Research Article

Ga-68- and Cu-64-Labeled NOTA-Albumin Conjugates for PET Sentinel Lymph Node Imaging

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Our objective was to develop and evaluate Ga-68- and Cu-64-labeled albumin conjugates for PET imaging of sentinel lymph nodes. Four different albumin conjugates were prepared starting from NOTA-HSA. The lymph node uptake of Ga-68- and Cu-64-labeled albumin conjugates was investigated after subcutaneous injection into the foot pad of Wistar rats. A pig model was utilized for further biological evaluation of the lymph node uptake. For all the four conjugates, radiolabeling with Ga-68 and Cu-64 resulted in >95% radiochemical yield. Denatured and mannosylated Ga-68 NOTA-HSA revealed the highest popliteal lymph node uptake in rats (2.78 ± 0.38% ID and 6.13 ± 1.13% ID 10 min and 60 min p.i., resp.). The popliteal lymph node reached its maximum activity after approximately 120 min and remained constant for denatured and mannosylated Cu-64 NOTA-HSA at least up to 240 min p.i. In a pig model, 2% of the injected dose of this compound was found in the sentinel lymph node 60 min after subcutaneous injection. In conclusion, PET imaging of sentinel lymph nodes with Ga-68- and Cu-64-labeled denatured NOTA-Man-HSA could be successfully demonstrated and deserves further investigations.

1. Introduction

Sentinel lymph node (SLN) diagnosis is an important field in nuclear medicine, and its benefit has been proven in many cancer types, mainly breast cancer and melanoma. The sentinel lymph node is the first lymph node in the drainage basin of a primary tumor. SLN detection is a two-step procedure. In the first step, SLNs are imaged by means of planar lymphoscintigraphy. Recently, the advantage of SPECT/CT in cases with difficult anatomical location of sentinel nodes was reported as an alternative imaging modality [1, 2]. In the second step of the procedure, SLNs are intraoperatively detected by a hand-held gamma camera, excised and pathologically examined. If the sentinel node is free of metastatic cells, the likelihood of general nodal metastasis is very low.

In general, technetium-99m-labeled colloids in combination with blue dyes are used to detect the sentinel lymph node. The labeled colloids are injected beneath the skin surrounding the primary tumor or alternatively directly into the tumor. After entering the lymphatic vessels, colloids use the same drainage pathway as potential metastatic cancer cells and are eventually retained in the first draining lymph node (sentinel node) by phagocytosis or mechanical trapping. In Europe, mainly the colloids based on denatured human serum albumin are used, whereas in USA; sulphur colloid is preferred.

There are two crucial factors in the performance of Tc-99m colloids in SLN mapping. First, the agent has to be quickly transported from the site of injection to ensure the detection of SLNs in close proximity. Second, a high and persistent lymph node accumulation is necessary for
the imaging and the intraoperative detection of the nodes. Both processes are mainly influenced by the size of the particles. As a compromise between fast transport of small particles from the site of injection and high lymph node uptake of larger colloids, nowadays a particle size of 10–200 nm is favoured [3–6].

A tailor-made Tc-99m radiopharmaceutical for SLN diagnosis based on dextran (Lymphoseek) is currently in clinical phase III studies [7, 8]. In this case, 7 nm-sized dextran as carrier molecule is additionally loaded with mannosyl units to enhance the receptor-mediated uptake of the radiopharmaceutical into the lymph nodes. Preliminary data indicated a similar lymph node uptake like filtered sulphur colloid, whereas injection site clearance is significantly faster [9]. The development of a positron-emitting agent would further improve the image resolution and the localization of SLNs by using PET/CT. Especially for SLNs in the abdomen, this is crucial for radiotherapy planning [10]. A recent paper by Heuveling et al. describes the preclinical results of Zr-89 nanocolloidal albumin PET/CT in head and neck cancer [11]. Choi and coworkers utilized the generator-derived Ga-68 as PET nuclide and labeled mannosylated human serum albumin (MSA) for SLN detection in rodents [12].

Most of the currently available commercial technetium-99m kits for sentinel lymph node scintigraphy contain colloids of denatured albumin. In our study, we investigated the effect of human serum albumin (HSA) denaturation in combination with mannosylation and compared the biological performance of this agent with that of the nondenatured HSA as well as probes without mannosyl units. We provide data on the biodistribution of Ga-68-labeled conjugates and the long-term biokinetic studies of one conjugate labeled with Cu-64.

2. Materials and Methods

2.1. General. p-SCN-Bn-NOTA \times 3 \text{HCl} (NOTA-NCS) was purchased from Macrocyclics. Tetra-O-acetyl mannosyl isothiocyanate was synthesized according to the procedure of Kühne et al. [13].

HSA solution (20%) was purchased from C.A.F.-D.C.F., Belgium. Poloxamer 238 was purchased from BTC, Cologne, Germany. Gallium-68 was eluted from a Ge-68/Ga-68 generator (iThemba LABS, South Africa). Copper-64 was produced with a cyclotron CYCLONE 18/9 via the $^{64}$Ni(p, n)$^{65}$Cu reaction according to the method of Thieme and co-workers [14].

TLC plates were scanned on a TLC analyzer (RITA, Raytest, Straubenhardt, Germany). For ultrafiltration, a Heraeus Labofuge 400R was used with a Jumbosep centrifugal device (Pall) with a molecular weight cut-off of 30 kDa. Samples were centrifuged at 2500 min$^{-1}$ and 10°C for 60 minutes. Mass spectrometry was performed on an Autoflex MALDI-TOF/TOF (Bruker Daltonics) with dithyroxyl benzoic acid (NOTA-HSA) and 3-hydroxypropionic acid (NOTA-Man-HSA) as matrices. Particle size distribution was determined with a Zetasizer Nano ZS (Malvern). PET data of rats was acquired using a microPET P4 (Siemens preclinical solutions, Knoxville, TN, USA) scanner. For the pig experiment, a Sensation Biograph 16 Hi-REZ Siemens/CTI, Knoxville, TN, USA, was utilized.

2.2. Synthesis of NOTA-HSA. Two milliliters of a sodium tetraborate solution (0.1 M, pH 9.2) was added to 3.4 mg (6.1 \text{μmol}) p-SCN-Bn-NOTA \times 3 \text{HCl}. After complete dissolution of the solid in an ultrasonic bath, the solution was mixed with 100 μL HSA solution (20 mg; 0.3 μmol) in a 2 mL Eppendorf Protein LoBind Tube. The reaction mixture was treated in an Eppendorf Thermomixer at 30°C and 300 min$^{-1}$ for 18 h. Low molecular weight impurities were separated by ultrafiltration at 10°C (fivefold washing with water). The residual solution was filled into 10 mL glass vials and lyophilized. The number of conjugated NOTA per HSA was determined by MALDI-MS. The difference between the peak abundance of NOTA-HSA (71,725 Da) and native HSA (67,000) was divided by the MW of p-SCN-Bz-NOTA as trisodium salt due to the basic reaction mixture ($M = 626$ g/mol). Eight NOTA chelators per HSA molecule were calculated.

2.3. Conjugation of Mannose to NOTA-HSA. Tetra-O-acetyl mannosyl isothiocyanate (33 mg) and 30 mg NOTA-HSA lyophilisate were dissolved in 30 mL 0.25 M carbonate buffer pH 9.0. The reaction mixture was slowly stirred at room temperature for 18 hours. The solution was filtered with a syringe filter (pore size 0.45 μm). Unbound mannose was removed by ultrafiltration at 10°C (fivefold washing with water). The residual solution was filled into 10 mL glass vials and lyophilized. The number of conjugated mannosyl units was determined by MALDI-MS. The difference between the peak abundance of NOTA-Man-HSA (75,585 Da) and NOTA-HSA (71,725 Da) as a starting material was divided by the MW of mannosyl isothiocyanate ($M = 221$ g/mol). Seventeen sugar units per HSA molecule were calculated.

2.4. Denaturation of HSA Conjugates. In a volume of 20 mL water, 10 mg of the HSA conjugate, 40 mg of Poloxamer 238, and 11 mg of Na$_2$HPO$_4 \times 2$ H$_2$O were subsequently dissolved. The pH was adjusted to 7.3 with 0.1 M HCl, and the volume of the solution was adjusted with water to give a total of 40 mL. The solution was then heated at 80°C for 30 min. After cooling down to room temperature, the solution was filtered with a syringe filter (pore size 0.45 μm). Aliquots of 2 mL were filled into 10 mL glass vials, and the solvent was removed by lyophilization.

2.5. Ga-68 and Cu-64 Labeling of NOTA-HSA. NOTA-HSA$^{\text{denat}}$, NOTA-Man-HSA, and NOTA-Man-HSA$^{\text{denat}}$. One millilitre of ammonium acetate buffer (0.5 M, pH 5.0) was added to a 10 mL vial containing 0.5 mg of the HSA conjugate. Subsequently, 0.3 mL of Ga-68 eluate (100–250 MBq) was added. After 15 min at room temperature, 0.25 mL of aqueous EDTA solution (0.04 M, pH 4) was added to the reaction mixture. Five minutes later, the radiochemical yield was determined by thin layer chromatography with chromatography paper MN128 (Macherey-Nagel) as a stationary phase and 0.1 M citrate buffer pH 5.0 as a mobile
phase. Radiolabeled HSA conjugates remain at the starting point, whereas unbound radionuclide (as citrate or EDTA complex) and Ga-68 NOTA-NCS move with the solvent front ($R_f = 0.9 – 1.0$). The size of the radiolabeled conjugates was analyzed by the filtration of an aliquot through a filter with 80 nm pore size (Whatman). In an analogous manner, NOTA-Man-HSA$_{denat}$ was labeled with Cu-64 by adding 100 μL of (ca. 100 MBq) $[^{64}\text{Cu}]\text{CuCl}_2$ solution (0.1 N HCl). For in vivo experiments, the addition of EDTA was omitted, whereas 1 mL of HEPES (1 M, pH 8) was added to the reaction mixture for pH adjustment.

2.6. In Vivo Studies. The animal research committee of the Regierungspräsidium Dresden approved the animal facilities and the experiments according to the institutional guidelines and the German animal welfare regulations. The experimental procedure used conforms to the European Convention for the Protection of Vertebrate Animals Used for Experimental and Other Scientific Purposes (ETS no. 123), to the Deutsches Tierschutzgesetz, and to the Guide for the Care and Use of Laboratory Animals published by the US National Institutes of Health (DHED Publication no. (NIH) 82-23, Revised 1996, Office of Science and Health Reports, DRR/NIH, Bethesda, MD, USA). The Wistar rats (Wistar Unilever, HsdCpb: Wu, Harlan Winkelmann GmbH, Borchen, Germany, 100–150 g body weight) were housed under standard conditions with free access to standard food and tap water.

The biodistribution of Ga-68- and Cu-64-labeled HSA conjugates was studied after subcutaneous application into the foot pad or i.v. application.

The biodistribution of the radiolabeled compounds was studied in 4 male rats of each group at 10 and 60 min after tracer injection. The animals were anesthetized with desflurane (Suprane, Baxter Healthcare Corporation, Deerfield, IL, USA) (7.0%–10.0% v/v in 30% oxygen), and i.v. Ga-68 108 ± 20 MBq/kg BW (BW 135 ± 28 g), i.v. Cu-64 4.0 ± 0.4 MBq/kg BW (BW 118 ± 3 g), and intrafoot Ga-68 55 ± 8 MBq/kg (BW 130 ± 21 g), and intrafoot Cu-64 6.4 ± 0.4 MBq/kg (BW 136 ± 7 g) aliquots were administered i.v. into a tail vein in 500 μL and subcutaneously into the foot in 100 μL isotonic NaCl or electrolyte solution E153 (Serumwerk Bernburg AG, Bernburg, Germany). After the recovery from anesthesia, rats were again anesthetized at 10 or 60 min after tracer injection, respectively. The blood was withdrawn by heart puncture, and the animals were euthanized. Organs and tissues were removed, dried, and weighed, and the radioactivity was measured in a cross-calibrated well counter (WIZARD, Automatic Gamma Counter, Perkin Elmer, Waltham, MA, USA) or dose calibrator (Aktivimeter Isomed 2000; MED-Nuklear-Medizintechnik, Dresden, Germany). The data were decay corrected and normalized to the amount of injected activity calculated from the activity of injection syringes before and after injection and expressed as the percentage of injected activity (%ID) or the standardized uptake values in biodistribution studies (SUVPET, SUVPET = (activity/mL tissue)/(injected activity/body weight, mL/g)) were calculated in ROI.

The pig experiment was performed according to a previously published procedure by Tafra et al. [15]. Briefly, an anesthetized pig was administered intradermally with 45 MBq Ga-68 NOTA-Man-HSA$_{denat}$ in 0.5 mL. The injection site was placed just distal to the hind and forelimb joints of each hind leg. Injection was made at four points equidistant around the mock site in each extremity. PET/CT scans were acquired 15 min and 60 min after injection.

3. Results and Discussion

3.1. Synthesis of NOTA-HSA, NOTA-HSA$_{denat}$, NOTA-Man-HSA, and NOTA-Man-HSA$_{denat}$. All conjugates were prepared starting from NOTA-HSA which was synthesized by...
conjugation of an isothiocyanate derivative of NOTA to ε-amino groups of HSA lysine side chains. The number of conjugated chelators was determined by MALDI-MS, comparing between the mean molecular weights of NOTA-HSA and native HSA and dividing the difference by the MW of NOTA-NCS. By using a 10-fold or 20-fold molar excess of NOTA, approximately 4 and 8 chelators could be conjugated per albumin, respectively. In subsequent studies, it turned out that using albumin with 4 NOTA units results in low radiochemical yield (ca. 87%) of the corresponding denatured conjugates. Therefore, further studies were conducted by using albumin with 8 conjugated chelators.

In the second step, in a similar manner, tetra-O-acetyl mannosyl isothiocyanate was bound to NOTA-HSA. A 200-fold molar excess of mannose was used to fix as much mannose as possible to all sterically available ε-amino groups of lysine side chains. The O-acetyl groups of the sugar are cleaved under the used reaction conditions (0.25 M carbonate buffer pH 9). This was proven by preliminary experiments with penta-O-acetyl mannose treated under identical conditions. 1H NMR of the reaction mixture of that compound was performed to show that no more proton signals of mannose-O-acetyl CH₃ groups were found. Instead, a singlet for acetate CH₃ was detected. For mannosylated NOTA-HAS, the number of attached mannose units was again calculated from the results of MALDI-MS. An average of 17 sugars per NOTA-HSA was calculated. In the work of Choi and co-workers, 7 NOTA chelators and 11 mannose units were conjugated to native albumin, we determined an average size of 5.8 ± 1.5 nm.

3.2. Ga-68 and Cu-64 Labeling of HSA Conjugates. Ferreira et al. reported on the labeling of NOTA with Ga-68 in acetate buffer [16]. A pH of 4-5 and a reaction time of 5 min (room temperature) were found to be optimal conditions for labeling. According to this published data, we performed Gallium-68 labeling in a similar manner using 0.5 M acetate buffer pH 4, 5, and 6, respectively. A buffer pH of 5 revealed the highest labeling yield. After 15 min at room temperature, the labeling yield after EDTA quenching of the remaining [Ga-68]Ga³⁺ exceeded 95% for NOTA-HSA, NOTA-HSA.denat, NOTA-Man-HSA, and NOTA-Man-HSA.denat, determined by thin layer chromatography.

To verify the particle size distribution of our labeled HSA conjugates, we filtered an aliquot of the labeling mixture. For these experiments, we used polycarbonate filters with a defined pore size of 80 nm. For all the four conjugates, 97%–99% of the radioactivity passed the filter membrane and was detected in the eluate, indicating a particle size smaller than 80 nm. HSA colloids were labeled with Cu-64 under identical
conditions with the same high yield. For animal experiments, 1 mL of HEPES buffer (1 M, pH 8) was added to the reaction mixture after labeling, resulting in a pH of ca. 7. This does not affect the integrity and the purity of the product, as determined by TLC and filtration.

3.3. Biodistribution Studies of Ga-68- and Cu-64-Labeled HSA Conjugates. The rat model was used to analyze the lymphatic transport of the tracers and their uptake in lymph nodes [17, 18]. Labeled colloids were subcutaneously injected into the foot pad of rats. Colloids are transported to some extent via the lymph vessels. The first passed lymph node (sentinel lymph node) is the popliteal node located behind the knee joint [19]. Analogously to Weiss et al. [18], we separated the whole knee to determine the activity of the popliteal lymph node. Biodistribution was analyzed 10 and 60 min after injection (Figure 2).

Noticeable activity was found in the foot pad (injection site), knee (popliteal lymph node), and liver. Activity in other organs and tissues was marginal. For all the tested compounds, the activity in the foot pad decreased by ca. 20%–25% ID from 10 to 60 min p.i., whereas an increase is observed in the popliteal lymph node (knee) and the liver. The uptake in the liver and the transport from the injection site are similar for all conjugates. On the other hand, differences are seen in the lymph node uptake. According to a receptor-mediated mechanism triggered by mannose, conjugates containing mannose are preferably taken up by phagocytosis in comparison to the corresponding conjugate (denatured/nondenatured) without mannose units. Furthermore, denaturation by heat treatment seems to have a positive influence on the uptake of the mannosylated conjugate 60 min p.i. compared to its nondenatured counterpart, although the difference is rather small. Among the four tested conjugates, NOTA-Man-HSA\textsubscript{denat} revealed the highest lymph node uptake of 2.78% ± 0.38% ID and 6.13% ± 1.13% ID 10 and 60 min p.i., respectively. This compound was subsequently labeled with Cu-64. Cu-64 was utilized to further characterize the colloids, taking advantage of better image resolution and longer half-life of Cu-64 in comparison with Ga-68. Essentially, the same biodistribution patterns were found after subcutaneous injection of Cu-64- and Ga-68-labeled NOTA-Man-HSA\textsubscript{denat} into the foot pad (Figure 3), indicating that the radiolabeling with either of the two radionuclides does not significantly alter the biodistribution. The small differences between the Ga-68- and the Cu-64-labeled agent in Figure 3 are most probably due to different mobility behavior of the rats in both groups. The transport of colloids within the lymph vessels is promoted by muscle contraction. Figure 5 reveals more precise results under equivalent conditions for both groups, because in this experiment animals were anaesthetized throughout the whole experiment (vide infra).

In further experiments, PET data of Ga-68- and Cu-64-labeled denatured NOTA-Man-HSA were acquired up to 120 and 240 min p.i., respectively (Figure 4). As expected, both tracers show similar distribution patterns, reflecting the results of the biodistribution studies. The main activity is found in the injection site (foot). At later time points, also
Ga-68 NOTA-Man-HSA<sub>delt</sub>

5 min  60 min  120 min

Scatter artefacts

Bladder

Popliteal LN

Injection site

Cu-64 NOTA-Man-HSA<sub>delt</sub>

5 min  120 min  240 min

Second echelon LNs

Bladder

Popliteal LN

Injection site

Figure 4: Coronal PET images of Ga-68- and Cu-64-labeled denatured NOTA-Man-HSA after subcutaneous injection into the foot pad of rats.

![Figure 4: Coronal PET images of Ga-68- and Cu-64-labeled denatured NOTA-Man-HSA after subcutaneous injection into the foot pad of rats.](image)

the bladder becomes slightly visible. The popliteal lymph node is visible 60 and 120 min p.i. for the Ga-68-labeled compound and 120 min and 240 min p.i. in the case of Cu-64. The normalized activity curves show a slow radioactivity accumulation within the lymph node. The maximum activity plateau is reached after approximately 120 min in all tested individuals, independent of the radiolabel (Figure 5).

![Figure 5: Normalized time-dependent radioactivity uptake in the foot pad and the popliteal lymph node after subcutaneous injection of Ga-68 (n = 2) and Cu-64 (n = 4) labeled denatured NOTA-Man-HSA into the foot pad of rats.](image)

Copper-64 images are characterized by higher resolution and lower scatter artefacts due to the lower $\beta^+$-energy in comparison to Ga-68. This might also be the reason why in the case of Cu-64 second echelon lymph nodes become visible 120 min p.i.

Lymph node uptake of Ga-68 NOTA-Man-HSA after subcutaneous injection is also presented by Choi et al. [12]. Their
data are obtained from dynamic micro-PET imaging up to 10 min p.i. A more or less constant popliteal uptake of approx. 5% ID is observed as soon as 1 min p.i., which is unusual for a biological transport mechanism and is not in accordance to our finding of slow increase of activity accumulation within the lymph node (Figure 5). Data on the uptake up to several hours p.i. are important for intraoperative detection of lymph nodes by hand-held beta probes. In contrast to Tc-99m-labeled colloids where lymph node dissection is usually performed 15–24 h p.i. [5, 20], intraoperative node detection has to be performed immediately after imaging in the case of gallium-68 due to its shorter physical half-life. We demonstrated high and persistent lymph node uptake of denatured NOTA-Man-HSA up to 240 min p.i.

Fast transport from the injection site is another crucial factor. For example, in a study of Vermeeren et al. in 6 out of 46 patients, sentinel lymph nodes could not be detected by planar imaging due to their close proximity to the injection area [2]. In our studies, the activity of denatured NOTA-Man-HSA at the site of injection decreased from 88.7% ± 4.5% ID 10 min p.i. to 64.9% ± 17.5% ID. This is significantly lower than that of the currently used Tc-99m-labeled colloid, where almost the entire injected dose remains at the injection site [20].

Furthermore, a pig model was utilized to create PET/CT images of the lymphatic transport and lymph node uptake. The conjugate with the highest uptake in the rat popliteal lymph node, Ga-68-labeled NOTA-Man-HSA\textsubscript{denat}, was injected in both extremities according to the procedure of Tafra et al. [15]. Clearly, the lymphatic transport and the uptake in the sentinel lymph nodes could be visualized on early images 15 min p.i. (Figure 7).

The mean uptake of both injections into the sentinel node was 1.2% after 15 min and 2.0% after 60 min. In another published study in which a pig model was used, lymph node uptakes of 1.32%–2.04% ID were found for Lymphoseek and 0.63%–2.35% ID for filtered Tc-99m sulphur colloid [21]. In comparison, average lymph node uptakes of 1% ID for HSA nanocolloids [20], 0.65% ID [22] and 0.85% ID for sulphur colloids [23], and 0.73% ID for Lymphoseek [23] are reported for patients with malignant melanoma, mammalian carcinoma, or prostate cancer. These similar values can be explained by saturation of the limited number of phagocytic cells (macrophages or dendritic cells) within the lymph node [24].

4. Conclusion

Four different conjugates based on NOTA-human serum albumin have been synthesized and labeled with Ga-68 and Cu-64, respectively. The effects of mannosyl units and denaturation were studied after subcutaneous injection into the foot pad of rats. Distinct uptake in the popliteal lymph node was observed for all tested compounds. Mannosylation could be identified to positively influence the uptake in the lymph node. The effect of denaturation by heat treatment in the case of mannosylated compounds is rather low. However, heat-treated HSA colloids bearing mannosyl units (NOTA-Man-HSA\textsubscript{denat}) revealed the highest lymph node uptake of 6.13%±1.13% ID 60 min p.i. After the intravenous injection in
rats, the majority of Ga-68- and Cu-64-labeled NOTA-Man-HSA\textsubscript{denat} were taken up quickly by the reticuloendothelial system of the liver.

PET experiments in rats revealed similar distribution pattern for Ga-68- and Cu-64-labeled NOTA-Man-HSA\textsubscript{denat}. The first lymph node in the drainage basin reaches its maximum activity after approx. 120 min and remained constant for the copper-64 compound at least up to 240 min p.i.

The feasibility of lymphatic mapping with Ga-68-labeled NOTA-Man-HSA\textsubscript{denat} was furthermore demonstrated by PET/CT imaging using a pig model, where 2% of the injected dose was found in the sentinel lymph node 60 min p.i.

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