Synthesis, Biodistribution, and Radiation Dosimetry of a Novel mGluR5 Radioligand: $^{18}$F-AZD9272

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ABSTRACT: The metabotropic glutamate receptor subtype mGluR5 has been proposed as a potential drug target for CNS disorders such as anxiety, depression, Parkinson’s disease, and epilepsy. The AstraZeneca compound AZD9272 has previously been labeled with carbon-11 and used as a PET radioligand for mGluR5 receptor binding. The molecular structure of AZD9272 allows one to label the molecule with fluorine-18 without altering the structure. The aim of this study was to develop a fluorine-18 analogue of AZD9272 and to examine its binding distribution in the nonhuman primate brain in vivo as well as to obtain whole body radiation dosimetry. $^{18}$F-AZD9272 was successfully synthesized from a nitro precursor. The radioligand was stable, with a radiochemical purity of >99% at 2 h after formulation in a sterile phosphate buffered solution (pH = 7.4). After injection of $^{18}$F-AZD9272 in two cynomolgus monkeys, the maximum whole brain radioactivity concentration was 4.9−6.7% of the injected dose ($n = 2$) and PET images showed a pattern of regional radioactivity consistent with that previously obtained for $^{11}$C-AZD9272. The percentage of parent radioligand in plasma was 59 and 64% ($n = 2$) at 120 min after injection of $^{18}$F-AZD9272, consistent with high metabolic stability. Two whole body PET scans were performed in nonhuman primates for a total of 231 min after injection of $^{18}$F-AZD9272. Highest uptakes were seen in liver and small intestine, followed by brain and kidney. The estimated effective dose was around 0.017 mSv/MBq. $^{18}$F-AZD9272 shows suitable properties as a PET radioligand for in vivo imaging of binding in the primate brain. $^{18}$F-labeled AZD9272 offers advantages over $^{11}$C-AZD9272 in terms of higher image resolution, combined with a longer half-life. Moreover, based on the distribution and the estimated radiation burden, imaging of $^{18}$F-AZD9272 could be used as an improved tool for quantitative assessment and characterization of AZD9272 binding sites in the human brain by using PET.

KEYWORDS: PET, mGluR5 radioligands, NHP, fluorine-18, kinetics, dosimetry

INTRODUCTION

Glutamate is the brain’s main excitatory neurotransmitter primarily located on the membranes of neuronal and glial cells and is present in over 50% of nervous tissue. It is particularly abundant in the human nervous system and mostly prominent in the human brain. Glutamate acts through ionotrophic (NMDA, kainite, and AMPA) and metabotropic glutamate

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Metabotropic glutamate receptors (mGluRs) are a family of G protein-coupled receptors. In 1992, mGluR5 was first cloned in animals and followed by humans several years later. Even though its actions are mostly excitatory, there are strong links and receptor interactions between mGluR5 and the NMDA receptor. It has also been reported that the activation of mGluR5 enhances GABA, especially in the nucleus accumbens. The density of mGluR5 is high primarily in the forebrain regions, striatum, and limbic regions. Moreover, the density of mGluR5 is much higher in younger animals than in adults suggests that early intervention targeting the mGluR5 may lead to prevent the neurodevelopmental disorders.

It is believed that presynaptic mGluR5 receptors participate in the regulation of synaptic plasticity and changes in neuronal excitability to maintain homeostasis. Because of its functions in different neuronal processes, mGluR5 has been suggested as a therapeutic drug target for several neurological disorders such as Parkinson’s diseases, anxiety, depression, schizophrenia, seizure disorder, addiction as well as various chronic pain states.

Positron emission tomography (PET) has been widely utilized in visualizing the localization of mGluR5. Imaging brain mGluR5 with PET in humans has been useful for studying neurodegenerative disorders and addiction. Several radioligands (Figure 1) have been developed and applied preclinically or clinically to image brain mGluR5. Most of those are diaryl alkyne, structural analogues of the prototype mGluR5 NAM MPEP. A noncompetitive mGluR5 antagonist AZD9272 which does not depend on the presence of alkyne moiety was developed by AstraZeneca. AZD9272 was previously labeled with carbon-11 and evaluated as a potential mGluR5 radioligand in nonhuman primates (NHPs) and human subjects. However, due to the short half-life of carbon-11 (20.4 min), the use of such a radioligand is restricted to imaging facilities that are close to the site of production. In addition, a three step synthesis of 11C-AZD9272 using palladium mediated 11C-cyanation was associated with difficulties in automation.

Since AZD9272 has two fluorine functional groups, it allows the radiolabeling with 18F from the corresponding nitro precursor without altering the structure. Fluorine-18 labeled PET tracers may provide advantages for use in PET imaging. Relatively long half-life (109.8 min) facilitates imaging at later time-points and lower positron energy (0.635 MeV) allows higher intrinsic resolution in the PET images. Fluorine-18...
 labeled radiotracers can be prepared in multiple clinical doses for long-distance distribution, enabling widespread clinical use.

The objectives of the present PET-study were to (i) to develop a fast and efficient synthetic method for labeling of a mGluR5 ligand AZD9272 with fluorine-18, (ii) to evaluate its binding in vivo in the NHP brain, and (iii) to provide dosimetry estimates for 18F-AZD9272 based on NHP whole-body PET measurements.

RESULTS AND DISCUSSION

Radiochemistry. The radiolabeling of 18F-AZD9272 (3-fluoro-5-(3-(5-[18F]fluoropyridin-2-yl)-1,2,4-oxadiazol-5-yl)benzonitrile) was achieved by nucleophilic substitution of the corresponding nitro-precursor (3-fluoro-5-(3-(5-nitrophenyl-2-yl)-1,2,4-oxadiazol-5-yl)benzonitrile) by 18F-fluoride in the presence of K2.2.2 and K2CO3 as shown in Figure 2. Different solvents such as acetonitrile, DMF, and DMSO were tested at different temperatures (Table 1). A combination of DMSO as the reaction solvent and 150 °C reaction temperature for 10 min resulted in the desired product with the best radiochemical yield. After HPLC purification, 1.9–3.2 GBq of 18F-AZD9272 was obtained from 15–20 min irradiation at 35 μA. The average radiochemical yield of the radiosynthesis was 23% (nondecay corrected), and the radiochemical purity was higher than 99%. The overall radiosynthesis including the fluorination reaction, HPLC purification, SPE isolation, and radiotracer formulation was completed within 70–75 min. The identity of the labeled compound was confirmed by co-injection of their corresponding fluorine-19 analogue using analytical HPLC. The radioligand 18F-AZD9272 was found to be stable in PBS buffered solution (pH = 7.4) for the duration of 120 min. 18F-AZD9272 was obtained with a molar activity (MA) of 94 ± 103 GBq/μmol (n = 6) at the time of injection.

Brain PET. The injected radioactivity of 18F-AZD9272 was 155 and 155 MBq for NHP1 and NHP2, respectively. The MA at the time of injection was 45 and 31 GBq/μmol, and the injected mass was 0.98 and 1.46 μg. Fusion images of MRI and summed PET of NHP2 are shown in Figure 3A. The total distribution volume (Vt) obtained by the two tissue compartment (2TC) model and Logan graphical analysis (GA) using metabolite corrected plasma radioactivity are shown in Table 2. Vt’s by GA are well correlated to those by 2TC although with slightly lower values (11% and 6%) (Figure 4A). Bias is quite small as within 3% at 75 min for both 2TC and GA (Figure 4B).

The brain uptake of 18F-AZD9272 is consistent with that previously reported for 11C-AZD9272.33 Also, the Vt values obtained by 2TC are similar between 18F-AZD9272 and 11C-AZD9272, e.g., 17.3 vs 18.0 mL/cm3 (ventral striatum) and 5.7 vs 6.5 mL/cm3 (occipital cortex).

Vt values estimated by GA were well correlated with those by 2TC, although slightly lower values were obtained. Additionally, a short length of measurement showed a small bias in Vt values obtained with both 2TC and GA. This indicates that GA with a short duration of measurement (e.g., 75 min) could be used for the estimation for Vt of 18F-AZD9272.

The consistent binding pattern observed with 18F- and 11C-labeled AZD9272 further confirms that the regional brain distribution for radiolabeled AZD9272 differs from that for other mGluR5 radioligands.33 Observations elsewhere suggest that AZD9272 displays additional, non-mGluR5-related binding that may represent specific binding to monoamine oxidase-B (MAO-B). Based on our previous observations,31 close to 90% of specific binding for 18F-AZD9272 in NHP brain can be inhibited by administration of a MAO-B-selective ligand. Moreover, the MAO-B component of 11C-AZD9272 binding has been estimated to be approximately 90–95% in human brain stem, thalamus, amygdala, cerebel- lum, and ventral striatum. The significant contribution MAO-B binding to the signal for AZD9272 suggests that 18F-AZD9272 could serve as a radioligand for assessment of MAO-B availability in these regions of the primate brain.

Figure 2. Radiosynthesis of 18F-AZD9272.

Table 1. Optimization of Radiolabeling of 18F-AZD9272

| entry | precursor (mg) | solvent  | reaction time (min) | reaction temperature (°C) | yield (%) |
|-------|----------------|---------|---------------------|--------------------------|-----------|
| 1     | 2              | acetonitrile | 5                   | 75                       | 1         |
|       |                |          |                     | 90                       | 1         |
|       |                |          |                     | 110                      | 3         |
|       |                |          |                     | 15                       | 75        |
|       |                |          |                     | 90                       | 1         |
|       |                |          |                     | 110                      | 5         |
|       |                |          |                     | 30                       | 90        |
|       |                |          |                     | 100                      | 2         |
|       |                | DMSO     | 5                   | 90                       | 0         |
|       | 2              | DMSO     | 5                   | 120                      | 1         |
|       |                |          |                     | 135                      | 3         |
|       |                |          |                     | 150                      | 5         |
|       |                |          |                     | 90                       | 0         |
|       |                |          |                     | 120                      | 1         |
|       |                |          |                     | 135                      | 5         |
|       |                |          |                     | 150                      | 5         |
|       |                |          |                     | 15                       | 120       |
|       |                |          |                     | 150                      | 23        |
|       |                |          |                     | 165                      | 22        |
|       |                |          |                     | 165                      | 23        |
| 3     | 4–6            | DMSO     | 15                  | 150                      | 23        |
| 4     | 2              | DMF      | 15                  | 125                      | 2         |
|       |                |          |                     | 145                      | 15        |
|       |                |          |                     | 145                      | 4         |
|       |                |          |                     | 155                      | 15        |
Nevertheless, characterization as a tracer for imaging brain MAO-B would require comparison of regional AZD9272 binding with that of established MAO-B radioligands in the same subjects. Taken together, 18F-AZD9272 shows favorable properties in terms of improved stability and image resolution and offers advantages over 11C-AZD9272 for further identification and characterization of AZD9272 binding to MAO-B.

Radiometabolite Analysis. The recovery of radioactivity from plasma into acetonitrile after deproteinization was higher than 95%. The parent compound was more abundant at 5 min representing approximately 95% and it decreased to 65% at 120 min (Figure 5A). The identity of the radio metabolite 18F-AZD9272 was confirmed by coinjection with the authentic nonradioactive AZD9272. The similar time-course for radio metabolism was observed for 11C-AZD9272 (Figure 5B). HPLC analysis of plasma following injection of 18F-AZD9272, parent compound and one radioactive metabolite was detected which eluted at 6.7 and 6.3 min (Figure 5C). The abundance of the radiometabolite at 6.3 min increases with time and representing approximately 4% at 5 min and it increased to 17% at 120 min. The identity of the radiometabolite was not analyzed. In principle, it is a possibility that the metabolite is sufficiently lipophilic to enter the brain. However, according to the time stability analysis, the bias to the duration of measurements was small (Figure 4B). This observation provides indirect support for the view that the radiometabolite is less likely to impact the quantitative estimates.

Whole Body PET. The injected radioactivity of 18F-AZD9272 was 217 and 199 MBq for the two NHPs (NHP3 and NHP4) respectively. The MA at the time of injection was 41 and 58 GBq/μmol, and the injected mass was 1.53 and 0.97 μg. CT and PET images of maximum intensity projection (MIP) over time of NHP4 are shown in Figure 6. High uptakes were seen in liver and small intestine, followed by brain and kidney. Human radiation dose estimates indicate that most organs appear to receive around 0.01−0.02 mSv/MBq (Table 2).

Table 2. Vt Using the 2TC Model and GA with Metabolite Corrected Plasma Activity

|          | 2TC (mL/cm³) | GA (mL/cm³) |
|---------|-------------|-------------|
|         | NHP1 | NHP2 | mean | NHP1 | NHP2 | mean |
| ACC     | 11.5  | 11.2  | 11.4  | 10.2  | 10.7  | 10.4  |
| AMG     | 15.0  | 14.4  | 14.7  | 12.8  | 13.1  | 13.0  |
| CAU     | 13.3  | 13.7  | 13.5  | 12.1  | 13.0  | 12.6  |
| CER     | 7.1   | 7.9   | 7.5   | 6.6   | 7.6   | 7.1   |
| HIP     | 10.7  | 11.9  | 11.3  | 9.7   | 11.2  | 10.5  |
| INS     | 9.0   | 10.0  | 9.5   | 8.5   | 9.6   | 9.1   |
| OC      | 5.4   | 6.0   | 5.7   | 5.5   | 5.7   | 5.6   |
| PFC     | 9.6   | 9.5   | 9.5   | 8.9   | 9.3   | 9.1   |
| PUT     | 9.5   | 10.7  | 10.1  | 9.3   | 10.1  | 9.7   |
| TC      | 7.6   | 8.4   | 8.0   | 7.2   | 8.1   | 7.7   |
| THA     | 11.7  | 11.4  | 11.6  | 10.6  | 10.8  | 10.7  |
| VM      | 11.0  | 8.9   | 9.9   | 9.1   | 8.4   | 8.8   |
| VST     | 17.3  | 17.3  | 17.3  | 14.5  | 15.6  | 15.1  |

ACC, anterior cingulate cortex; AMG, amygdala; CAU, caudate; CER, cerebellum; HIP, hippocampus; INS, insular cortex; OC, occipital cortex; PFC, prefrontal cortex; PUT, putamen; TC, temporal cortex; THA, thalamus; VM, ventral midbrain; VST, ventral striatum.

Figure 3. (A) PET images of 18F-AZD9272 coregistered with MRI and averaged between 9 and 120 min in the transaxial (left), sagittal (middle), and coronal (right) projections. (B) Time–activity curves for 18F-AZD9272 in different brain regions.
Materials and Methods

General. Liquid chromatographic analysis (LC) was performed with a Merck-Hitachi gradient pump and a Merck-Hitachi, L-4000 variable wavelength UV-detector. LC-MS was performed using a Waters Quatta-ToF Premier micro mass spectrometer, or Waters SQD 3001 single quadrupole mass spectrometer, coupled to Waters Acquity UPLC instruments. The precursor (3-fluoro-5-(3-(5-nitropyridin-2-yl)-1,2,4-oxadiazol-5-yl)benzonitrile) and the reference standard nonradioactive AZD9272 (3-fluoro-5-(3-(5-fluoropyridin-2-yl)-1,2,4-oxadiazol-5-yl)benzonitrile) were supplied by AstraZeneca AB. All other chemicals and reagents were obtained from commercial sources and used without any further purification. Solid-phase extraction (SPE) cartridges SepPak QMA light and SepPak C18 Plus were purchased from Waters (Milford, MA). The C18 Plus cartridge was activated using EtOH (10 mL) followed by sterile water (10 mL). The SPE cartridge SepPak QMA light was activated using K₂CO₃ solution (0.5M, 10 mL) followed by water (15 mL, 18 MΩ).

Fluorine-18 fluoride was produced at Karolinska Hospital (Stockholm, Sweden)

Production of ¹⁸F-Fluoride (¹⁸F–F⁰). Fluorine-18 fluoride (¹⁸F–F⁰) was produced from a GEMS PET trace Cyclotron using 16.4 MeV protons via the ¹⁸O(p,n)¹⁸F nuclear reaction on ¹⁸O-enriched water (¹⁸O–H₂O). After irradiation, ¹⁸F–F⁰ was isolated from ¹⁸O–H₂O on a light anion exchange cartridge (SepPak QMA) and subsequently eluted from the cartridge with a solution of K₂CO₃ (9.75 μmol, 1.35 mg), Kryptofix₂₂₂ (4,7,13,16,21,24-hexaoxa-1,10-diazabicyclo[8.8.8]octadec-7-yl) (19.5 μmol, 7.35 mg) in water (64 μL, 18 MΩ), and MeCN (1.5 mL) to a 10 mL screw cap reaction vessel. The solvents were evaporated at 150 °C for 10 min under continuous nitrogen flow (70–80 mL/min) to form a dry complex of ¹⁸F–F⁰/K₂CO₃/K₂₂₂. The residue was cooled to room temperature (RT) before the precursor (3-fluoro-5-(3-(5-nitrophenyl)-1,2,4-oxadiazol-5-yl)benzonitrile) was added.

Synthesis of ¹⁸F-AZD9272 (3-Fluoro-5-(3-(5-[¹⁸F]fluoropyridin-2-yl)-1,2,4-oxadiazol-5-yl)benzonitrile). To the dry complex of ¹⁸F–F⁰/K₂CO₃/K₂₂₂, nitro-precursor (3-fluoro-5-(3-(5-nitrophenyl)-1,2,4-oxadiazol-5-yl)benzonitrile) (2 mg) in DMSO (500 μL) was added at 150 °C and left for 15 min to produce ¹⁸F-AZD9272. The reaction mixture was cooled to RT and was diluted with water to a total volume of 3 mL before injecting to a semipreparative reverse phase μ-Bondapak HPLC column (C18, 7.8 Ω × 300 mm, 10 μm, Waters) for purification. The column outlet was connected to an UV absorbance detector (λ= 270 nm) coupled to a radioactive detector (μ-Bondapak HPLC column (C18, 7.8 Ω × 300 mm, 10 μm, Waters) for purification. The column outlet was connected to an UV absorbance detector (λ = 270 nm) in series with a GM tube for radioactivity detection. Elution with mobile phase CH₃CN/0.1% TFA with a gradient (RT) before the precursor (3-fluoro-5-(3-(5-nitrophenyl)-1,2,4-oxadiazol-5-yl)benzonitrile) was added.

Quality Control of ¹⁸F-AZD9272. The radiochemical purity, identity, and stability of ¹⁸F-AZD9272 was determined by using an analytical HPLC system which included a ACE RP column (C18, 3.9 Ω × 250 mm, 5 μm particle size), Merck-Hitachi L-7100 Pump, L-7400 UV detector, and GM tube for radioactivity detection (YVR International). The mobile phase CH₃CN/0.1% TFA with a gradient HPLC method (10–90% in 8 min) and flow rate of 3 mL/min was used to elute the product. The effluent was monitored with a UV absorbance detector (λ = 270 nm) coupled to a radioactive detector (b-flow, Beckman, Fullerton, CA). The retention time (RT) of ¹⁸F-AZD9272 was 4–5 min. The identity of ¹⁸F-AZD9272 was confirmed by using HPLC with the coinjection of the authentic nonradioactive AZD9272 (3-fluoro-5-(3-(5-fluoropyridin-2-yl)-1,2,4-oxadiazol-5-yl)benzonitrile) standard.

Molar Activity (MA) Determination. The MA of the final product was measured by analytical HPLC which included a ACE RP column.
column (C18, 3.9 mm I.D. × 250 mm, 5 μm particle size) using mobile phase CH3CN/50 mM H3PO4 (35/65) and flow rate of 2 mL/min. MA was calibrated for UV absorbance (λ = 270 nm) response per mass of ligand and calculated as the radioactivity of the radioligand (GBq) divided by the amount of the associated carrier substance (μmol). Each sample was analyzed three times and compared to a reference standard also analyzed three times.

PET Measurements in a Cynomolgus Monkey. The study was approved by the Animal Ethics Committee of the Swedish Animal Welfare Agency (Dnr N185/14) and was performed according to “Guidelines for planning, conducting and documenting experimental research” (Dnr 4820/06-600) of Karolinska Institutet. The NHPs were housed in the Astrid Fagraeus Laboratory (AFL) of the Swedish Institute for Infectious Disease Control, Solna, Sweden.

Brain PET. Two male cynomolgus monkeys (NHP1: 5460 g and NHP2: 7080 g) were used in this study. Parts of the data have been reported.32 Anesthesia was induced by intramuscular injection of ketamine hydrochloride (approximately 10 mg/kg) at AFL and maintained by intravenous infusion of ketamine (4 mg/kg/h) and xylazine (0.4 mg/kg/h) with a pump. The NHP head was immobilized with a fixation device. Body temperature was maintained by using a Bair Hugger system (model 505, Arizant Healthcare, MN) and monitored with an esophageal thermometer. Heart rate, blood pressure, respiratory rate, and oxygen saturation were continuously monitored throughout the experiments. Fluid balance was maintained by continuous infusion of saline.

PET measurements were conducted using a High Resolution Research Tomograph (HRRT) PET scanner (Siemens Molecular Imaging, Knoxville, TN). A transmission scan of 6 min using a single 137Cs source was performed before the 18F-AZD9272 injection. List-mode data were reconstructed with a series of 34 frames (20 s × 9, 1 min × 3, 3 min × 5, and 6 min × 17), using the ordinary Poisson-3D-ordered subset expectation maximization (OP-3D-OSEM) algorithm with 10 iterations and 16 subsets including modeling of the point spread function (PSF). Only a 105 min image of NHP1 was obtained due to the technical issue.

An automated blood sampling system (ABSS) was used to measure the continuous radioactivity for the first 3 min after the radioligand injection. Then, blood sampling was performed manually for the measurement of metabolism and radioactivity at 5, 15, 30, 60, 90, and 120 min after the injection. A reversed-phase radio-HPLC method was used to determine the amount of unchanged 18F-AZD9272 and its radioactive metabolites in NHP plasma.35

Figure 5. Radiometabolite analysis during the course of the PET measurements. It is shown as the relative plasma composition of radiometabolites and parent compound in percent of total plasma radioactivity that was injected. (A) In vivo radiometabolism of 18F-AZD9272 was detected during the 120 min PET scan. (B) In vivo radiometabolism of 11C-AZD9272 was detected during the 30 min PET scan. (C) Representative HPLC chromatogram 15 min after injection of 18F-AZD9272.
Figure 6. Whole body images of CT and PET.

Figure 7. Time–activity curves of percent of injected dose: (A) high uptake organs and (B) low uptake organs.
Whole Body PET. Two female cynomolgus monkeys (NHP3: 7420 g and NHP4: 6360 g) were used in this study. Anesthesia was induced by intramuscular injection of ketamine hydrochloride (approximately 10 mg/kg) at AFL and maintained by intravenous infusion of ketamine (4 mg/kg/h) and xylazine (0.4 mg/kg/h) with a pump. The body of the NHP was immobilized using a vacuum pad. Body temperature was maintained by using a Bair Hugger system (approximately 10 mg/kg) at AFL and maintained by intravenous infusion of saline.

Whole-body PET scans were conducted using a GE Discovery PET/CT 710 system (GE Healthcare, Waukesha, WI). One low-dose CT scan was performed before intravenous administration of $^{18}$F-AZD9272 for attenuation correction. Then five series of PET acquisitions, each covering five axial fields of view (AFOV), were conducted. The five PET series consisted of two 20 s × 5 AFOV, three 40 s × 5 AFOV, three 80 s × 5 AFOV, three 160 s × 5 AFOV, and eight 240 s × 5 AFOV. The total duration of the whole-body PET measurement was around 4 h. PET images were reconstructed with a 3D ordered-subset expectation maximization (OSEM) algorithm with three iterations and 18 subsets, including the time-of-flight information (VUE Point FX) and the point spread function correction (Sharp IR). A 2D Gaussian filter with 5.5 mm cutoff was used.

ROIs were drawn for the brain, heart, liver, lung, kidney, gall bladder, bone (limbar vertebrae), urinary bladder, stomach, small intestine, spleen, and pancreas with the help of the CT images for anatomic landmarks. The time—activity curve was expressed as the percentage of the injected dose (%ID) calculated as follows: decay-corrected radioactivity (Bq/cc) × ROI volume (cc)/injected dose (Bq) × 100.

Estimates of the absorbed radiation dose in humans were calculated with OLINDA/EXM 1.1 (Organ Level INternal Dose Assessment Code) software, using the adult male (70 kg) reference model. The fractional uptake in NHP organs was assumed to be equal to the uptake in human organs.

Radiometabolite Analysis. Radiometabolite analysis was performed following a previously published method. In short, a reverse-phase HPLC method was used for the determination of the percentages of radioactivity corresponding to unchanged radioligand $^{18}$F-AZD9272 and its radioactive metabolites during the course of a PET measurement. Venous blood samples (2 mL) were obtained from the monkey at different time point such as 5, 15, 30, 45, 60, 90, and 120 min after injection of $^{18}$F-AZD9272. Collected blood (2 mL) was centrifuged at 2000 g for 2 min to obtain the plasma (0.5 mL). The plasma obtained after centrifugation of blood at 2000 g for 2 min was mixed with a 1.4 times volume of acetonitrile. The mixture was then centrifuged at 2000 g for 4 min, and the extract was separated from the pellet and was diluted with water before injecting it into the HPLC system coupled to an online radioactivity detector. An Agilent binary pump (Agilent 1200 series) coupled to a manual injection valve (7725i, Rheodyne), 1–3.0 mL loop and a radiation detector (Oyokoken, S-24932) housed in a shield of 50 mm thick lead was used for metabolite measurements. Data collection and control of the LC system was performed using chromatographic software (ChemStation Rev. B.04.03; Agilent). The accumulation time of the radiation detector was 10 s. Chromatographic separation was achieved on an ACE C18 column (250 mm × 10 mm ID) by gradient elution. Acetonitrile (A) and 10 mM ammonium formate (B) were used as the mobile phase at 5.0 mL/min, according to the following program: 0–8.5 min, (A/B) 50:50 → 95:5 v/v; 8.5–11.0 min, (A/B) 95:5 v/v. Peaks for radioactive compounds eluting from the column were integrated, and their areas were expressed as a percentage of the sum of the areas of all detected radioactive compounds (decay-corrected to the time of injection on the HPLC).

To calculate the recovery of radioactivity from the system, an aliquot (2 mL) of the eluate from the HPLC column was measured and divided with the amount of total injected radioanalytes.

### Table 3. Estimated Radiation Dose at Different Organs

| Organ             | NHP3 (μSv/MBq) | NHP4 (μSv/MBq) | mean (μSv/MBq) |
|-------------------|----------------|----------------|---------------|
| adrenals          | 14.6           | 14.1           | 14.4          |
| brain             | 13.0           | 17.4           | 15.2          |
| breasts           | 8.0            | 7.9            | 7.9           |
| gallbladder wall  | 21.3           | 21.1           | 21.2          |
| LLI wall          | 26.8           | 27.9           | 27.4          |
| small intestine   | 58.9           | 62.1           | 60.5          |
| stomach wall      | 13.2           | 13.1           | 13.2          |
| ULI wall          | 65.7           | 69.4           | 67.6          |
| heart wall        | 17.5           | 19.0           | 18.3          |
| kidneys           | 42.3           | 29.6           | 36.0          |
| liver             | 47.0           | 45.7           | 46.4          |
| lungs             | 12.7           | 11.8           | 12.3          |
| muscle            | 10.1           | 10.0           | 10.1          |
| ovaries           | 18.2           | 18.6           | 18.4          |
| pancreas          | 16.4           | 15.4           | 15.9          |
| red marrow        | 10.7           | 10.7           | 10.7          |
| osteogenic cells  | 14.5           | 14.5           | 14.5          |
| skin              | 7.4            | 7.4            | 7.4           |
| spleen            | 9.6            | 8.4            | 9.0           |
| testes            | 8.7            | 8.6            | 8.7           |
| thymus            | 9.7            | 9.6            | 9.6           |
| thyroid           | 9.1            | 9.1            | 9.1           |
| urinary bladder wall | 11.3         | 11.4           | 11.4          |
| uterus            | 16.1           | 16.3           | 16.2          |
| total body        | 12.2           | 12.1           | 12.2          |
| effective dose    | 17.4           | 17.4           | 17.4          |

The regions of interest (ROIs) were delineated manually on the MRI images of each NHP for the anterior cingulate cortex, amygdala, caudate, cerebellum, hippocampus, insular cortex, occipital cortex, prefrontal cortex, putamen, temporal cortex, thalamus, ventral midbrain, and ventral striatum. The MRI of the individual NHP was co-registered to summed PET images of the whole measurement. By applying the co-registration parameters to ROIs, the time—activity curves of brain regions were generated from dynamic PET data.

As the main outcome measure, the total distribution volume ($V_T$) defined as ($K_i/k_e$)($k_i/k_e + 1$) by the two tissue compartment (2TC) model and Logan graphical analysis (GA) was calculated with the metabolite corrected plasma radioactivity as the input function. The acquisition data was truncated to 63 min from 105 min (NHP1) or 123 min (NHP2) to check the time stability of $V_T$. The procedure has been described in detail in a previous publication on $^{[15]}$C]AZD9272 binding.33

### Research Article

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ABBREVIATIONS
mGluRS, metabotropic glutamate receptor; PET, positron emission tomography; PD, Parkinson’s disease; NHP, nonhuman primate; HPLC, high performance liquid chromatography; LC-MS, liquid chromatography—mass spectrometry; DMSO, dimethyl sulfoxide; DMF, dimethylformamide; PBS, phosphate buffered solution; MA, molar activity; HRRT, High Resolution Research Tomograph; SUV, standard uptake value; ROI, region of interest; \( V_t \), distribution volume; TC, tissue compartment; GA, graphical analysis

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