 0.11 MBq/μg (78–92 GBq/μmol). At 4 nM total antibody concentration and 200 \(\times\) 10^3 cells/mL, \(^{67}\text{Ga}\)-Ga-THP-trastuzumab showed higher percentage of cell association (10.7 \(\pm\) 1.3%) than \(^{111}\text{In}\)-DOTA-trastuzumab (6.2 \(\pm\) 1.6%; \(p = 0.01\)). The proportion of bound activity that was internalised did not differ significantly for the two tracers (62.1 \(\pm\) 1.4% and 60.8 \(\pm\) 15.5%, respectively). At 100 nM, percentage cell binding of both radiopharmaceuticals was greatly reduced compared to 4 nM and did not differ significantly between the two (1.2 \(\pm\) 1.0% \(^{67}\text{Ga}\)-Ga-THP-trastuzumab and 0.8 \(\pm\) 0.9% for \(^{111}\text{In}\)-DOTA-trastuzumab). Viability and clonogenicity of HER2-positive cells decreased when each radionuclide was incorporated into cells by conjugation with trastuzumab, but not when the same level of radioactivity was confined to the medium by omitting the antibody conjugation, suggesting that \(^{67}\text{Ga}\) needs to be cell-bound or internalised for a therapeutic effect. Microautoradiography showed that radioactivity bound to individual cells varied considerably within the population.

Conclusions: \(^{67}\text{Ga}\)-Ga-THP-trastuzumab reduced cell viability and clonogenicity only when cell-bound, suggesting \(^{67}\text{Ga}\) holds promise as a therapeutic radionuclide as part of a targeted radiopharmaceutical. The causes and consequences of non-homogeneous uptake among the cell population should be explored.

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1. Introduction

With regulatory approval of beta particle-emitting radiopharmaceuticals such as \(^{[177}\text{Lu}]\text{Lu-DOTATATE}\) [1], early successes in trials of alpha particle-emitting therapy \(^{223}\text{Ra}\) [2] and an increasing choice of therapeutic radionuclides, chelators and radiochemistry techniques to choose from, this is an exciting time for molecular radiotherapy. Examples of therapeutic radiopharmaceuticals in routine clinical use include \(^{131}\text{I}\) for thyroid cancer, \(^{[111}\text{In}]\text{mIBG}\) for neuroblastoma, \(^{90}\text{Y}\)- and \(^{177}\text{Lu}\)-radiopeptides for neuroendocrine tumours, and \(^{90}\text{Y}\)-labelled microspheres (albeit defined for regulatory purposes as a medical device rather than a radiopharmaceutical) for hepatic tumours [3,4]. Most prominent among new beta particle-emitting radiopharmaceuticals undergoing clinical trials is \(^{[177}\text{Lu}]\text{Lu-PSMA}\) [5]. The potential of high linear energy transfer alpha particle-emitters, such as \(^{213}\text{Bi},\ 225\text{Ac},\ 211\text{At},\ \text{and}\ 227\text{Th}\) [6–8] to treat low volume tumours effectively while limiting healthy tissue toxicity is being explored. By comparison with alpha particles (<100 μm particle range), the short range (often <1 μm) of Auger electrons and related secondary electrons would be expected to increase the therapeutic ratio and further mitigate non-target toxicity [9]. Although likely to be less effective than beta emitters in large tumours because of the lack of “cross-fire” contribution to the tumour radiation dose, Auger electron-emitters could be effective against single cells and micrometastases which are ineffectively treated by beta-
emitters [10] and are responsible for recurrence of several tumour types [11,12].

Among therapeutic Auger electron-emitters, $^{111}$In has so far been closest to clinical translation and has been evaluated in both breast cancer [13] and neuroendocrine tumours [14]. $^{67}$Ga, which has been used clinically for SPECT imaging of inflammation and infection [15,16] also emits Auger electrons but its use as a therapeutic radionuclide is relatively unexplored [17–22]. Despite promising in vitro results, including its use in a radiopharmaceutical targeted towards the CD74 receptor on human B-lymphoma cells [21,22], low specific activities and a lack of purpose-designed gallium chelators have hindered progress with $^{67}$Ga therapy. The recent development of several new chelators [23–30], including the tris(hydroxypyridinone) chelators, allow kit-based radiolabelling of biomolecules, including proteins, with gallium radionuclides at high specific activities [31–33]. This development provides renewed opportunities for clinical translation to assess the tumouricidal potential of $^{67}$Ga preclinically and clinically as a radionuclide for molecular radionuclide therapy. Our recent studies have shown that $^{67}$Ga caused more DNA damage per Bq than $^{111}$In in a cell-free system. The cellular radioactivity required to kill 50% and 90% of HCC1954 breast cancer cells was 6- and 1.5-fold less, respectively, for $^{67}$Ga than for $^{111}$In [9,34]. Lower plasmid DNA damage was caused by $^{67}$Ga than $^{111}$In when the radionuclides were physically separated from the DNA, suggesting that $^{67}$Ga would cause less unwanted damage to surrounding non-targeted cells than $^{111}$In [34]. These data suggest that $^{67}$Ga could be selectively cytotoxic if targeted to tumour cells, with minimal damage to nearby non-targeted cells.

A widely explored tumour-specific molecular target is HER2, a receptor expressed on breast cancer cells. The antibody trastuzumab binds to HER2 and so is a potential basis for targeted delivery of radionuclides and evaluation of the therapeutic potential of $^{67}$Ga. Even in patients with primary tumours classified as HER2-negative, their metastases can still prove HER2-positive [35]. HER2-positivity in patients with breast cancer is associated with aggressive disease, poor prognosis and shortened overall survival [36]. A combination of trastuzumab, pertuzumab and docetaxel was able to partially overcome treatment failure with trastuzumab alone [37] was 1010 ± 388 MBq/mL [111In]InCl3 (110 MBq). The mixture was left for 15 min at room temperature. Unchelated $^{67}$Ga was scavenged with 2 μL 50 mM EDTA. Alteratively, $^{67}$GaCl3 (58.3 ± 11.9 MBq in 80 μL) was added to 100 μg THP-trastuzumab in 0.2 M ammonium acetate (3.5 mg/mL), pH was adjusted to 6.5 with 0.1 M NaHCO3 and was almost always >95%. Preparations in which the radiochemical purity was ≥95% were purified using a PD-10 column in 1 mL fractions of PBS containing 0.1% BSA.

2.2. Chelator conjugation to trastuzumab

Trastuzumab was conjugated to chelators tris(hydroxypyridinone) (H3-THP-Ph-SCN; THP, synthesised as previously described [40]) and S-2-(4-isothiocyanatobenzyl)-1,4,7,10-tetraazacyclododecane tetracetic acid (p-SCN-Bn-DOTA, Macrocycles, USA; DOTA) for radiolabelling with $^{67}$Ga and $^{111}$In, respectively.

2.2.1. THP-trastuzumab

Trastuzumab (435 μL, 22.96 mg/mL) in physiological saline was incubated with 50 mM ethylenediaminetetraacetate sodium (EDTA) solution for 30 min to remove any adventitious metal ions. The mixture was filtered at 4000 g in a Vivaspin 2 filter (Sartorius, Germany). The filter was washed with metal-free HEPES and the antibody thus recovered had a concentration of 12.46 ± 0.05 mg/mL (as measured by NanoDrop spectrophotometer) in metal-free HEPES. This solution of trastuzumab (2.32 mg, 260 μL, pH 8.7) was incubated with THP (21 μL of 10 mM solution in DMSO) at 10:1 chelator:antibody molar ratio at room temperature (2 h) and then 4°C (2 h). The mixture was applied to a Sephadex PD MidiTrap G-25 size exclusion column (GE Healthcare) and eluted with 0.2 M ammonium acetate. The fraction eluting between 1 mL and 2.5 mL was applied to a Sephadex PD-10 size exclusion column (GE Healthcare), and eluted with 0.2 M ammonium acetate solution in 500 μL fractions. Fraction 8 (protein concentration 3.5 mg/mL, measured by NanoDrop) was used for radiolabelling.

2.2.2. DOTA-trastuzumab

850 μL trastuzumab (23.5 mg/mL in saline) was dialysed overnight in a 3.5 kDa MWCO Slide-a-lyzer™ cassette (Thermofisher, USA) against 0.2 M, pH 5.5 sodium acetate containing 2 g/L chelax (Sigma, UK). DOTA conjugation to trastuzumab (10 mg, 12.8 mg/mL) was achieved by incubating a reaction solution containing a 40:1 M ratio of chelator to antibody, i.e. 1.1 mg DOTA (p-SCN-Bn-DOTA, Macrocycles, USA; 110 mg/mL in 0.25 M pH 5.5 sodium acetate) to which 83 μL 1 M NaHCO3, pH 9.5 was added and the mixture stirred for 60 min at room temperature. The conjugated antibody was dialysed against 0.2 M, pH 5.5 ammonium acetate containing 2 g/L chelax to remove unbound p-SCN-Bn-DOTA, furnishing a 13.5 mg/mL DOTA-trastuzumab solution (measured by NanoDrop).

2.3. Radiolabelling and quality control

2.3.1. $^{67}$Ga[Ga-THP-trastuzumab

$^{67}$Ga[Ga-citrate (48.6 Bq/mL) was added as aliquots of 25 μL (1.2 MBq), 100 μL (4.9 MBq) and 200 μL (9.7 MBq) to 100 μg THP-trastuzumab in 0.2 M ammonium acetate (3.5 mg/mL). pH was adjusted to 6.5 with 0.1 M NaHCO3. The mixture was left for 15 min at room temperature. Unchelated $^{67}$Ga was scavenged with 2 μL 50 mM EDTA. Alternatvily, $^{67}$GaCl3 (58.3 ± 11.9 MBq in 80 μL) was added to 100 μg THP-trastuzumab in 0.2 M ammonium acetate (3.5 mg/mL), pH was adjusted to 6.5 with 0.1 M NaHCO3 and the mixture was left for 15 min at room temperature.

2.3.2. $^{111}$In[In-DOTA-trastuzumab

$^{111}$In[InCl3 (94.1 ± 29.4 MBq in 120 ± 30 μL) was mixed with 250–430 μL 0.1 M ammonium acetate, pH 5.5, and added to 150 μg DOTA-trastuzumab. The mixture was kept at 40°C for 2 h. Unchelated $^{111}$In was scavenged with 2 μL of 50 mM EDTA.

2.3.3. Quality control

Labelling efficiencies were determined by paper chromatography on Whatman 1 paper (1 cm × 6.5 cm) with a mobile phase of 1 mM DTPA and were almost always >95%. Preparations in which the radiochemical purity was ≥95% were purified using a PD-10 column in 1 mL fractions of PBS containing 0.1% BSA.
2.4. Cell culture

MDA-MB-231 (HER2-negative) and HCC1954 (HER2-positive) breast cancer cell lines were grown as monolayers at 37 °C in a humidified atmosphere with 5% CO₂ in high glucose (4.5 g/L) Dulbecco’s Modified Eagle Medium (DMEM; PAA Laboratories) and RPMI-1640, respectively. Media were supplemented with 1.5 mM l-glutamine (PAA Laboratories, Austria), 10% foetal bovine serum (Invitrogen) and penicillin/streptomycin (Invitrogen) to create full medium. Cells were harvested using trypsin-EDTA for experimental use.

2.5. Binding assay

5 × 10⁴ HCC1954 cells per well were seeded in a 6-well plate 24 h prior to the experiment and incubated in a 5% CO₂ incubator at 37 °C. 4 nM [⁶⁷Ga]Ga-THP-trastuzumab or [¹¹¹In]In-DOTA-trastuzumab (0.08 and 0.09 MBq, respectively) were added to cells in a final volume of 250 μL full medium followed by incubation for 1 h at 37 °C. Medium was collected and combined with supernatant from two subsequent PBS wash steps to measure unbound activity. Cells were lysed in their wells with 1 mL 0.1 M NaOH and the solutions transferred to counting vials with 1 mL 0.1 M NaCl and 200 mM NaOAc (pH 2.5) for 5 min at 4 °C [41]. Cells were subjected to an acid wash by resuspending in 1 mL 500 mM PBS wash steps after centrifuging at 1500 rpm for 5 min. Cell pellets were subjected to an acid wash by resuspending in 1 mL 500 mM NaCl and 200 mM NaOAc (pH 2.5) for 5 min at 4 °C [41]. Cells were then centrifuged to separate membrane-associated activity (supernatant) from internalised activity (pellet). Radioactivity was measured as described in the methods section.

2.6. Internalisation assay

1 × 10⁶ HCC1954 cells in suspension were treated with 4 nM [⁶⁷Ga]Ga-THP-trastuzumab or [¹¹¹In]In-DOTA-trastuzumab (0.08 and 0.09 MBq, respectively) in 1 mL medium for 1 h at 37 °C. Unbound activity in the medium was collected and combined with washings from two PBS wash steps after centrifuging at 1500 rpm for 5 min. Cell pellets were subjected to an acid wash by resuspending in 1 mL 500 mM NaCl and 200 mM NaOAc (pH 2.5) for 5 min at 4 °C [41]. Cells were then centrifuged to separate membrane-associated activity (supernatant) from internalised activity (pellet). Radioactivity was measured as described above.

2.7. Viability

[⁶⁷Ga]Ga-THP-trastuzumab or [¹¹¹In]In-DOTA-trastuzumab (0.18 MBq/μg, 27 GBq/μmol) was added to 50 × 10³ HCC1954 cells in 250 μL full medium per well at 0, 4, 10, 40 and 100 nM, i.e. 0, 0.02, 0.06, 0.2 and 0.6 MBq, respectively. Controls included untreated HCC1954 cells, HCC1954 cells incubated with 100 nM non-radiolabelled trastuzumab, non-cell-binding activity (⁶⁷Ga]Ga-THP and [¹¹¹In]In-DOTA; 0.5 MBq) in HCC1954 cells, and HER2-negative MDA-MB-231 cells incubated with 4 nM [⁶⁷Ga]Ga-THP-trastuzumab or [¹¹¹In]In-DOTA-trastuzumab.

2.8. Clonogenic assay

2 × 10³ cells per well were plated in 6-well plates 24 h prior to the experiment. HCC1954 cells were incubated with 0.18 MBq/μg (27 GBq/μmol) radiolabelled trastuzumab in 500 μL for 1 h at 37 °C. The radioactive supernatant was then removed and cells were incubated in 2 mL full medium in a 5% CO₂ incubator at 37 °C. The medium was changed every 3 days. After 9–14 days, cells were fixed and stained with 1:1 methanol/1% crystal violet (Sigma, UK). Surviving fractions were calculated by counting colonies consisting of ≥50 cells. Controls included HCC1954 cells incubated with 100 nM non-radiolabelled trastuzumab, non-cell-binding radiopharmaceutical ([⁶⁷Ga]Ga-THP and [¹¹¹In]In-DOTA; at 1 MBq) and MDA-MB-231 cells incubated with 100 nM [⁶⁷Ga]Ga-THP-trastuzumab or [¹¹¹In]In-DOTA-trastuzumab. Clonogenic survival values were normalised to the untreated group (set at 1.0).

2.9. Microautoradiography

1 × 10⁶ HCC1954 cells in suspension were treated with 0 or 4 nM [⁶⁷Ga]Ga-THP-trastuzumab (0 and 0.04 MBq, respectively; 0.07 MBq/μg, 11 GBq/μmol) in 1 mL full medium. Preparations were incubated in a 5% CO₂ incubator at 37 °C for 1 h and washed in PBS, fixed in 4% formaldehyde in PBS for 20 min at room temperature, washed and resuspended in 200 μL PBS. Gelatine was dissolved in PBS in a microwave oven. Once cooled to about 37 °C, 0.4% of the gelatine solution was added to cells (200 μL) in a 1 cm × 1 cm mould and kept at 4 °C for 15 min. Once solidified, the gelatine block was trimmed to into cubes of 4 × 4 × 4 mm, which were snap frozen in isopentane in liquid nitrogen. Sections 10 μm thick were cut in a cryotome (Bright Instrument Company Ltd., UK) and placed on positively-charged polylysine-coated slides. In a dark room, the slides were coated by dipping in K2Ifford photographic emulsion (Ilford, Germany), which was prepared by adding 10 mL emulsion to 16.7 mL distilled water containing 0.01% glycerol and gently heating to 37 °C until dissolved. Dipped slides were left to dry horizontally at 4 °C for 24 h in a light-protected case, and developed at room temperature for 5 min in Kodak D19 (Kodak, USA; 15.8% (w/v) in distilled water). Slides were then dipped in 1% acetic acid for 30 s and then in 0.5% gelatin and dried at room temperature. Finally, autoradiographs were fixed in 30% sodium thiosulfate for 5 min, washed for 10 min and left to dry. Prior to imaging, 50 μL DAPI-containing mounting solution was applied to the slide and a coverslip was placed on top. Images were acquired with an EVOS FL Cell imaging microscope (Life Technologies, USA) and analysed with Image J (NIH, USA). The total number of silver grains around each cell was measured (within a circle 25 μm in diameter around the cell centre) in cells identified as DAPI-positive. The amount of signal in cells was corrected for background in an equivalent cell-free area of the image.

2.10. Statistical analysis

Statistical analysis was performed using Graphpad Prism Version 7.0c. An unpaired two-tailed t-test was used to analyse binding assay results comparing [⁶⁷Ga]Ga-THP-trastuzumab uptake with [¹¹¹In]In-DOTA-trastuzumab. A one-sample two-tailed t-test and a paired two-tailed t-test were used to analyse viability and survival fraction data. Data, except Fig. 1B, are average ± standard deviation.

3. Results

3.1. Radiolabelling

The labelling efficiencies for THP-trastuzumab labelling with [⁶⁷Ga] Ga-citrate were 100%, 84% and 33% for 25 μL (1.2 MBq), 100 μL (4.9 MBq), and 200 μL (9.7 MBq), respectively. This led to low specific activities of between 0.01 and 0.04 MBq/μg (2–6 GBq/μmol). However, radiolabelling THP-trastuzumab with [⁶⁷Ga]GaCl₃ achieved ~90% radiochemical yield and a specific activity of 0.52 ± 0.16 MBq/μg (78 GBq/μmol). [⁶⁷Ga]GaCl₃ was therefore used for all further radiolabellings. [¹¹¹In]In-DOTA-trastuzumab gave a radio labelling yield of 98% with specific activity of 0.61 ± 0.11 MBq/μg (92 GBq/μmol).
[67Ga]Ga-THP-trastuzumab showed higher binding (10.7 ± 1.3%) than HER2-positive HCC1954 cells. At 4 nM total antibody concentration, [111In]In-DOTA-trastuzumab (0.61 ± 0.11 MBq/μg) speciﬁcally bound to HER2-positive HCC1954 cells at increasing trastuzumab concentrations (n = 3). B. Calculated mean activity per HCC1954 HER2-positive cell (Bq/cell) for [67Ga]Ga-THP-trastuzumab and [111In]In-DOTA-trastuzumab. C. Binding of [67Ga]Ga-THP-trastuzumab and [111In]In-DOTA-trastuzumab to HCC1954 cells at 4 and 100 nM. Controls include non-internalised activity ([67Ga]Ga-THP and [111In]In-DOTA; 0.5 MBq/μg) and HER2-negative MDA-MB-231 incubated with 4 nM [67Ga]Ga-THP-trastuzumab and [111In]In-DOTA-trastuzumab (control cells) (n = 3).

3.2. Binding and internalisation assay

Both [67Ga]Ga-THP-trastuzumab (0.52 ± 0.16 MBq/μg) and [111In]In-DOTA-trastuzumab (0.61 ± 0.11 MBq/μg) speciﬁcally bound to HER2-positive HCC1954 cells. At 4 nM total antibody concentration, [67Ga]Ga-THP-trastuzumab showed higher binding (10.7 ± 1.3%) than [111In]In-DOTA-trastuzumab (6.2 ± 1.6%; p = 0.02; 50 × 103 cells, Fig. 1A). The percentage binding of both preparations decreased with increasing trastuzumab concentration. On increasing total antibody concentration to 100 nM, binding of [67Ga]Ga-THP-trastuzumab and [111In]In-DOTA-trastuzumab was inhibited and reduced to 1.2 ± 1.0% and 0.8 ± 0.9% (p = 0.85), respectively, indicating target-speciﬁc binding. However, the total activity bound to cells was higher at the highest antibody concentrations (Fig. 1B). Binding of antibody-free [67Ga]Ga-THP and [111In]In-DOTA to HCC1954 cells was minimal (0.07 ± 0.02% and 0.03 ± 0.00%, respectively). Binding of [67Ga]Ga-THP-trastuzumab and [111In]In-DOTA-trastuzumab to HER2-negative MDA-MB-231 cells was 0.18 ± 0.10% and 0.21 ± 0.23%, respectively, again conﬁrming that binding was HER2-speciﬁc.

Binding of 4 nM [67Ga]Ga-THP-trastuzumab or [111In]In-DOTA-trastuzumab to 1 × 106 HCC1954 cells (19.71 ± 1.32% and 11.93 ± 3.32%, respectively) was higher than that of 5 × 106 cells (10.7 ± 1.3% and 6.2 ± 1.6%, respectively). Of the cell-bound fraction, 62.1 ± 14.4% and 60.8 ± 15.5% was internalised for [67Ga]Ga-THP-trastuzumab and [111In]In-DOTA-trastuzumab, respectively.

4. Viability (trypan blue staining)

Treatment of HER2-positive HCC1954 cells with 100 nM non-radiolabelled trastuzumab did not affect viability (relative viability 101 ± 3.7% compared to untreated control cells whose relative viability was deﬁned as 100%). Following treatment of HER2-positive HCC1954 cells with either [67Ga]Ga-THP-trastuzumab or [111In]In-DOTA-trastuzumab, their viability, as measured with trypan blue staining, decreased as the [67Ga]Ga-THP-trastuzumab concentration and the activity added to the medium (and hence the cell-bound activity per cell) increased (Fig. 2A, B). Treatment of HCC1954 cells with [67Ga]Ga-THP-trastuzumab at 100 nM (2 MBq/mL) gave an average cellular radioactivity of 0.14 Bq/cell and produced signiﬁcant reduction in cell viability (to 66.5 ± 4.8% of the control value which was deﬁned as 100 ± 8.6% at 0 nM antibody concentration, p = 0.007) (Fig. 2A–C). Under the same conditions, [111In]In-DOTA-trastuzumab gave an average cellular uptake of 0.10 Bq/cell, and reduced viability to 66.2 ± 6.7% of the control value. Thus, the effect of the two radiopharmaceuticals (at similar average cell-bound activity per cell) on viability was not signiﬁcantly different (p > 0.9). Treatment of HER2-positive cells with non-antibody-conjugated [67Ga]Ga-THP (which did not bind to cells) did not measurably reduce cell viability, while non-antibody-conjugated [111In]In-DOTA (which also was not cell-bound) marginally decreased cell viability to 85.2 ± 4.0% of the control (Fig. 2D). Thus, both radionuclides were signiﬁcantly less toxic to HER2-positive HCC1954 cells when not cell-bound (p = 0.0003 for [67Ga]Ga and p = 0.0146 for [111In]). The difference in toxicity of non-cell-bound [67Ga]Ga and [111In]In was not signiﬁcant (p = 0.09). The results suggest that [67Ga]Ga and [111In]In need to be bound to cells to achieve signiﬁcant reduction in cell viability.

4.1. Clonogenicity

Treatment of HER2-positive HCC1954 cells with 100 nM non-radiolabelled trastuzumab did not affect clonogenicity (relative clonogenicity 0.97 ± 0.13 compared to untreated control cells whose relative clonogenicity was deﬁned as 1.0). Treatment with either [67Ga]Ga-THP-trastuzumab or [111In]In-DOTA-trastuzumab decreased their relative clonogenicity as concentration of radiolabelled trastuzumab, and consequently the amount of activity per cell, increased (Fig. 3A–C). Treatment of HCC1954 cells with 100 nM [67Ga]Ga-THP-trastuzumab and [111In]In-DOTA-trastuzumab reduced the clonogenic fraction to 0.43 ± 0.04 and 0.60 ± 0.07, respectively, compared to the untreated control (1.0 ± 0.18; p = 0.002 and 0.01, respectively). The effect of [67Ga]Ga on clonogenicity, at similar average activity per cell, was signiﬁcantly greater than that of [111In]In (p = 0.01). Treatment of HER2-positive cells with non-antibody-conjugated (and hence non-internalised) [67Ga]Ga-THP or [111In]In-DOTA reduced clonogenicity only marginally (0.88 ± 0.08, p = 0.13 and 0.82 ± 0.09, p = 0.04, respectively) compared to untreated control HCC1954 cells (Fig. 3D). The difference between clonogenic fraction of cells treated with 100 nM [67Ga]Ga-THP-trastuzumab (0.43 ± 0.04) and [111In]In-DOTA-trastuzumab and (0.60 ± 0.07) and non-cell-bound radioactivity (0.88 ± 0.08 and 0.82 ± 0.09 for [67Ga]Ga and [111In]In, respectively) was also signiﬁcant (p = 0.01 and 0.05, respectively). Clonogenic survival was signiﬁcantly impaired by treatment with [67Ga]Ga-THP-trastuzumab even at very low average activity per cell (<0.05 Bq/cell) but the effect of increasing the activity per cell beyond this level was less dramatic additional effect (Fig. 3B, C).

5. Microautoradiography

Of the activity added in this experiment (0.04 MBq, 0.07 MBq/μg, 11 GBq/μmol), 14.86 ± 0.38% of [67Ga]Ga-THP-trastuzumab was cell-bound. Microautoradiography revealed that radioactivity attached to individual cells varied considerably among the cell population; 15% of cells were classiﬁed as unlabelled (≤1 silver grain per
Fig. 2. Viability assay (n = 3) using trypan blue 3 days after a one-hour incubation of [67Ga]Ga-THP-trastuzumab or [111In]In-DOTA-trastuzumab with HER2-positive HCC1954 cells at 0.18 MBq/μg over a range of concentrations (A), activity per mL during incubation (B), and activity bound per cell (C). Controls for the viability assay in D are non-internalised activity in HER2-positive HCC1954 cells incubated with [67Ga]Ga-THP and [111In]In-DOTA; 0.5 MBq, 2 MBq/mL), HER2-negative MDA-MB-231 cells incubated with 100 nM radiolabelled trastuzumab, and HER2-positive HCC1954 cells incubated with 100 nM non-radiolabelled trastuzumab (n = 6). (For interpretation of the references to color in this figure legend, the reader is referred to the web version of this article.)

Fig. 3. Clonogenic assay (n = 3) of HER2-positive HCC1954 cells incubated for 1 h with [67Ga]Ga-THP-trastuzumab or [111In]In-DOTA-trastuzumab at 0.18 MBq/μg over a range of concentrations (A), activity per mL during incubation (B), and activity bound per cell (C). In D, controls non-internalised activity in HER2-positive HCC1954 cells incubated ([67Ga]Ga-THP and [111In]In-DOTA; 1 MBq, 2 MBq/mL), HER2-negative MDA-MB-231 cells incubated with 100 nM radiolabelled trastuzumab, and HER2-positive HCC1954 cells incubated with 100 nM non-radiolabelled trastuzumab (n = 6).
cell, e.g. Fig. 4A), 40% showed low levels of labelling (≤10 silver grains per cell, e.g. Fig. 4B), 25% medium labelling (41–90 silver grains per cell, e.g. Fig. 4C) and 20% high labelling (≥91 per cell, e.g. Fig. 4D).

6. Discussion

This study extends our previous cell-free system studies using pBR322 plasmids and cells non-specifically labelled using [67Ga]Ga-oxine, which showed that 67Ga causes more DNA damage, and a greater decrease in cancer cell clonogenicity when bound to cells, than the same activity of 111In [34]. Here, we have exploited the targeted delivery of radionuclides to HER2-positive cells using the antibody trastuzumab to investigate the selectivity and cytotoxicity of targeted radionuclide therapy with 67Ga compared to 111In [10,42–44].

Initial binding studies were carried out to determine the selectivity of uptake and the concentrations of radiolabelled antibodies required to achieve sufficient uptake of radionuclide in cells to affect measurable impact on cell viability and clonogenicity.

Due to difficulty experienced in radiolabelling trastuzumab with 67Ga to a sufficiently high specific activity (i.e. comparable to that achievable with 111In) under mild conditions using the DOTA chelators, we used the gallium-specific chelator THP, which can be radiolabelled with gallium radionuclides in under 5 min, at pH 6.5 and room temperature [23] allowing mild and efficient radiolabelling of proteins [33]. Similar specific activities of 0.52–0.61 MBq/μmol (78–92 GBq/μmol) could thus be achieved for both [67Ga]Ga-THP-trastuzumab and 111In-DOTA-trastuzumab.

At both low and high concentrations of radiolabelled trastuzumab, the percent binding of [67Ga]Ga-THP-trastuzumab to HER2-positive HCC1954 cells was somewhat higher (about 1.5 to 1.7-fold) than that of [111In]In-DOTA-trastuzumab. The choice of radionuclide did not affect the fraction of cell-bound radioactivity that was internalised within the cell (as opposed to bound to the cell surface), with over 60% of cell-bound radioactivity being internalised in both cases.

Viability studies using trypan blue staining showed that 67Ga induced significant toxicity, but only if incorporated into the cell; at similar concentrations in the media, non-antibody-conjugated [67Ga]Ga-THP, which did not bind to cells, did not affect cell viability. [111In]In-DOTA, which did not bind to cells, diminished cell viability only slightly, possibly due to a higher fraction of radiation dose coming from gamma emissions in the case of 111In. This is consistent with our previous observations [34].

Both [67Ga]Ga-THP-trastuzumab and [111In]In-DOTA-trastuzumab at 100 nM (2.5 MBq/mL) decreased clonogenic survival significantly compared to untreated HCC1954 cells and compared to HCC1954 cells incubated with 100 nM non-radiolabelled trastuzumab (Fig. 3), suggesting that the mechanism of cell death is mediated by radiation damage. Interestingly, the trends seen in Fig. 3A–C match what one might expect for a survival curve of tumour cells affected through the bystander response [45]. However, further studies will need to be carried out to determine whether the bystander response is indeed activated by these Auger-emitting radiopharmaceuticals. Our data using 100 nM non-radiolabelled trastuzumab (14 μg/mL, 1 h incubation) are in agreement with previous findings that trastuzumab induces no change in viability of HCC1954 cells even after 7 days of incubation at 15 μg/mL [46].

At similar radioactivity concentrations in the medium, [67Ga]Ga-THP-trastuzumab is 1.4-fold more toxic (measured as effect on clonogenicity) than [111In]In-DOTA-trastuzumab (Fig. 3, A and B). It is likely that only part of this apparent difference in toxicity is associated with the 1.5–1.75-fold more activity of [67Ga]Ga-THP-trastuzumab taken up per cell, because when clonogenicity is plotted against the measured average activity per cell (Fig. 3, C), 67Ga still shows significantly greater suppression of clonogenicity than 111In.

The differences in toxic effects as measured by viability (trypan blue) and clonogenicity indicate that viability and clonogenic assays measure different aspects of cytotoxicity and are complementary rather than alternative methods.

The modest difference in toxicological profile between the two radioimmunoconjugates could originate from differences in the cellular trafficking, or from differences in the electron and photon emission spectra of the two radionuclides. The 67Ga conjugate shows greater apparent affinity and uptake than the 111In conjugate, and although this does not cause any significant difference in the measured internalised fraction, it implies that differences in subsequent intracellular trafficking, including to the cell nucleus, cannot be excluded. Electron emissions (Auger and Coster-Kronig electrons) of 67Ga and 111In are similar in total energy (6.24 MeV and 6.75 MeV, respectively) and both also emit internal conversion electrons (0.32 and 0.16 per decay respectively, with a total energy of 28.078 KeV and 25.957 KeV, respectively). However, the energy distribution, and hence range in tissue, of the Auger and Coster-Kronig electrons differs markedly between the two radionuclides. The Supplementary information provides relevant data in the form of plots of yield of electrons per decay against the logarithm of the range [47]. The total energy of the Auger and Coster-Kronig emissions of 67Ga is distributed among fewer electrons than is the case for 111In (4.7 and 14.7, respectively [47]) and consequently, electrons from 67Ga have higher energy per electron, and hence longer range. Fewer Auger electrons emissions per decay than 14.7 [47] have been estimated for 111In, namely between 5.84 and 7.43, however these values are still higher than for 67Ga (between 4.56 and 4.96) [48]. 67Ga emits no electrons with a range below 1 nm, whereas 111In emits 8; and 67Ga emits around 0.6 electrons per decay with a range encompassing multiple compartments of the cell (1–10 μm), whereas 111In emits none. Thus, 111In decay is likely to be biologically effective over a much smaller range (<1 μm) than 67Ga decay, and the biological effects of decay of 111In are likely to be much more dependent on its sub-cellular location than those of 67Ga.

Although DNA damage is known to be an important mechanism by which radionuclide decay induces cell killing, it is a currently open and actively investigated question whether the cell nucleus and DNA is the most important target for radionuclide emission, or whether other organelles such as the mitochondria or cell
membrane [49,50] are also important radiosensitive targets. Although not explored here, it is known that $^{111}$In–In–trastuzumab can localise to the nucleus [42,44]. The extent to which this happens is likely to be an important factor in the therapeutic efficacy of the radiopharmaceutical, considering the very short range of the majority of its electron emissions. Nuclear localisation peptide sequences have often been used to further increase therapeutic efficacies of Auger-emitting radiopharmaceuticals [42–44], including some labelled with $^{67}$Ga [51]. Although other non-nuclear-associated pathways are known that can influence therapeutic efficacy [50], and despite the longer range of the $^{67}$Ga electron emissions, it is possible that an approach using nuclear localisation sequences will also be of benefit for $^{67}$Ga–trastuzumab. The data presented here, however, do not provide insight into the mechanism of cell killing or sub-cellular target of these radioimmunoconjugates.

Microautoradiography demonstrates a wide range of $^{67}$Ga–THP–trastuzumab binding among HCC1954 cells, despite the presumption that the cells are a clonally identical population. In fact, 20% of cells in the highest activity group received a radiation dose that was 15% in the lowest activity group, 20% of cells in the highest activity group received around a hundred times higher radiation dose than the 15% in the lowest activity group, and a large sub-population of the cells would have received negligible radiation dose. This may account for the observation (Fig. 3B, C) that the surviving cell fraction does not greatly diminish below 0.5 as activity of $^{67}$Ga–THP–trastuzumab per cell increases above 0.05 Bq/cell. It implies that the average activity per cell is likely to have very limited utility as a predictor of toxicity. Heterogeneity of target expression is likely to be even greater in cell populations within patients’ tumours. This is likely to be the overriding limitation in targeted Auger emitter therapy. The impact of non-homogeneous radiopharmaceutical uptake in limiting anti-tumour effectiveness will need to be considered for any radiopharmaceutical undergoing in vitro and in vivo evaluation and requires further investigation if the antitumour effects of targeted Auger electron-emitting radioligands can be better understood.

7. Conclusion

$^{67}$Ga–THP–trastuzumab specifically reduced cell viability and clonogenicity of HER2-expressing cells, confirming the potential of $^{67}$Ga as a therapeutic radionuclide as part of a targeted radiopharmaceutical. Diminished viability and clonogenicity only occurred when radioactivity was directly bound to cells and was not caused by radioactivity in the medium. Highly heterogeneous cellular uptake may compromise the efficacy of Auger electron irradiation and the interpretation of data derived from average cellular radioactivity. This merits further investigation using methods, such as microautoradiography, capable of measuring the radioactivity of individual cells.

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