Comparison of image quality and spatial resolution between $^{18}$F, $^{68}$Ga, and $^{64}$Cu phantom measurements using a digital Biograph Vision PET/CT

Anja Braune*1, Liane Oehme1, Robert Freudenberg1, Frank Hofheinz2, Jörg van den Hoff2, Jörg Kotzerke1,2,3 and Sebastian Hoberück1,4

Abstract

Background: PET nuclides can have a considerable influence on the spatial resolution and image quality of PET/CT scans, which can influence diagnostics in oncology, for example. The individual impact of the positron energy of $^{18}$F, $^{68}$Ga, and $^{64}$Cu on spatial resolution and image quality was compared for PET/CT scans acquired using a clinical, digital scanner.

Methods: A Jaszczak phantom and a NEMA PET body phantom were filled with $^{18}$F-FDG, $^{68}$Ga-HCl, or $^{64}$Cu-HCl, and PET/CT scans were performed on a Siemens Biograph Vision. Acquired images were analyzed regarding spatial resolution and image quality (recovery coefficients (RC), coefficient of variation within the background, contrast recovery coefficient (CRC), contrast–noise ratio (CNR), and relative count error in the lung insert). Data were compared between scans with different nuclides.

Results: We found that image quality was comparable between $^{18}$F-FDG and $^{64}$Cu-HCl PET/CT measurements featuring similar maximal endpoint energies of the positrons. In comparison, RC, CRC, and CNR were degraded in $^{68}$Ga-HCl data despite similar count rates. In particular, the two smallest spheres of 10 mm and 13 mm diameter revealed lower RC, CRC, and CNR values. The spatial resolution was similar between $^{18}$F-FDG and $^{64}$Cu-HCl but up to 18% and 23% worse compared with PET/CT images of the NEMA PET body phantom filled with $^{68}$Ga-HCl.

Conclusions: The positron energy of the PET nuclide influences the spatial resolution and image quality of a digital PET/CT scan. The image quality and spatial resolution of $^{68}$Ga-HCl PET/CT images were worse than those of $^{18}$F-FDG or $^{64}$Cu-HCl despite similar count rates.

Keywords: PET/CT, $^{18}$F-FDG, $^{68}$Ga, $^{64}$Cu, NEMA PET body phantom, Jaszczak phantom, Spatial resolution, Image quality
Background

Currently, different radiotracers are available for the examination of the same disease, such as \(^{18}\text{F}-\text{PSMA-1007}\) and \(^{68}\text{Ga}-\text{PSMA-11}\) for the diagnosis of prostate cancer [1–3] or \(^{68}\text{Ga}-\text{DOTA-TOC}\) and \(^{18}\text{F}-\text{SiFAlin-TATE}\) for the diagnosis of neuroendocrine tumors [4]. In a clinical study examining 102 patients, in repeated PET measurements of the same patient using the same PET/CT scanner, \(^{18}\text{F}-\text{PSMA-1007-PET}\) revealed approximately five times as many PSMA-positive lesions attributed to benign origin as \(^{68}\text{Ga}-\text{PSMA-11-PET}\) [1, 2]. The same study revealed that the maximum standardized uptake value \((\text{SUV}_{\text{max}})\) of lesions attributed to benign origin was higher for \(^{18}\text{F}-\text{PSMA-1007-PET}\) than for \(^{68}\text{Ga}-\text{PSMA-11-PET}\) [2]. These differences between PET scans of different radionuclides might be caused by a combination of different effects: Radiotracers differ in their biokinetics and affinities [5]. In addition, state-of-the-art digital PET/CT scanners allow PET measurements with higher sensitivity and spatial resolution. This could reveal differences in PET scans of different nuclides that were previously undetectable. The more precise measurements also offer new possibilities for analyses of the impact of different radionuclides and their physical properties on image quality and detectability of small lesions. The maximum positron energy and thus the mean range in tissue differ between \(^{68}\text{Ga}, \ ^{18}\text{F}\), and \(^{64}\text{Cu}\): Positrons arising from the decay of \(^{68}\text{Ga}\) feature an endpoint energy three times higher than those of \(^{18}\text{F}\) or \(^{64}\text{Cu}\) and therefore have a much greater mean range in tissue \((^{68}\text{Ga}: 3.5 \text{ mm}; \ ^{18}\text{F}: 0.6 \text{ mm}; \ ^{64}\text{Cu}: 0.7 \text{ mm})\) (Table 1) [6]. This degrades the spatial resolution of \(^{68}\text{Ga-PET}\).

The individual impact of different physical properties of PET nuclides on PET image quality and spatial resolution has already been studied in detail, including the use of preclinical PET/CT scanners with a higher intrinsic spatial resolution than clinical PET/CT scanners [7–10]. In addition, different simulations have been performed on this topic [6, 10–12]. However, no analyses have been performed on PET data recorded on PET/CT scanners of the newest generation, which allow for more accurate PET measurements by applying time-of-flight (ToF) measurements and point spread function (PSF) reconstruction. Such devices might reveal new details on the influence of different positron energies of \(^{68}\text{Ga}, \ ^{18}\text{F}\), or \(^{64}\text{Cu}\) on image quality and detectability of small lesions.

Therefore, the aim of this work was to compare the individual impact of the positron energy of the most commonly used PET isotopes \(^{18}\text{F}\) and \(^{68}\text{Ga}\) as well as \(^{64}\text{Cu}\) on the image quality and spatial resolution of scans acquired with one of the latest clinical digital PET/CT scanners. This study focused specifically on the influence of the positron

|                              | \(^{18}\text{F}\) | \(^{68}\text{Ga}\) | \(^{64}\text{Cu}\) |
|------------------------------|-----------------|-----------------|----------------|}
| \(\beta^+\) yield (%)        | 96.7            | 88.0 (1899 keV) | 17.4          |
| \(1.1 (822 \text{ keV})\)   |                 |                 |               |
| Half-life (min)              | 109.8           | 67.6            | 762           |
| Maximum endpoint \(\beta^+\) energy (keV) | 633.5          | 1899.1          | 653.1         |
| Average \(\beta^+\) energy (keV) | 249.3          | 836.0           | 278           |
| Maximum range of positrons in water (mm) | 2.4            | 9.2             | 2.5           |
| Mean range of positrons in water (mm) | 0.6            | 3.5             | 0.7           |

Table 1 Physical properties of the positron-emitting radioisotopes \(^{18}\text{F},\ ^{68}\text{Ga},\) and \(^{64}\text{Cu}\) [16]
energy of the PET nuclide on image quality. All other factors that could influence PET image quality in addition to the positron energy, such as count rate, PET device, phantom setup, and reconstruction parameters, were kept in a comparable range. The scans were recorded with a Biograph Vision PET/CT scanner (Siemens Healthineers) enabling ToF measurements with a timing resolution of 214 ps and an axial spatial resolution of 3.7 mm full width at half maximum (FWHM) for a point source positioned 10 cm from the center of the plane [13]. Due to the very good imaging properties of the scanner, deterioration is to be expected in the $^{68}$Ga images due to the increased positron range.

Methods

A Jaszczak phantom (Model ECT/DLX/P, Data Spectrum Corporation, Durham, USA) was used to qualitatively evaluate spatial resolution and image quality. It is illustrated in Fig. 1. The Jaszczak phantom features cold rods (4.8 mm, 6.4 mm, 7.9 mm, 9.5 mm, 11.1 mm, and 12.7 mm in diameter) and spheres (9.5 mm, 12.7 mm, 15.9 mm, 19.1 mm, 25.4 mm, and 31.8 mm in diameter) surrounded by an activity-filled background compartment (0.3 mol hydrochloric acid (HCl)). PET measurements of the Jaszczak phantom are common for the analysis of spatial resolution and enable a visual assessment of the separability of the cold rods. However, focal hot spots cannot be imitated as in real patient data with the Jaszczak phantom. We additionally determined the spatial resolution semiquantitatively using the NEMA PET body phantom (PTW Freiburg), as shown in Fig. 1. The NEMA PET body phantom allows hot spot imaging of spheres of different sizes (10 mm, 13 mm, 17 mm, 22 mm, 28 mm, and 37 mm in diameter) at different sphere-to-background contrast ratios.

All PET measurements were performed on a digital Biograph Vision PET/CT system (Siemens Healthineers). Following a low-dose CT (120 kVp, 78 mAs, spiral pitch factor
of 1.5, 512 × 512 matrix with a pixel size of 0.98 mm × 0.98 mm) used for attenuation correction of the subsequent PET scan, the PET data were acquired in the list mode over a single bed position, covering an axial field of view (FoV) of 26 cm [13]. The duration of the PET data acquisition of the Jaszczak phantom filled with ¹⁸F-FDG was determined based on the recommendations in the NEMA NU 2-2018 protocol for the characterization of image quality [14]. In the corresponding measurements with the NEMA PET body phantom, the standards in clinical routine according to the German Guideline for ¹⁸F-FDG PET/CT in oncology were used as a reference [15]. The duration of the acquisition of the ⁶⁸Ga-HCl and ⁶⁴Cu-HCl PET data was adjusted to that of the ¹⁸F-FDG-PET data. The respective actual activity concentration in the phantom at the timepoint of imaging and the decay probability of the nuclides (Table 1) were taken into account to achieve similar count statistics between scans with different nuclides. PET data were reconstructed according to the standards in our clinical routine for ¹⁸F-FDG using an ordered subset expectation maximization (OSEM) 3D iterative reconstruction algorithm with 6 iterations and 5 subsets (6i5s), applying PSF and ToF (TrueX algorithm) with an image matrix size of 440 × 440, resulting in a voxel size of (1.65 × 1.65 × 1.5) mm³. No postfiltering was applied (all-pass filter). Reconstructions were performed with attenuation correction and relative scatter correction.

Qualitative evaluation of spatial resolution using the Jaszczak phantom

The background compartment of the Jaszczak phantom was filled with ¹⁸F-FDG, ⁶⁸Ga-HCl, or ⁶⁴Cu-HCl aiming at an activity concentration of 5.3 kBq/mL [14].

The acquisition duration of the Jaszczak phantom filled with ¹⁸F-FDG, ⁶⁸Ga-HCl, or ⁶⁴Cu-HCl was 546 s, 611 s, or 2629 s, respectively. PET/CT scans of the Jaszczak phantom were analyzed visually to determine spatial resolution. In each PET/CT scan, the narrowest rods and spheres that were still distinguishable from one another were determined visually.

Semiquantitative evaluation of spatial resolution and image quality using the NEMA PET body phantom

The NEMA PET body phantom was filled with ¹⁸F-FDG, ⁶⁴Cu-HCl, or ⁶⁸Ga-HCl. According to the recommendations in the NEMA NU 2–2018 protocol, we aimed at an activity concentration of 5.3 kBq/ml in the background compartment and a sphere-to-background activity concentration ratio of approximately 4:1 or 8:1 [14]. The actual activity concentrations at the timepoint of imaging differed slightly from the target and are specified in Table 2. The phantom was also scanned after wrapping it in gel cooling packs 1 cm thick containing propylene glycol to simulate attenuation and scatter conditions comparable with those in an obese patient. The acquisition durations of all PET/CT scans of the NEMA PET body phantom are specified in Table 2.

Image analysis

Spatial resolution was evaluated semiquantitatively according to Hofheinz et al. [17] using the software Rover (version 3.0.60 h, ABX, Radeberg, Germany). Briefly, the resolution was determined based on the analysis of radial activity profiles of the homogeneously filled phantom spheres and the assessment of the FWHM of the PSF in the
reconstructed images. PSF was modeled by a 3D Gaussian function, and FWHM was determined by applying the method described in detail in [17]. This method is based on fitting the analytic solution for the radial activity profile of a homogeneous sphere convolved with a 3D Gaussian function to the reconstructed data. In this process, the full 3D vicinity of each sphere is evaluated by transforming the data to spherical coordinates relative to the center of the sphere. The analytic solution has five parameters: signal (true activity within the sphere), background level, FWHM of the PSF, sphere radius, and wall thickness of the spherical inserts. The radius and wall thickness of the spheres were fixed to their known values. The remaining three parameters were determined by nonlinear least-squares fitting. With this method, the spatial resolution can be determined at a finite background as well as for extended objects. Therefore, it allowed us to study the size and contrast dependence of the resolution. Note that this method assumes a Gaussian PSF, which is never exactly the case. However, the method still leads to a reasonable approximation of the spatial resolution as long as the slope at the object boundary (signal decline) is modeled correctly by the fit function. The means and standard deviations of the FWHM of all six spheres were compared between PET measurements with different nuclides.

Image quality was evaluated semiquantitatively using Rover (version 3.0.60 h, ABX, Radeberg, Germany). Three uniform background volumes of interest (VOIs) of at least 61 ml volume were delimited as illustrated in Additional file 1: Fig. S1A. According to [18], a 3D isocontour at 50% of the maximum pixel value was used for the segmentation of each sphere, taking into account the activity concentration in the background of the phantom. For each sphere, the mean, maximum, and peak recovery coefficients (RCmean, RCmax, and RCpeak, respectively) were determined as the ratio of the measured mean, maximum, or peak SUV of the VOI to the actual activity concentration in the phantom sphere at the timepoint of imaging. SUVpeak was determined as the mean SUV of a
12-mm-diameter spherical region around the maximum voxel, considering only voxels within the 3D isocontour of the sphere [19]. The actual activity concentration was determined as the amount of activity filled in the phantom (measured with an activity meter, which was calibrated for the nuclide and syringe used to fill the phantom with the radioactivity) divided by the volume of the phantom compartment. The means and standard deviations of the RC values of all six spheres were compared.

Image noise, called percent background variability in the NEMA NU 2-2018 protocol [14], was calculated as the coefficient of variation within each of the three background VOIs (CoVBG). The mean CoVBG values of the three VOIs were compared between PET images of different nuclides.

The contrast between each sphere and the background (contrast recovery coefficient (CRC)) was calculated according to the definitions in the NEMA NU 2-2018 protocol [14]. The means and standard deviations of the CRC of all spheres were compared between PET images.

The contrast–noise ratio (CNR) was calculated as the difference in SUVmean between each sphere and the background divided by the standard deviation of the activity concentration in the background compartment. The means and standard deviations of the CNRs of all spheres were compared between PET images.

According to the NEMA NU 2-2018 protocol [14], the relative count error in the lung insert of the NEMA PET body phantom was determined as the ratio of the average number of counts in a cylindrical VOI with a 30 mm diameter in the lung insert of the phantom not filled with radioactivity relative to the average number of counts within the three VOIs placed in the activity-filled phantom background.

Results

Qualitative evaluation of spatial resolution using the Jaszczak phantom

Having adjusted the reconstruction time to the radionuclide-specific half-life, positron yield (Table 1), and activity concentration in the phantom at the timepoint of imaging (Table 2), the total number of true events detected by the PET/CT scanner was similar in the 18F-FDG (86.57 Mcts), 68Ga-HCl (81.65 Mcts), and 64Cu-HCl measurements (85.81 Mcts).

Figure 2 compares the resolution of PET images of the Jaszczak phantom filled with 18F-FDG, 68Ga-HCl, or 64Cu-HCl. In the 18F-FDG and 64Cu-HCl measurements, the resolvable rods were those separated 4.8 mm apart (rods with the smallest distance in the phantom) as well as all of the more widely spaced rods. In comparison, in the 68Ga-HCl measurements, the minimum resolvable rods were those separated 6.4 mm apart (the rods with the second smallest distance in the phantom). The smallest of the six cold spheres featuring a diameter of 9.5 mm was clearly recognizable in PET images of all three nuclides.

Analysis of NEMA PET body phantom measurement

With a comparable weight setup of the phantom, the number of true counts detected by the PET/CT scanner was similar for scans of the NEMA PET body phantom filled with 18F-FDG, 68Ga-HCl, or 64Cu-HCl (Table 2). Wrapping the phantom in cooling packs for simulation of attenuation and scatter conditions similar to those in an obese patient
reduced the number of true events detected. The decrease in the number of true events detected was similar for all nuclides and ranged between 36.1% for $^{68}$Ga-HCl measurements at the low contrast ratio and 38.4% for $^{68}$Ga-HCl measurements at the high contrast ratio (Table 2).

### Semiquantitative evaluation of spatial resolution

When comparing scans of the NEMA PET body phantom at similar sphere-to-background contrast ratios and weight setups, the spatial resolution was similar between $^{18}$F-FDG and $^{64}$Cu-HCl PET measurements but worse in the respective $^{68}$Ga-HCl measurements (Table 3). For each nuclide, spatial resolution was degraded in scans of the phantom mimicking an obese weight setup compared with the respective scan of the phantom not wrapped with cooling packs. The spatial resolution was up to 13% better in the PET image of the phantom filled with a higher versus lower sphere-to-background activity concentration ratio (Table 3: $^{18}$F-FDG measurements and obese setup).

### Semiquantitative evaluation of image quality

Representative of all sphere-to-background activity concentration ratios and weight setups, Fig. 3 compares the $RC_{mean}$, $RC_{max}$, $RC_{peak}$, CRC, and CNR of all spheres of the NEMA PET body phantom filled with $^{18}$F-FDG, $^{68}$Ga-HCl, or $^{64}$Cu-HCl at a sphere-to-background activity concentration ratio of approximately 4:1 and without the simulation of additional attenuation and scattering, as in an obese patient. For all parameters describing image quality, the means and standard deviations of all six spheres are given in Table 4.
The mean activity concentrations recovered from the PET/CT images were comparable with the true activity concentrations for all activity-filled spheres of different diameters in $^{18}$F-FDG measurements of the NEMA PET body phantom, as indicated by $R_{\text{mean}} \approx 1$ for all spheres (Fig. 3). These findings were independent of the contrast ratio...
or weight setup (Table 4). Similarly, the $R_{\text{mean}}$ values were comparable between spheres of different diameters in all $^{64}$Cu-HCl measurements. In comparison, $R_{\text{mean}}$ differed between spheres of different sizes in the $^{68}$Ga-HCl PET/CT scans. Spheres with a diameter of 10 mm showed the largest differences compared with the true activity concentration within the sphere, while recovery improved with increasing sphere size, and $R_{\text{mean}}$ reached $\approx 1$ for the largest spheres (Fig. 3). For example, in $^{68}$Ga-HCl PET/CT scans of the phantom filled with a sphere-to-background activity concentration ratio of approximately 8:1 and with normal weight setup, $R_{\text{mean}}$ differed between 0.87 and 1.02 for the sphere of the smallest and largest diameter.

While the measurements of the NEMA PET body phantom filled with $^{64}$Cu-HCl at low sphere-to-background activity concentration ratios revealed an $R_{\text{mean}}$ of approximately 1 for all spheres, the mean activity concentration recovered from all spheres was on average 10 to 12% higher than the assumed true activity concentration in the $^{64}$Cu-HCl measurements at higher sphere-to-background activity concentration ratios (Table 4).

The mean CRC values of spheres of different diameters were similar in the $^{64}$Cu-HCl and $^{18}$F-FDG measurements but lower in the $^{68}$Ga-HCl measurements (Table 4). For each nuclide, the mean CRC values were comparable between measurements of the phantom with different weight setups and sphere-to-background activity concentration ratios. Figure 3 illustrates that the CRC was independent of the size of the sphere in $^{64}$Cu-HCl and $^{18}$F-FDG measurements but decreased continuously with decreasing sphere diameter in $^{68}$Ga-HCl measurements. In all $^{64}$Cu-HCl and $^{18}$F-FDG measurements, CNR was comparable between spheres of different sizes. In comparison, in $^{68}$Ga-HCl measurements, CNR decreased with decreasing sphere diameter (Fig. 3). For example, in $^{68}$Ga-HCl measurements with a normal weight setup and a high contrast ratio, the CNR of the smallest sphere was 33% lower than that of the largest sphere. $\text{CoV}_B$ was similar for all nuclides when comparing PET images of the NEMA PET body phantom with the same setup (weight setup and sphere-to-background activity concentration ratio). $\text{CoV}_B$ was lower in the normal weight setup than in the specific obese setup.

For measurements with all phantom setups, the relative count error in the lung insert was higher for $^{68}$Ga-HCl measurements than for $^{18}$F-FDG or $^{64}$Cu-HCl measurements (Table 4).

**Discussion**

The main finding of this study is that the image quality of PET scans of a clinical, digital PET/CT scanner (especially spatial resolution, recovery coefficients, and image noise) is strongly affected by the positron energy of the PET isotope used and by attenuation.
Qualitative and semiquantitative evaluation of spatial resolution

Both the qualitative analyses of the Jaszczak phantom and the semiquantitative analyses of the NEMA PET body phantom revealed that spatial resolution was degraded in $^{68}$Ga-HCl versus $^{18}$F-FDG and $^{64}$Cu-HCl PET measurements. Similarly, studies in preclinical [10] and clinical PET scanners [20, 21] have found worse spatial resolution in $^{68}$Ga measurements than in $^{18}$F PET measurements. In contrast to the study by Sanchez-Crespo et al., who used similar activity concentrations and acquisition times for $^{18}$F and $^{68}$Ga PET measurements, in this study the count rates between PET scans with the different nuclides were kept the same. The fact that differences are still visible implies that the higher endpoint energy of positrons originating from the decay of $^{68}$Ga (and an associated increase in positron range) compared with those originating from the decay of $^{18}$F or $^{68}$Ga worsens spatial resolution in clinical, digital PET/CT scans. The image resolution was similar between $^{64}$Cu-HCl and $^{18}$F-FDG PET/CT measurements, as the positrons arising from the decay of those two nuclides feature similar endpoint energies.

Consistent with Rogasch et al. [22], we found that the spatial resolution of PET data that were reconstructed using iterative reconstruction and PSF degrades when the signal-to-background ratios decrease. This is most likely a combined effect of the dependence of the spatial resolution on the optimized number of iterations and subsets during iterative PET reconstruction and a dependency of the magnitude of the Gibbs artifacts during PSF reconstruction on the signal-to-background ratio [22]. Similarly, degraded spatial resolution in scans of the NEMA PET body phantom mimicking an obese weight setup is most likely a result of the iterative reconstruction of the PET data and a nonoptimized number of iterations and subsets. In this study, PET data were reconstructed according to the standards in our clinical routine for $^{18}$F-FDG to best replicate clinical practice. This implies an iterative reconstruction of the PET data and a fixed number of iterations and subsets. An optimization of the reconstruction parameters is beyond the scope of this analysis.

Semiquantitative evaluation of image quality

The similar number of true events detected by the PET/CT scanner in the measurements of the NEMA PET body phantom filled with the different nuclides at the normal or obese weight setup (Table 2) indicates that the adjustment of the analyzed recording time accurately compensated for nuclide-specific differences in positron yield and half-life. There were small deviations between targeted and actual activity concentrations or contrast ratios in the phantom at the timepoint of the PET/CT scans. However, this did not affect the analyses, as the activity concentrations recovered from the PET/CT images were normalized to actual activity concentrations in the phantom and as the statistics were kept comparable between the scans of the phantom at similar weight setups.

In all $^{18}$F-FDG and $^{64}$Cu-HCl measurements, the mean activity concentrations recovered from the PET/CT images were comparable between spheres of different sizes. In comparison, in the $^{68}$Ga-HCl measurements, the $RC_{\text{mean}}$ values were lower for spheres of $\leq 13$ mm diameter but reached $\approx 1$ for spheres $\geq 22$ mm diameter. The lower $RC$
values for the smallest spheres filled with $^{68}$Ga-HCl are most likely a result of the degraded spatial resolution of smaller objects in $^{68}$Ga-PET/CT scans. In agreement with the presented $^{68}$Ga-HCl measurements but differing from the $^{18}$F-FDG measurements, both Sanchez-Crespo et al. [20] and Ryu et al. [21] found that the scanned object should be at least 20 mm in diameter to accurately recover the true mean activity concentration in $^{18}$F-FDG and $^{68}$Ga-HCl images of a clinical PET/CT scanner. The different findings in the $^{18}$F-FDG measurements presented in this study are most likely due to the use of the latest PET/CT scanning technology. In this study, a ToF-enabled PET/CT scanner was used, which has been shown to exhibit less size-based partial volume bias than non-ToF scanners [23].

While the mean activity concentrations were accurately recovered in all $^{18}$F-FDG measurements and in $^{64}$Cu-HCl measurements at low sphere-to-background activity concentration ratios ($\text{RC}_{\text{mean}} \approx 1$), the recovered mean activity concentrations were approximately 10% higher than the true activity concentration in all spheres when the phantom was filled with $^{64}$Cu-HCl at sphere-to-background activity concentration ratios of approximately 8:1. As $\text{RC}_{\text{mean}}$ was increased equally for all spheres of this measurement series, we assume that inaccuracies have occurred in determining the true activity concentration. Measurement inaccuracies can occur when measuring low radioactivity values with the gamma counter. These inaccuracies are larger for $^{64}$Cu-HCl than for $^{18}$F-FDG or $^{68}$Ga-HCl due to the much lower $\beta^+$ yield (Table 1). In addition, the determination of the volume of solution in which the radioactivity was diluted may have been incorrect. This may have resulted in an overestimation of the activity concentration. However, small underestimations of the true activity concentration are not decisive for the core message of this study: the consistency of the image quality parameters for spheres of different diameters.

The PSF reconstruction used in this study leads not only to an improved sharpness of hot spheres and higher spatial resolution [25] but also to an artificial edge overemphasis, which is called the Gibbs artifact [24, 26]. In addition, ToF and the use of an inappropriate kernel can cause edge artifacts and affect contrast recovery and spatial resolution [27]. The artificial increase in the retrieved activity concentration, which is particularly pronounced at the edge of a lesion, can be seen in the radial profiles shown in Fig. 4. The overestimation of the retrieved activity concentration is nonlinear and increases with decreasing lesion size (Fig. 4) [24]. This explains the higher $\text{RC}_{\text{max}}$ values of the smallest sphere of 10 mm diameter compared with larger spheres, as found in our $^{18}$F-FDG and $^{64}$Cu-HCl measurements.

The magnitude of the Gibbs artifacts might have been comparable between the measurements with the different nuclides as the same reconstruction parameters were used. However, partial volume effects were presumably higher in the $^{68}$Ga-HCl measurements than in the $^{18}$F-FDG or $^{64}$Cu-HCl measurements as the mean range of positrons in water originating from the decay of $^{68}$Ga is higher (Table 1). This may have outweighed the nonlinear overestimation of the retrieved activity concentration, which in combination resulted in lower $\text{RC}_{\text{mean}}$ and $\text{RC}_{\text{max}}$ values in the $^{68}$Ga-HCl measurements than in the $^{18}$F-FDG or $^{64}$Cu-HCl measurements, especially for the smallest sphere.
The higher positron range and partial volume effects in PET measurements with \( ^{68}\text{Ga} \) potentially reduce CRC and CNR in \(^{68}\text{Ga-HCl}\) measurements compared with \(^{64}\text{Cu-HCl}\) or \(^{18}\text{F-FDG}\) measurements. This is particularly relevant for small lesions and might explain the decrease in CNR and CRC with decreasing sphere diameter. Similarly, Ryu et al. reported lower CRC values in \(^{68}\text{Ga-PET}\) scans than in \(^{18}\text{F-PET}\) scans for spheres of 10 mm to 37 mm diameter \([21]\). While Ryu et al. reported a decrease in CRC with decreasing sphere size in \(^{68}\text{Ga-HCl}\) and \(^{18}\text{F-FDG}\) measurements, the CRC values in our \(^{18}\text{F-FDG}\) measurements were comparable between all spheres and approximately 100%. The particularly low CRC values for spheres \(\leq 13\) mm in diameter filled with \(^{18}\text{F-FDG}\) in \([21]\) are potentially due to a reconstruction of the data without resolution recovery by PSF or worse ToF performance \([21]\).

The higher relative count error in the lung insert of the NEMA PET body phantom filled with \(^{68}\text{Ga-HCl}\) compared to \(^{18}\text{F-FDG}\) or \(^{64}\text{Cu-HCl}\) was probably caused by a combination of factors. The range of the positrons originating from the decay of \(^{68}\text{Ga}\) is higher, which results in a greater penetration of the lung insert. In addition, single-photon emission and scattered photons arise during the decay of \(^{68}\text{Ga}\) but not during the decay of \(^{18}\text{F}\) or \(^{64}\text{Cu}\).

Wrapping the phantom in cooling packs increased the scattering and absorption of photons and thus reduced the detected number of true counts, which increased \( \text{CoV}_{\text{BG}} \) to a similar extent in PET measurements with all nuclides.
Conclusion

In summary, the image quality was found to be similar between $^{18}$F-FDG and $^{64}$Cu-HCl PET/CT images featuring similar maximal endpoint energies of positions and thus a similar maximum range of positrons in tissue before annihilation. In comparison, the much higher endpoint energy of positrons arising from the decay of $^{68}$Ga degrades quantitative parameters describing image quality (RC, CRC, and CNR) and spatial resolution, especially of small lesions ≤ 13 mm in diameter. As a result, in $^{68}$Ga-HCl images but not in $^{18}$F-FDG and $^{64}$Cu-HCl PET/CT images, quantitative image parameters differ between lesions of different sizes.

Table 4 Means and standard deviations of image quality parameters determined for NEMA PET body phantom measurements

| Parameter | Contrast: ≈ 4:1: normal weight setup | Contrast: ≈ 4:1: obese setup | Contrast: ≈ 8:1: normal weight setup | Contrast: ≈ 8:1: obese setup |
|-----------|-------------------------------------|-------------------------------|-------------------------------------|-------------------------------|
| $^{18}$F-FDG | $1.06 \pm 0.01$ | $1.08 \pm 0.02$ | $1.00 \pm 0.04$ | $1.02 \pm 0.05$
| $^{68}$Ga-HCl | $0.97 \pm 0.09$ | $1.03 \pm 0.08$ | $0.95 \pm 0.06$ | $0.96 \pm 0.05$
| $^{64}$Cu-HCl | $1.05 \pm 0.03$ | $1.07 \pm 0.02$ | $1.10 \pm 0.03$ | $1.12 \pm 0.03$
| $^{18}$F-FDG | $1.38 \pm 0.05$ | $1.45 \pm 0.04$ | $1.33 \pm 0.09$ | $1.37 \pm 0.11$
| $^{68}$Ga-HCl | $1.28 \pm 0.13$ | $1.41 \pm 0.15$ | $1.23 \pm 0.04$ | $1.28 \pm 0.07$
| $^{64}$Cu-HCl | $1.38 \pm 0.06$ | $1.43 \pm 0.05$ | $1.42 \pm 0.09$ | $1.51 \pm 0.06$
| $^{18}$F-FDG | $1.12 \pm 0.23$ | $1.10 \pm 0.25$ | $1.00 \pm 0.33$ | $1.03 \pm 0.34$
| $^{68}$Ga-HCl | $1.01 \pm 0.13$ | $1.08 \pm 0.16$ | $1.00 \pm 0.16$ | $0.99 \pm 0.21$
| $^{64}$Cu-HCl | $1.10 \pm 0.22$ | $1.10 \pm 0.27$ | $1.09 \pm 0.33$ | $1.13 \pm 0.34$
| CRC (%) | $^{18}$F-FDG | $100.99 \pm 1.72$ | $101.57 \pm 2.99$ | $96.46 \pm 4.07$ | $97.86 \pm 5.24$
| $^{68}$Ga-HCl | $81.99 \pm 10.88$ | $76.48 \pm 8.32$ | $80.98 \pm 5.75$ | $91.20 \pm 5.06$
| $^{64}$Cu-HCl | $98.97 \pm 3.67$ | $98.79 \pm 2.18$ | $103.90 \pm 3.32$ | $106.09 \pm 3.09$
| CNR | $^{18}$F-FDG | $34.26 \pm 0.58$ | $24.60 \pm 0.72$ | $86.55 \pm 3.65$ | $64.19 \pm 3.44$
| $^{68}$Ga-HCl | $27.83 \pm 3.69$ | $19.36 \pm 2.11$ | $70.48 \pm 5.00$ | $61.10 \pm 3.39$
| $^{64}$Cu-HCl | $46.86 \pm 1.74$ | $33.24 \pm 0.73$ | $76.40 \pm 2.44$ | $57.87 \pm 1.69$
| CoVBG (%) | $^{18}$F-FDG | $8.96 \pm 0.33$ | $12.55 \pm 0.15$ | $9.72 \pm 0.21$ | $13.29 \pm 0.77$
| $^{68}$Ga-HCl | $9.60 \pm 0.40$ | $12.87 \pm 0.41$ | $9.67 \pm 0.43$ | $12.56 \pm 0.81$
| $^{64}$Cu-HCl | $9.30 \pm 0.64$ | $13.09 \pm 0.63$ | $9.68 \pm 0.69$ | $13.04 \pm 0.77$
| Lung count error [%] | $^{18}$F-FDG | $3.82$ | $5.42$ | $4.19$ | $5.22$
| $^{68}$Ga-HCl | $7.85$ | $13.48$ | $8.23$ | $5.90$
| $^{64}$Cu-HCl | $3.72$ | $5.11$ | $3.76$ | $4.95$

Means and standard deviations of the mean, maximum and peak recovery coefficients ($R_{C_{mean}}$, $R_{C_{max}}$, and $R_{C_{peak}}$ respectively), percent contrast between each sphere and the background (contrast recovery coefficient (CRC)), contrast–noise ratio (CNR), percent background variability (CoVBG) and the relative count error in the lung insert (lung count error). Except for the relative count error in the lung insert, each parameter is presented as the mean ± standard deviation of the six spheres for PET/CT measurements of the NEMA PET body phantom filled with $^{18}$F-FDG, $^{68}$Ga-HCl, or $^{64}$Cu-HCl with sphere-to-background activity concentration ratios of approximately 4:1 or 8:1 and with (obese setup) or without (normal weight setup) the simulation of an obese patient by applying cooling packs around the phantom.
Abbreviations
SUV Standardized uptake value
ToF Time of flight
PSF Point spread function
HCl Hydrochloric acid
FoV Field of view
OSEM Ordered subset expectation maximization
FWHM Full width at half maximum
VOI Volume of interest
RC Recovery coefficient
CoVBG Coefficient of variation within the background VOI
CRC Contrast recovery coefficient
CNR Contrast–noise ratio

Supplementary Information
The online version contains supplementary material available at https://doi.org/10.1186/s40658-022-00487-7.

Additional file 1: Transversal (left), coronal (middle), and sagittal plane of the NEMA PET body phantom filled with 18F-FDG and illustration of the regions of interest.

Acknowledgements
Not applicable.

Author contributions
A.B., R.F., L.O., J.K., and S.H. contributed to the design of the study and discussed the data. A.B. planned, performed, and analyzed the measurements. S.H. is the senior author. All authors discussed the results and implications and commented on the manuscript. All authors read and approved the final manuscript.

Funding
Open Access funding enabled and organized by Projekt DEAL.

Availability of data and material
The datasets used and/or analyzed during the current study are available from the corresponding author on reasonable request.

Declarations
Ethics approval and consent to participate
Not applicable.

Consent for publication
Not applicable.

Competing interests
The authors declare that they have no competing interests.

Received: 4 January 2022 Accepted: 22 August 2022

References
1. Kroenke M, Mirzoyan L, Horn T, Peeken JC, Wurzer A, Wester HJ, et al. Matched-pair comparison of (68)Ga-PSMA-11 and (18)F-rhPSMA-7 PET/CT in patients with primary and biochemical recurrence of prostate cancer: frequency of non-tumor-related uptake and tumor positivity. J Nucl Med. 2021;62(8):1082–8.
2. Rauscher I, Kronke M, Konig M, Gafita A, Maurer T, Horn T, et al. Matched-pair comparison of (68)Ga-PSMA-11 PET/CT and (18)F-PSMA-1007 PET/CT: frequency of pitfalls and detection efficacy in biochemical recurrence after radical prostatectomy. J Nucl Med. 2020;61(1):51–7.
3. Hoberuck S, Lock S, Borkowitz A, Sommer U, Winzer R, Zophel K, et al. Intraindividual comparison of [(68) Ga]-Ga-PSMA-11 and [(18)F]-F-PSMA-1007 in prostate cancer patients: a retrospective single-center analysis. EJNMMI Res. 2021;11(1):109.
4. Ilhan H, Lindner S, Todicca A, Cyran CC, Tiling R, Auernhammer CJ, et al. Biodistribution and first clinical results of (18)F-SiFAlin-TATE PET: a novel (18)F-labeled somatostatin analog for imaging of neuroendocrine tumors. Eur J Nucl Med Mol Imaging. 2020;47(4):870–80.
5. Kotzerke J, van den Hoff J. Quantification: there is more to worry about than good scanner hardware and reliable calibration. Eur J Nucl Med Mol Imaging. 2017;44(12):1955–7.
6. Carbine P, Vandendriessche S, Diogo A, Perez-Benito D, Efthimiou N, Thyssen C, et al. Monte Carlo Simulations of the GE Signa PET/MR for Different Radioisotopes. Front Physiol. 2020;11: 525575.
7. Disselhorst JA, Brom M, Laverman P, Slump CH, Boerman OC, Oyen WJ, et al. Image-quality assessment for several positron emitters using the NEMA NU 4–2008 standards in the Siemens Inveon small-animal PET scanner. J Nucl Med. 2010;51(4):610–7.
8. Teuho J, Riehakainen L, Honkaniemi A, Moisio O, Han C, Terri M, et al. Evaluation of image quality with four positron emitters and three preclinical PET/CT systems. EJNMMI Res. 2020;10(1):155.
9. Cal-Gonzalez J, Vaequero JJ, Herrera JL, Perez-Liva M, Soto-Montenegro ML, Pena-Zalbidea S, et al. Improving PET quantification of small animal [(68)Ga]DOTA-labeled PET/CT studies by using a CT-based positron range correction. Mol Imaging Biol. 2018;20(4):584–93.
10. Carter LM, Kesner AL, Pratt EC, Sanders VA, Massicano AVF, Cutler CS, et al. The impact of positron range on PET resolution, evaluated with phantoms and PHITS Monte Carlo simulations for conventional and non-conventional radionuclides. Mol Imaging Biol. 2020;22(1):73–84.
11. Bertollì O, Eleftheriou A, Cecchetti M, Camarlinghi N, Belcari N, Tsoumpas C. PET iterative reconstruction incorporating an efficient positron range correction method. Phys Med. 2016;32(2):323–30.
12. Emond EC, Groves AM, Hutton BF, Thielemans K. Effect of positron range on PET quantification in diseased and normal lungs. Phys Med Biol. 2019;64(20):205010.
13. Specifications Sheet: Biograph Vision 600 System Specifications. 01.2019. Contract No.: MI-4235.KBK.JV.
14. NEMA Standards Publication NU 2–2018 Performance Measurements of Positron Emission Tomographs (PETS). National Electrical Manufacturers Association; 2018.
15. Krause BJ, Beyer T, Bockisch A, Delbeke D, Kotzerke J, Minkov V, et al. FDG-PET/CT in oncology. German guideline. Nuklearmedizin. 2007;46(6):291–301.
16. Conti M, Eriksson L. Physics of pure and non-pure positron emitters for PET: a review and a discussion. EJNMMI Phys. 2016;3(1):8.
17. Hofheinz F, Dittrich S, Potzsch C, Hoff J. Effects of cold sphere walls in PET phantom measurements on the volume reproducing threshold. Phys Med Biol. 2010;55(4):1099–113.
18. Boellaard R, O’Doherty MJ, Weber WA, Mottaghy FM, Lonsdale MN, Stoobants SG, et al. FDG PET and PET/CT: EANM procedure guidelines for tumour PET imaging: version 1.0. Eur J Nucl Med Mol Imaging. 2010;37(1):181–200.
19. Lodge MA, Chaudhry MA, Wahl RL. Noise considerations for PET quantification using maximum and peak standardized uptake value. J Nucl Med. 2012;53(7):1041–7.
20. Sanchez-Crespo A. Comparison of Gallium-68 and Fluorine-18 imaging characteristics in positron emission tomography. Appl Radiat Isot. 2013;76:55–62.
21. Ryu H, Meikle SR, Willowson KP, Eilick EM, Bailey DL. Performance evaluation of quantitative SPECT/CT using NEMA NU 2 PET methodology. Phys Med Biol. 2019;64(14):145017.
22. Rogasch JM, Hofheinz F, Lougovski A, Furth C, Ruf J, Grosser OS, et al. The influence of different signal-to-background ratios on spatial resolution and F18-FDG-PET quantification using point spread function and time-of-flight reconstruction. EJNMMI Phys. 2014;1(1):12.
23. Sunderland JJ, Christian PE. Quantitative PET/CT scanner performance characterization based upon the society of nuclear medicine and molecular imaging clinical trials network oncology clinical simulator phantom. J Nucl Med. 2015;56(1):145–52.
24. Kaalep A, Huisman M, Sera T, Vugs D, Boellaard R, et al. Feasibility of PET/CT system performance harmonisation for quantitative multicentre (89)Zr studies. EJNMMI Phys. 2018;5(1):16.
25. Nuyts J. Unconstrained image reconstruction with resolution modelling does not have a unique solution. EJNMMI Phys. 2014;1(1):98.
26. Lason C, Desmonts C, Quaek E, Gervais R, Do P, Dubos-Argis C, et al. Harmonizing SUVs in multicentre trials when using different generation PET systems: prospective validation in non-small cell lung cancer patients. Eur J Nucl Med Mol Imaging. 2013;40(7):985–96.
27. Efthimiou N, Thielemans K, Emond E, Cawthorne C, Archibald SJ, Tsoumpas C. Use of non-Gaussian time-of-flight kernels for image reconstruction of Monte Carlo simulated data of ultra-fast PET scanners. EJNMMI Phys. 2020;7(1):42.

Publisher’s Note
Springer Nature remains neutral with regard to jurisdictional claims in published maps and institutional affiliations.