Feasibility of equivalent performance of 3D TOF $[^{18}\text{F}]$-FDG PET/CT with reduced acquisition time using clinical and semiquantitative parameters

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Abstract

Background: High-performance time-of-flight (TOF) positron emission tomography (PET) systems have the capability for rapid data acquisition while preserving diagnostic image quality. However, determining a reliable and clinically applicable cut-off of the acquisition time plays an important role in routine practice. This study aimed to assess the diagnostic equivalence of short acquisition time of 57 with routine 75 seconds per bed position (s/BP) of $[^{18}\text{F}]$-fluoro-deoxy-glucose (FDG) PET.

Phantom studies applying EARL criteria suggested the feasibility of shortened acquisition time in routine clinical imaging by 3D TOF PET/CT scanners. Ninety-six patients with melanoma, lung or head and neck cancer underwent a standard whole-body, skull base-to-thigh or vertex-to-thigh $[^{18}\text{F}]$-FDG PET/CT examination using the 3D TOF Ingenuity TF PET/CT system (Philips, Cleveland, OH). The $[^{18}\text{F}]$-FDG activity applied was equal to 4MBq per kg body weight. Retrospectively, PET list-mode data were used to calculate a second PET study per patient with a reduced acquisition time of 57 s instead of routine 75 s/BP. PET/CT data were reconstructed using a 3D OSEM TOF algorithm. Blinded patient data were analysed by two nuclear medicine physicians. The number of $[^{18}\text{F}]$-FDG-avid lesions per body region (head&neck, thorax, abdomen, bone, extremity) and image quality (grade 1–5) were evaluated. Semiquantitative analyses were performed by standardized uptake value (SUV) measurements using 3D volume of interests (VOI).

The visual and semiquantitative diagnostic equivalence of 214 $[^{18}\text{F}]$-FDG-avid lesions were analysed in the routine standard (75 s/BP) as well as the calculated PET/CT studies with short acquisition time. Statistical analyses were performed by equivalence testing and Bland–Altman plots.

Results: Lesion detection rate per patient's body region agreed in > 98% comparing 57 s/BP and 75 s/BP datasets. Overall image quality was determined as equal or superior to 75 s in 80% and 69%, respectively. In the semiquantitative lesion-based analyses, a significant equivalence was found between the 75 s/BP and 57 s/BP PET/CT images both for $\text{SUV}_{\text{max}}$ ($p = 0.004$) and $\text{SUV}_{\text{mean}}$ ($p = 0.003$).

Conclusion: The results of this study demonstrate significant clinical and semiquantitative equivalence between short acquisition time of 57 s/BP and standard 75 s/BP 3D TOF $[^{18}\text{F}]$-FDG PET/CT scanning, which may improve the patient's workflow in routine practice.

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Introduction

Positron emission tomography/computed tomography (PET/CT) has been widely implemented as a diagnostic tool in the field of oncology, cardiology and neurology in clinical routine [1–4]. The most commonly employed PET radiopharmaceutical is $[^{18}\text{F}]$-fluoro-deoxy-glucose (FDG), a glucose analogue [2, 5, 6].

In 2015, the European Association of Nuclear Medicine (EANM) published the current guidelines for the calculation of $[^{18}\text{F}]$-FDG activity to be applied based on the patient’s body weight (bw), scanner type and PET acquisition time [2]. However, current EANM guidelines defining PET acquisition time and $[^{18}\text{F}]$-FDG activity calculation may not exactly reflect the ongoing technical improvements in PET/CT imaging [3, 7–10], which allow reductions in PET acquisition time or $[^{18}\text{F}]$-FDG activity applied while keeping high image quality [7, 11–13]. One of the major technical advancements in PET/CT scanners was the implementation of the time-of-flight approach leading to better performance of this modality and more accurate localization of the annihilation process [14]. Hence, image reconstruction and image quality could be substantially improved [15].

These advances could enable further reduction of $[^{18}\text{F}]$-FDG activity applied, which is highly desirable in clinical routine keeping the radiation exposure for patients and hospital staff as low as reasonably achievable (ALARA principle) [12, 16].

In addition, the possibility to reduce the PET acquisition time per bed position will improve the workflow of the PET/CT centres as well as the patient comfort [15]. Based on our previous phantom study results, the shortest EARL approved and equivalent time of acquisition was equal to 57 s per bed position [17]. In this study, we have tried to assess the clinical performance of the short 57 s/BP PET acquisition time with validated standard PET acquisition time of 75 s/BP. Furthermore, semiquantitative data were compared between short and standard PET acquisitions using “equivalence testing”, as a novel approach.

Material and methods

Patient population

The $[^{18}\text{F}]$-FDG PET/CT data of 96 consecutive patients with histopathologically verified melanoma, lung or head and neck cancer were retrospectively analysed. These three cancers types were selected for the following reasons: Firstly, those are highly referred cancer patients for PET/CT imaging in our centre, secondly, $[^{18}\text{F}]$-FDG-PET/CT studies are routinely integrated into the work-up of these cancer types and thirdly, there exists an integrated common and standardized histopathologic work-up of $[^{18}\text{F}]$-FDG-avid lesions. Patients were divided into three different weight classes (< 75 kg, 75–100 kg, > 100 kg). Patient demographics are shown in Table 1. Exclusion criteria were pregnancy, age < 18 years and a glucose level > 195 mg/dl measured before tracer application. This research was conducted according to the principles of the Declaration of Helsinki and all subsequent revisions and was approved by the routing Ethics Committee of the province (EC-number: 415-E/2491/2-2019). All data were carefully anonymized to fulfil the regulations regarding data protection.

PET/CT examination and data reconstruction

All 6-h fasting cancer patients underwent a standard whole-body (melanoma), vertex-to-thigh (head and neck) or skull base-to-thigh (lung) $[^{18}\text{F}]$-FDG PET/CT examination using our EARL accredited 3D TOF Ingenuity TF PET/CT system (Philips, Cleveland, OH) including a low-dose CT with 100 kV and 45 mAs for anatomical localization and attenuation correction purposes. Each patient received 4 MBq per kg bw (range 246–479 MBq; mean ± SD 345 ± 71 MBq) of $[^{18}\text{F}]$-FDG. The mean uptake time was 60 min ± 12 min as recommended by the European Association of Nuclear Medicine Research Ltd (EARL) [2]. All patients had to empty their bladder before PET/CT scanning. Following low-dose CT scan, PET acquisition was performed with a duration of 75 s per bed position and a bed overlap of 50% over the same anatomical area. PET data were reconstructed using the

| Patient # | Disease                   | Male | Female | Mean age (y) ± SD | < 75 kg bw | 75–100 kg bw | > 100 kg bw | Mean applied MBq ± SD |
|-----------|---------------------------|------|--------|------------------|------------|--------------|-------------|----------------------|
| 33        | Melanoma                  | 18   | 15     | 65.9 ± 13.5      | 11         | 12           | 10          | 341.6 ± 69.3         |
| 34        | Lung cancer               | 21   | 13     | 65.8 ± 8.3       | 10         | 10           | 14          | 351.9 ± 73.8         |
| 29        | Head & neck cancer        | 21   | 8      | 63.3 ± 12.6      | 10         | 11           | 8           | 342.0 ± 69.5         |
vendor-recommended blob-based ordered-subset expectation maximization (OSEM) time of flight (TOF) algorithm [18] with the default setting of 3 iterations and 33 subsets and a matrix of $144 \times 144$ with a voxel size of $4 \times 4 \times 4$ mm$^3$.

Transverse and axial spatial resolution of our PET scanner was equal to 4.7 mm, the TOF system sensitivity in the centre was $> 18830$ cps/MBq and the timing resolution of TOF performance was equal to 495 ps.

No post-reconstruction smoothing filter was used. All image data received correction for random coincidences, normalization, dead time losses, scatter and attenuation as recommended by the EANM guidelines version 1.0 [5].

Additionally, a second imaging dataset per patient with a short acquisition time of 57 s per bed position was reconstructed using PET list-mode data. All image reconstruction settings were identical.

**Imaging analysis**

The data of 192 anonymized PET/CT studies, two datasets per patient (75 s and 57 s PET acquisition time), were analysed using the Intellispace software version 10.1 (Philips Medical Systems, The Netherlands) by two experienced board-certified nuclear medicine physicians blinded to any information concerning patient, indication, PET acquisition time or applied radiotracer activity. [$^{18}$F]-FDG-avid lesions per body region (head & neck, thorax, abdomen, bone, extremity) and image quality (grade 1-5), further demonstrated in Table 2, were evaluated. Criteria for image quality receiving grade $\geq 3$ were image reconstruction artefacts, noisy images, irregular shape or contour of lesions and a low lesion to background ratio. Reader 1 and Reader 2 categorized the [$^{18}$F]-FDG-avid lesions per body regions in three groups, 0, 1–4 and $> 4$ lesions in order to generate reports for all 192 PET/CT datasets. After the completion of the reading, the report results with the categorized lesions were compared with the standard and recalculated PET/CT studies with reduced acquisition time.

SUV$_{\text{max}}$ and SUV$_{\text{mean}}$ were calculated with the help of 3D VOI for 214 [$^{18}$F]-FDG-avid lesions matched per patient between the 75 s and 57 s PET/CT studies. The matched lesions were mainly defined based on their anatomical localization and their morphological pattern on CT. For each matched lesion, the SUV was determined using the same threshold- and volume-based VOI determination in order to compare the SUV as precisely as possible. Firstly, we defined a 40% threshold for drawing an automatic VOI over the abnormal matched lesions. Secondly, we tried to adjust this threshold manually to receive the same volume on both matched lesions and therefore avoid any possible influence of different VOIs on the results.

As a measure for image noise, SUV$_{\text{mean}}$ plus standard deviation were additionally measured by drawing a circular region of interest (ROI) in normal liver tissue and the mediastinum in order to calculate the coefficient of variation (COV). The diameter of the ROIs was equal to $7.4 \pm 0.9$ cm (liver) and $4.3 \pm 0.8$ cm (mediastinum). The COV, which is defined as the standard deviation divided by the SUV$_{\text{mean}}$ [19], was evaluated for liver and mediastinum region for each PET/CT dataset.

**Statistical analysis**

A comparison of the mean of differences of SUV$_{\text{max}}$ and SUV$_{\text{mean}}$ of the 75 s and 57 s PET/CT datasets was performed by equivalence testing (TOST) [20] using the R software version 3.6.3 [21]. Due to the lack of objective and clinically sensitive equivalence margins on the original scale of SUV$_{\text{max}}$ and SUV$_{\text{mean}}$, standardized equivalence margins (i.e. Cohen’s d) equal to 0.2 were used. This means that only very small SUV differences between 75 s and 57 s PET/CT studies would be tolerated. For all equivalence tests, the alpha level was set to 0.05.

Descriptive statistics of continuous data were represented by the calculation of the mean $\pm$ SD, minimum and maximum. Frequencies in percent were evaluated for categorical variables, e.g. image quality.

Bland–Altman plots, the method of choice for interpreting comparison studies [22], were created with GraphPad Prism 6 for assessing the agreement of SUV$_{\text{max}}$ and SUV$_{\text{mean}}$ between the 75 s and 57 s PET/CT datasets.

We would like to mention that from a subject-matter point of view, it might seem counterintuitive to use equivalence tests instead of non-inferiority tests. However, we prefer the former approach, because it aligns well with the two-sided concept of the visual inspections (i.e. the Bland–Altman plots). From a statistical point of view, both tests are equivalent in our setting anyway, because the superiority case is highly unlikely if not impossible at all, and therefore, the $p$ values of the TOST and the non-inferiority tests will be the same.

| Table 2 | Description of different grades for the assessment of image quality |
|---------|---------------------------------------------------------------|
| Grade 1 | Excellent |
| Grade 2 | Good |
| Grade 3 | Average |
| Grade 4 | Poor but interpretable |
| Grade 5 | Poor not interpretable |
Results
The imaging data of 96 oncological patients (60 men, 36 women, age 27–87, mean ± SD 65 ± 11.6 years) were analysed in this retrospective study.

Visual analyses
Equal report results were found in > 98% of the 480 investigated body regions (5 body regions per patient: head & neck, thorax, abdomen, bone, extremity) for both readers comparing 75 and 57 s PET/CT datasets. Of 480 analysed patient’s body regions, 9 (1.88%) were categorized differently concerning the number of [18F]-FDG-positive lesions by reader 1, whereas reader 2 investigated 5 (1.04%) differently (see Table 3). Table 4 summarizes, for both investigators, the results of lesion classification per patient’s body region. An additional visual analyses of the images showed that all matched PET/CT datasets, 75 s and the corresponding 57 s per patient, showed the same number of [18F]-FDG-positive lesions.

The image quality of short acquisition was assessed as equal in 62.5% and superior in 17.7% by reader 1. Moreover, reader 2 rated the image quality of short acquisition as even in 42.7% and superior in 26%. Therefore, the overall subjective image quality was described as equal or superior in 80.2 and 68.7%, respectively. It is noteworthy that no PET/CT dataset with short acquisition time was graded as not interpretable. All image quality grades of both readers are provided in Table 5.

Figures 1, 2 and 3 show examples of PET/CT images of patients suffering from melanoma, lung or head and neck cancer with < 75 kg, 75–100 kg and > 100 kg.

Semiquantitative analyses
SUV_{max} and SUV_{mean} were calculated for 214 [18F]-FDG-avid lesions (164 malignant: mean SUV_{max} 7.9 ± 4.0, mean SUV_{mean} 4.8 ± 1.9, 50 benign: mean SUV_{max} 6.0 ± 3.3, mean SUV_{mean} 4 ± 1.4) matched per patient between the 75 s and 57 s [18F]-FDG PET/CT studies. Mean difference of SUV_{max} ± SD was equal to 0.0074 ± 0.49, and the 90% confidence interval (CI) ranged from −0.048 to −0.062. Moreover, the mean difference of SUV_{mean} ± SD was equal to 0.0015 ± 0.14. The 90% CI ranged from −0.015 to −0.017. Since the CIs lay within the equivalence bounds, the equivalence tests were significant. Accordingly, the obtained p values were 0.004 for SUV_{max} and 0.003 for SUV_{mean}. An illustrative visualization for the equivalence tests is visualized in Fig. 4.

Bland–Altman plots show good agreement between the SUVs of the 75 s and 57 s [18F]-FDG PET/CT datasets (see Figs. 5 and 6). Furthermore, no systematic bias was observed in the quantification of SUV_{max} and SUV_{mean}.

Image noise
The overall COV_{liver} was equal to 0.12 ± 0.02 for 75 s PET/CT datasets and 0.13 ± 0.02 for the 57 s PET/CT datasets, while the COV_{mediastinum} resulted in 0.12 ± 0.02 for the 75 s PET/CT datasets and 0.14 ± 0.03 for the 57 s PET/CT datasets. As expected, the COV was increasing for patients with high body weight. Details of the calculated COVs for patient groups with different body weights are demonstrated in Fig. 7.

| Table 3 | Blinded rating of 96 PET datasets per acquisition time |
|---------|-----------------------------------------------------|
|         | Total number of investigated body regions rated equally | Investigated body regions rated differently |
| Reader 1 | 480 | 471 | 9 |
| Reader 2 | 480 | 475 | 5 |

Each patient’s investigation is divided into 5 different body regions leading to 480 body regions to investigate per acquisition time.

| Table 4 | Analysis of 480 body-region classifications: reader 1 and reader 2 |
|---------|-----------------------------------------------------|
|         | 0 lesions | 1–4 lesions | > 4 lesions |
| Reader 1 |          |            |            |
| Head & Neck | 69 | 24 | 3 |
| Thorax | 62 | 21 | 13 |
| Abdomen | 86 | 6 | 4 |
| Bone | 85 | 7 | 4 |
| Extremity | 92 | 3 | 1 |
| Reader 2 |          |            |            |
| Head & Neck | 76 | 18 | 2 |
| Thorax | 64 | 20 | 12 |
| Abdomen | 87 | 6 | 3 |
| Bone | 86 | 8 | 2 |
| Extremity | 94 | 1 | 1 |

| Table 5 | Two crosstabs of image quality: grading of the 75 s and 57 s PET dataset - reader 1 and reader 2 |
|---------|-----------------------------------------------------|
|         | Grade 1 (75 s) | Grade 2 (75 s) | Grade 3 (75 s) | Grade 4 (75 s) |
| Reader 1 |          |            |            |            |
| Grade 1 (57 s) | 29 | 13 | 0 | 0 |
| Grade 2 (57 s) | 15 | 29 | 4 | 0 |
| Grade 3 (57 s) | 0 | 4 | 2 | 0 |
| Grade 4 (57 s) | 0 | 0 | 0 | 0 |
| Reader 2 |          |            |            |            |
| Grade 1 (57 s) | 17 | 22 | 1 | 0 |
| Grade 2 (57 s) | 26 | 22 | 1 | 1 |
| Grade 3 (57 s) | 0 | 3 | 2 | 0 |
| Grade 4 (57 s) | 0 | 0 | 1 | 0 |
Discussion

Since experimental [19, 23] and clinical [23–25] evidence suggested the feasibility to shorten PET acquisition times beyond EANM recommendations, we explored the effect of PET scan time reduction on quantitative segmentation parameters, lesion detectability and image quality using the Ingenuity TF PET/CT system.

Our retrospective patient study, including 96 patients with very common cancers, e.g. melanoma, lung or head and neck cancer, demonstrates diagnostic equivalence of PET imaging datasets with 75 s and 57 s acquisition time. In contrast to other studies dealing with PET scan time reduction, this paper is novel since the comparison of quantitative segmentation parameters between the standard and reduced PET acquisition times are validated by equivalence testing. Our findings seem very plausible due to improvements in PET technology and image reconstruction, e.g. TOF [7], implemented in PET/CT scanners used in clinical routine. TOF technology allows a more accurate localization of the annihilation process leading to increased contrast and reduced noise levels of PET/CT studies [7]. These advantages allow shorter PET acquisition times [7], e.g. 57 s per bed position, while still maintaining good image quality and diagnostic accuracy. Another strength of our study is the inclusion of patients over a wide range of body weight (40–145 kg) demonstrating the diagnostic equivalence of the 57 s and 75 s acquisitions throughout much of the range of clinically presenting body weight.

Previous studies aiming to optimize PET acquisition time or [18F]-FDG activity applied often focused on the calculation of noise equivalent count rate (NECR) [24, 26–29], which is a measure for image signal-to-noise ratio of PET/CT scanners. However, clinical experience suggests that quantitative measurements (e.g. SUV) of imaging performance are more appropriate guides for the assessment of lesion detectability and PET image quality [30]. Comparable studies [15, 23, 25] (see Table 6) either included less PET patient data, evaluated different PET/CT scanners, used other methods for statistical analyses
or demonstrated a higher cut-off for PET acquisition time.

A similar study from Halpern et al. stated that a PET acquisition time of 60 s per bed position is only sufficient for patients with a body weight < than 59kg, although they injected 7.7 MBq/kg/bw [25]. However, the PET/CT scanner described in this paper, which has already been published in 2004, did not have TOF technology. Our study specifically included 32 patients weighing more than 100 kg, and the activity applied was equal to 4 MBq/kg/bw. Figure 7 shows that there is no significant COV difference for PET datasets of patients with a bw > 100 kg between the 75 s and 57 s acquisition times.

On the other hand, Murray et al. demonstrated that even with 15-s acquisition time per bed position \(^{18}\text{F}\)-FDG-avid lesions larger than 2 cm were visible and 2D calculated SUV values were comparable. However, this study compared non-TOF with TOF image reconstruction, and for the statistical analysis no equivalence test was performed [23]. The SUV calculation of our 214 \(^{18}\text{F}\)-FDG-avid lesions is based on 3D volumetric data (range 2.1–253312 mm\(^3\) = 0.002 ml–253.312 ml) used also for equivalence testing. We would like to emphasize that equivalence testing is considered as the state of the art and statistically reliable method for demonstrating equality of two investigated methods [20]. Furthermore, our lesion detection rate per patient’s body region matched in > 98% for both investigators comparing 75 and 57 s PET/CT datasets. All \(^{18}\text{F}\)-FDG-positive lesions were visible in the 75 s as well as the corresponding 57 s PET/CT dataset, and the image quality of short acquisition datasets was rated equal or superior in 80 and 69%.

Another study with the aim to evaluate the scanner performance of a digital PET/CT system concluded a
minimal PET acquisition time of 90s/BP [15]. However, this newest generation scanner shows higher sensitivity, better spatial resolution and highly improved TOF resolution compared to our Ingenuity TF PET/CT system. It is important to mention though that the mean $[^{18}\text{F}]}$-FDG uptake time was equal to 101 min. This increased delay was caused by a performance of an analog PET/CT examination beforehand.[15]

The results of this study are promising when applying a short PET acquisition time of 57 s/BP with comparable diagnostic accuracy and adequate image quality to standard acquisition time. This is highly beneficial since a reduced PET scan time improves patient comfort, which is especially important for many elder or anguished patients, who are more likely to move during the imaging procedure [25]. Accelerated PET imaging can avoid motion artefacts in the latter patient population and increase patient throughput, which meets the ever-increasing demand for PET studies [1] due to an increasing number of well-evidenced clinical indications.

Further benefits can be experienced in the paediatric field where PET/CT studies with short acquisition times might lead to a decrease in anaesthesia or sedation time [15, 31].

Fig. 3 MIP, fused axial and coronal PET/CT images (from left to right) with 75 s (upper row) and 57 s (lower row) PET acquisition time per bed position of a 57-year-old male patient with 105 kg bw suffering from head & neck cancer—no significant difference in image quality, lesion detectability and quantification.
Certain limitations of our study should be taken into account. As already mentioned, the PET acquisition time reduction was investigated by a retrospectively designed patient study. PET/CT datasets with 57 s acquisition time per bed position were calculated using PET-listmode data. Further prospective investigations are necessary for defining the lowest cut-off of PET acquisition time, especially when reducing tracer activity. Furthermore, this study protocol was planned for the Ingenuity TF PET/CT scanner. Although we believe that the described methodology of this study can be easily translated to other 3D TOF PET/CT scanner, these results may not be generalized for all PET/CT scanners from different vendors.

Since the focus of this study was to assess the impact of the PET results on routine clinical practice, reader 1 and reader 2 were asked to perform a global disease assessment (0, 1-4 or > 4 lesions per body region), and therefore, no counting of exact lesion numbers was required for their blind readings.

Fig. 4 Comparison of $SUV_{\text{max}}$ 75 s and 57 s PET/CT datasets and $SUV_{\text{mean}}$ 75 s and 57 s PET/CT images of 214 $[^{18}\text{F}]$-FDG-avid lesions by performing the TOST equivalence test [20].
Conclusion

This retrospective study of clinical PET/CT data successfully demonstrates the diagnostic equivalence of [18F]-FDG PET/CT studies using short acquisition time of 57 s per bed position showing equivalent lesion...
detectability and adequate image quality comparing to standard PET acquisition time.

Abbreviations
ALARA: As low as reasonably achievable; Bw: Body weight; CI: Confidence interval; COV: Coefficient of variation; EANM: European Association of Nuclear Medicine; EARL: EANM Research Ltd; [18F]-FDG: Fluor-18 fluoro-deoxy-glucose; NECR: Noise equivalent count rate; OSEM: Ordered-subset expectation maximization; PET/CT: Positron emission tomography/computed tomography; PSF: Point spread function; ROI: Region of interest; s/BP: Seconds per bed position; SD: Standard deviation; SUV: Standardized uptake value; TOF: Time-of-flight; VOI: Volume of interest.

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Authors’ contributions
JP, LH and CP designed the study. JP and LH performed the PET/CT data recalculation and SUV quantification. GR and GSZ analysed the blinded patient data. GZ supervised the statistical analyses. JP drafted the manuscript. MB and CP supervised the study. All authors read and approved the final manuscript.

Table 6  Comparison of studies focusing on PET acquisition time

| Author          | Study type | [18F]-FDG activity | PET/CT scanner & reconstruction | PET acquisition time | Findings                        |
|-----------------|------------|--------------------|---------------------------------|----------------------|---------------------------------|
| Halpern et al.  | patient    | 7.77 MBq/kg/bw     | Reveal RT PET/CT + 3D OSEM      | 120s, 60s            | 60s/BP only for patients < 59 kg |
| Murray et al.   | phantom + patient | 350 ± 40 MBq       | Gemini TF PET/CT + 3D-RAMLA & OSEM TOF | 60s, 30s, 15s, 10s   | 15s/BP: approved lesion detectability |
| Sonni et al.    | patient    | 356 ± 37 MBq       | Discovery MI PET/CT + 3D PSF TOF | 120s, 90s, 60s, 30s  | 90s/BP with good image quality  |
| Pilz et al.     | patient    | 4 MBq/kg/bw        | Ingenuity TF PET/CT + 3D OSEM TOF | 75s, 57s             | 57 s/BP = Approved acquisition time |

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Availability of data and materials
The datasets used and analysed during this study are available from the corresponding author on reasonable request.

Declarations
Ethics approval
This retrospective study was conducted according to the principles of the Declaration of Helsinki and all subsequent revisions and was approved by the routing Ethics Committee of the province Salzburg (EC number: 415-E/2491/2-2019).

Consent for publication
Not applicable.

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

1. Prieto E, García-Velloso MJ, Rodríguez-Fraile M, Morán V, García-García B, Guillén F, et al. Significant dose reduction is feasible in FDG PET/CT protocols without compromising diagnostic quality. Phys Med. 2018;46:134–9.

2. Boellaard R, Delgado-Bolton R, Oyen WJG, Giammarile F, Tatsch K, Eschner W, et al. FDG PET/CT. EANM procedure guidelines for tumour imaging: version 2.0. Eur J Nucl Med Mol Imaging. 2015;42(2):328–54.

3. Slomka PJ, Pan T, Germann G. Recent advances and future progress in PET instrumentation. Semin Nucl Med. 2016;46(1):1–19.

4. van Sluis J, Boellaard R, Somasundaram A, van Snick PH, Bozza RH, Dierckx RAJO, et al. Image quality and semiquantitative measurements on the biograph vision PET/CT system: initial experiences and Comparison with the Biograph mCT. J Nucl Med. 2020;61(1):129–35.

5. Boellaard R, O’Doherty MJ, Weber WA, Mortzagh 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.

6. Baratto L, Park SY, Hatami N, Davidzon G, Srinivas S, Gambhir SS, et al. 18F-FDG silicon photomultiplier PET/CT: a pilot study comparing semi-quantitative measurements with standard PET/CT. PloS One. 2017;12(6):e0178936.

7. van der Vos CS, Koopman D, Rijnsdorp S, Arends AJ, Boellaard R, van Sluis J, et al. Optimizing injected dose in clinical PET by accurately modeling the radioactivity of 18F-FDG for reliable semiquantification in PET/CT imaging. Ann Nucl Med. 2015;29(11):693–705.

8. Watson CC, Case ME, Bendriem B, Carney JP, Townsend DW, Eberl S, et al. Optimizing injected dose in clinical PET by accurately modeling the counting-rate response functions specific to individual patient scans. J Nucl Med. 2005;46(11):1825–34.

9. Vandenberghe S, Mikhaylova E, Giammarile F, Tatsch K, Eschner W, et al. Standards for PET image acquisition and quantitative data analysis. EJNMMI Phys. 2017;4(1):9.

10. Berg E, Cherry SR. Innovations in instrumentation for positron emission tomography. Semin Nucl Med. 2018;48(4):311–31.

11. Boellaard R. Standards for PET image acquisition and quantitative data analysis. J Nucl Med. 2009;50(Suppl 1):115–20.

12. Karakatsanis NA, Fokou E, Tsoumpas C. Dosage optimization in positron emission tomography. Semin Nucl Med. 2018;48(4):311–31.

13. van Sluis J, Boellaard R, Oyen WJG, Giammarile F, Tatsch K, Eschner W, et al. FDG PET/CT. EANM procedure guidelines for tumour imaging: version 2.0. Eur J Nucl Med Mol Imaging. 2015;42(2):328–54.

14. Willowson KP, Bailey EA, Bailey DL. A retrospective evaluation of radiation exposure to staff and distributed time-of-flight list mode PET reconstruction. In: 2006 IEEE nuclear science symposium. 2006. p. 1715–22.

15. Soni I, Baratto L, Park S, Hatami N, Srinivas S, Davidzon G, et al. Initial experience with a SiPM-based PET/CT scanner: influence of acquisition time on image quality. EJNMMI Phys. 2018;5(1):9.

16. Zargan S, Ghafarian P, Shabestani Monfared A, Sharafi AA, Bakshayeshkaram M, Ay MR. Evaluation of radiation exposure to staff and environment dose from 18F-FDG in PET/CT and cyclotron center using thermoluminescent dosimeters. J Biomed Phys Eng. 2017;7(1):12.

17. Annual Congress of the European Association of Nuclear Medicine. October 12-16, 2019 Barcelona, Spain. Eur J Nucl Med Mol Imaging. 2019;46(5):1–952.

18. Wang W, Hu Z, Gualtieri EE, Parma MJ, Walsh E, Sebok D, et al. Systematic and distributed time-of-flight list mode PET reconstruction. In: 2006 IEEE nuclear science symposium. 2006. p. 1715–22.

19. Boellaard R, van Oorschot JAC, Jager PL, Tenbergen CJA, Knollema S, Slump CH, et al. Technical note: how to determine the FDG activity for tumour PET imaging that satisfies European guidelines. EJNMMI Phys. 2016;3(1):22.

20. van Sluis J, Boellaard R, Somasundaram A, van Snick PH, Bozza RH, Dierckx RAJO, et al. Digital PET compliance to EARL accreditation specifications. J Nucl Med. 2020;61(5):764–71.

21. Martin Bland J, Altman D. STATISTICAL METHODS FOR ASSESSING AGREEMENT BETWEEN TWO METHODS OF CLINICAL MEASUREMENT. The Lancet. 1986;327(8476):307–10.

22. Murray I, Kalemis A, Glennon J, Hasan S, Quraishi S, Beyer T, et al. Time-of-flight PET/CT using low-activity protocols: potential implications for cancer therapy monitoring. Eur J Nucl Med Mol Imaging. 2010;37(9):1643–53.

23. Wickham F, McMeekin H, Burns M, McCool D, Pencharz D, Skillen A, et al. Patient-specific optimisation of administered activity and acquisition times for 18F-FDG PET imaging. EJNMMI Res. 2017;7(1):3.

24. Halpern BS, Dahlbom M, Quon A, Schiepers C, Waldherr C, Silverman DH, et al. Impact of patient weight and emission scan duration on PET/CT image quality and lesion detectability. J Nucl Med. 2004;45(5):797–801.

25. McDermott GM, Chowdhury FU, Scarsbrook AF. Evaluation of noise equivalent count parameters as indicators of adult whole-body FDG-PET image quality. Ann Nucl Med. 2013;27(9):855–61.

26. Fukukita H, Suzuki K, Matsumoto K, Terauchi T, Daisaki H, Ikari Y, et al. Japanese guideline for the oncology FDG-PET/CT data acquisition protocol: synopsis of version 2.0. Ann Nucl Med. 2014;28(7):693–705.

27. Martin Bland J, Altman D. STATISTICAL METHODS FOR ASSESSING AGREEMENT BETWEEN TWO METHODS OF CLINICAL MEASUREMENT. The Lancet. 1986;327(8476):307–10.

28. Watson CC, Case ME, Bendriem B, Carney JP, Townsend DW, Eberl S, et al. Optimizing injected dose in clinical PET by accurately modeling the counting-rate response functions specific to individual patient scans. J Nucl Med. 2005;46(11):1825–34.

29. Masuda Y, Kondo C, Matsuo Y, Uetani M, Kusakabe K. Comparison of imaging protocols for 18F-FDG PET/CT in overweight patients: optimizing scan duration versus administered dose. J Nucl Med. 2009;50(6):844–8.

30. Chen MK, Menard DH, Cheng DW. Determining the minimal required radioactivity of 18F-FDG for reliable semiquantification in PET/CT imaging: a phantom study. J Nucl Med Technol. 2016;44(1):26–30.

31. van Sluis J, Boellaard R, Dierckx RAJO, Stormezand GN, Glaudemans AWJM, Noordzij W. Image quality and activity optimization in oncologic 18F-FDG PET using the digital biograph vision PET/CT system. J Nucl Med. 2020;61(5):764–71.

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