RESEARCH

Region-specific blood–brain barrier transporter changes leads to increased sensitivity to amisulpride in Alzheimer’s disease

Gayathri Nair Sekhar1, Alice L. Fleckney1, Sevda Tomova Boyanova1, Huzefa Rupawala1, Rachel Lo1, Hao Wang1, Doaa B. Farag1,2, Khondaker Miraz Rahman1, Martin Broadstock3,4, Suzanne Reeves5 and Sarah Ann Thomas1*

Abstract

Background: Research into amisulpride use in Alzheimer’s disease (AD) implicates blood–brain barrier (BBB) dysfunction in antipsychotic sensitivity. Research into BBB transporters has been mainly directed towards the ABC superfamily, however, solute carrier (SLC) function in AD has not been widely studied. This study tests the hypothesis that transporters for organic cations contribute to the BBB delivery of the antipsychotics (amisulpride and haloperidol) and is disrupted in AD.

Methods: The accumulation of [3H]amisulpride (3.7–7.7 nM) and [3H]haloperidol (10 nM) in human (hCMEC/D3) and mouse (bEnd.3) brain endothelial cell lines was explored. Computational approaches examined molecular level interactions of both drugs with the SLC transporters [organic cation transporter 1 (OCT1), plasma membrane monoamine transporter (PMAT) and multi-drug and toxic compound extrusion proteins (MATE1)] and amisulpride with the ABC transporter (P-glycoprotein). The distribution of [3H]amisulpride in wildtype and 3× transgenic AD mice was examined using in situ brain perfusion experiments. Western blots determined transporter expression in mouse and human brain capillaries.

Results: In vitro BBB and in silico transporter studies indicated that [3H]amisulpride and [3H]haloperidol were transported by the influx transporter, OCT1, and efflux transporters MATE1 and PMAT. Amisulpride did not have a strong interaction with OCTN1, OCTN2, P-gp, BCRP or MRP and could not be described as a substrate for these transporters. Amisulpride brain uptake was increased in AD mice compared to wildtype mice, but vascular space was unaffected. There were no measurable changes in the expression of MATE1, MATE2, PMAT OCT1, OCT2, OCT3, OCTN1, OCTN2 and P-gp in capillaries isolated from whole brain homogenates from the AD mice compared to wildtype mice. Although, PMAT and MATE1 expression was reduced in capillaries obtained from specific human brain regions (i.e. putamen and caudate) from AD cases (Braak stage V–VI) compared to age matched controls (Braak stage 0–II).

Conclusions: Together our research indicates that the increased sensitivity of individuals with Alzheimer’s to amisulpride is related to previously unreported changes in function and expression of SLC transporters at the BBB (in particular PMAT and MATE1). Dose adjustments may be required for drugs that are substrates of these transporters when prescribing for individuals with AD.

Keywords: Amisulpride, MATE1, PMAT, OCT1, Blood–brain barrier, Alzheimer’s

© The Author(s) 2019. This article is licensed under a Creative Commons Attribution 4.0 International License, which permits use, sharing, adaptation, distribution and reproduction in any medium or format, as long as you give appropriate credit to the original author(s) and the source, provide a link to the Creative Commons licence, and indicate if changes were made. The images or other third party material in this article are included in the article’s Creative Commons licence, unless indicated otherwise in a credit line to the material. If material is not included in the article’s Creative Commons licence and your intended use is not permitted by statutory regulation or exceeds the permitted use, you will need to obtain permission directly from the copyright holder. To view a copy of this licence, visit http://creativecommons.org/licenses/by/4.0/. The Creative Commons Public Domain Dedication waiver (http://creativecommons.org/publicdomain/zero/1.0/) applies to the data made available in this article, unless otherwise stated in a credit line to the data.
(parkinsonism, postural hypotension, stroke) and mortality than other diagnostic groups [1, 2]. This has led to restrictions on the national health service (NHS) use of this class of drugs in the pharmacological management of psychosis and agitation in dementia. Emerging evidence from research into amisulpride use in older people with Alzheimer’s disease (AD) psychosis suggests that blood–brain barrier (BBB) dysfunction may be an important contributor to this heightened sensitivity [3, 4].

Amisulpride is a benzamide derivative, second-generation antipsychotic drug, used to treat schizophrenia [5] and a drug for which the optimal dose (400–800 mg/day), blood concentration (100–319 ng/ml) and striatal dopamine D2/3 receptor occupancy range to avoid non-response and parkinsonism (40–70%) are well established in young adults with schizophrenia [6–8]. Despite being highly selective for dopamine D2/3 receptors [in vitro \(K_i=2.8\) nM] and D3 \(K_i=3.2\) nM] amisulpride has a low propensity to induce parkinsonism, due to its poor BBB penetration and mesolimbic selectivity [9]. In an open treatment study which used amisulpride use in older people with AD psychosis and very late-onset (>60 years) schizophrenia-like psychosis (VLOSLP), treatment response and parkinsonism occurred at very low doses (25–75 mg/day AD, 50–100 mg/day VLOSLP), and at correspondingly low blood drug concentrations (40–100 ng/ml AD, 40–169 ng/ml VLOSLP) due to higher than anticipated striatal dopamine D2/3 receptor occupancies (caudate occupancy, steady state treatment, 50 mg/day amisulpride; 41–83% AD, 41–59% VLOSLP) [3, 10–12]. These findings strongly implicate age and AD-specific changes in central pharmacokinetics in antipsychotic drug sensitivity, particularly at the BBB, which controls drug entry through the expression of transporters [13]. Furthermore, they suggest that amisulpride [14, 15] is a sufficiently sensitive tool with which to probe BBB functionality.

The majority of research into BBB transporters has been directed towards the ABC superfamily, which are ATP-dependent efflux transporters such as P-glycoprotein (P-gp) [16], whose action is compromised in age [17], and more markedly so in AD [18–21]. It has been suggested that amisulpride is a weak P-gp substrate [22, 23], but the importance of P-gp relative to other transporters, especially members of the SLC superfamily, remains unclear. Amisulpride is predominately positively charged (98.9%) at physiological pH (pKa 9.37), and is likely a substrate for the organic cation transporters (OCT) and organic cation transporters novel (OCTN); as observed using the immortalized human cerebral microvessel endothelial cell line (hCMEC/D3) [14]. However, it is also possible that other SLC transporters of organic cations, such as plasma membrane monoamine transporter (PMAT) and multdrug and toxic compound extrusion proteins (MATEs), are involved. Haloperidol, a first generation antipsychotic drug which is highly selective for D2 receptors [24], was chosen as a clinically relevant comparator, as it is a positively charged molecule at pH7.4 and a OCT1 substrate and inhibitor [25, 26] and likely to utilize similar transporters to amisulpride to cross the BBB. Although the drug is no longer used as a first line treatment for psychosis, due to its propensity to cause parkinsonism, it is used at very low doses in the treatment of delirium, which most commonly occurs in those aged over 65 years and with an underlying cognitive impairment (https://www.nice.org.uk/, accessed 15.11.2019). The drug is also extensively used in palliative care and, as a result, is one of the 20 drugs on the WHO list of essential medications.

This study tested the hypothesis that there was an interaction between amisulpride and influx and/or efflux transporters at the BBB which was relevant from a pharmacodynamic perspective by:

1. identifying the transporter involved in the CNS distribution of amisulpride and haloperidol, by examining their kinetic characteristics and inhibitor sensitivity at the human and mouse BBB in vitro.
2. confirming the molecular level interactions of amisulpride and haloperidol with the selected BBB transporters using an in silico computational approach.
3. establishing whether amisulpride access to the CNS is increased in transgenic AD mice, which harbour the human amyloid precursor protein (APP)-Swedish mutation (KM670/671NL), tau mutation (P301L), and presenilin-1 mutation (M146V) compared to wildtype mice.
4. investigating transporter expression in human (and mouse) brain endothelium from age-matched post-mortem AD and healthy aged controls.
5. Examining the type of medications prescribed to individuals with AD and age matched controls.

Overall BBB dysfunction in the AD process and its potential impact on drug delivery in particular on antipsychotic medication will be explored (Fig. 1). The results can also be used to inform further studies [27]. Abstracts of this work have been presented [28, 29].

**Materials and methods**

**Materials**

- \[O\text{-methyl-}^3\text{H}]amisulpride (MW374.8; specific activity 77 Ci/mmol; 97% radiochemical purity) was tritiated (TRQ41291 Quotient, UK). \[^3\text{H}(\text{G})\text{haloperidol (MW375.9; specific activity, 20 Ci/mmol; 99% radiochemical purity: cat# ART1729)}\text{ was purchased from American Radiolabelled Chemicals Inc, St. Louis, Missouri, USA.}\[^{14}\text{C(U)}\text{-sucrose (MW359.48; specific activity 536 mCi/}

**Materials**

- \[O\text{-methyl-}^3\text{H}]amisulpride (MW374.8; specific activity 77 Ci/mmol; 97% radiochemical purity) was tritiated (TRQ41291 Quotient, UK). \[^3\text{H}(\text{G})\text{haloperidol (MW375.9; specific activity, 20 Ci/mmol; 99% radiochemical purity: cat# ART1729)}\text{ was purchased from American Radiolabelled Chemicals Inc, St. Louis, Missouri, USA.}\[^{14}\text{C(U)}\text{-sucrose (MW359.48; specific activity 536 mCi/}

**Materials**

- \[O\text{-methyl-}^3\text{H}]amisulpride (MW374.8; specific activity 77 Ci/mmol; 97% radiochemical purity) was tritiated (TRQ41291 Quotient, UK). \[^3\text{H}(\text{G})\text{haloperidol (MW375.9; specific activity, 20 Ci/mmol; 99% radiochemical purity: cat# ART1729)}\text{ was purchased from American Radiolabelled Chemicals Inc, St. Louis, Missouri, USA.}\[^{14}\text{C(U)}\text{-sucrose (MW359.48; specific activity 536 mCi/}

**Materials**

- \[O\text{-methyl-}^3\text{H}]amisulpride (MW374.8; specific activity 77 Ci/mmol; 97% radiochemical purity) was tritiated (TRQ41291 Quotient, UK). \[^3\text{H}(\text{G})\text{haloperidol (MW375.9; specific activity, 20 Ci/mmol; 99% radiochemical purity: cat# ART1729)}\text{ was purchased from American Radiolabelled Chemicals Inc, St. Louis, Missouri, USA.}\[^{14}\text{C(U)}\text{-sucrose (MW359.48; specific activity 536 mCi/}

**Materials**

- \[O\text{-methyl-}^3\text{H}]amisulpride (MW374.8; specific activity 77 Ci/mmol; 97% radiochemical purity) was tritiated (TRQ41291 Quotient, UK). \[^3\text{H}(\text{G})\text{haloperidol (MW375.9; specific activity, 20 Ci/mmol; 99% radiochemical purity: cat# ART1729)}\text{ was purchased from American Radiolabelled Chemicals Inc, St. Louis, Missouri, USA.}\[^{14}\text{C(U)}\text{-sucrose (MW359.48; specific activity 536 mCi/}

**Materials**

- \[O\text{-methyl-}^3\text{H}]amisulpride (MW374.8; specific activity 77 Ci/mmol; 97% radiochemical purity) was tritiated (TRQ41291 Quotient, UK). \[^3\text{H}(\text{G})\text{haloperidol (MW375.9; specific activity, 20 Ci/mmol; 99% radiochemical purity: cat# ART1729)}\text{ was purchased from American Radiolabelled Chemicals Inc, St. Louis, Missouri, USA.}\[^{14}\text{C(U)}\text{-sucrose (MW359.48; specific activity 536 mCi/}
**a In vitro BBB models (human and mouse) followed by in silico and physicochemical assessments.**

**In vitro BBB models**
- hCMEC/D3
- bEnd.3

**Test molecule**
- Amisulpride
- Haloperidol
- Sucrose

**EXPERIMENT**
- Interaction with transporters
- MTT assay
- Transporter use
- Extracellular space
- Transporter protein expression

**In silico**
- Computational study to explore interaction of test molecules with selected transporters (OCT1, MATE1, PMAT and P-gp).

**In vitro**
- Octanol-saline partition coefficient to assess lipophilicity.

**b In vivo BBB studies in wild type and transgenic mice.**

**GENOTYPE**
- Wildtype (WT; n=6)
- 3xTg AD (n=4)

**AGE**
- WT 3 groups
- 12-13 mths (n=4)
- 16 mths (n=3)
- 24 mths (n=6)

**EXPERIMENT**
- Brain Perfusion Endpoints
- Amisulpride accumulation
- Vascular space
- Two brain regions
- Transporter total protein expression

**C Human tissue studies**

**PHENOTYPE**
- HUMAN
- 2 groups
- Age matched

**EXPERIMENT**
- Total and individual transporter protein expression
- OCT1
- OCTN1 & 2
- MATE1
- MATE2
- PMAT

**Medication history examined**

---

**Fig. 1** Flow charts to provide an overview of the experimental design for the in silico, in vitro and in vivo approaches. Experiments from the three approaches were performed in parallel.
mmol; 99% radiochemical purity: cat# MC266) was purchased from Moravek Biochemicals, USA. Aminopyrine (MW369.5, > 98% purity) was purchased from Cayman Chemicals, UK (cat#71675-85-9). Haloperidol (MW375.9; > 98% purity) was purchased from Sigma-Aldrich, Dorset, UK (cat#H1512). Anti-SLC22A1 antibody (Cat#ab55916; RRID:AB_882579), Anti-SLC22A2 antibody (Cat#ab170871; RRID:AB_2751021), Anti-SLC22A3 antibody (Cat#ab180701; RRID:AB_2751016), Anti-SLC22A4 antibody (Cat#ab200641; RRID:AB_2751017), Anti-SLC22A5 antibody (Cat#ab180757; RRID:AB_2751018), Anti-SLC47A1 antibody (Cat#ab104016: RRID:AB_10711136), Anti-SLC47A2 (Cat#ab174344; RRID:AB_2751019), SLC29A4 antibody (Cat#ab56554: RRID:AB_2190909), Goat Anti-Rabbit IgG H&L (cat#ab6728: RRID:AB_955440) and Rabbit anti-mouse HRP (Cat#ab6728: RRID:AB_955440) were purchased from Abcam, UK. Anti-SLC47A1 (Cat#ab174344; RRID:AB_2751019) was purchased from Alomone Laboratories, Israel. Anti-SLC29A4 (Cat#: bs-4176R: RRID:AB_11108960) was purchased from Bioss antibodies, USA. Goat anti-rabbit (IgG)-HRP was purchased from Cell Signalling. Cat#7074S:AB_2099233). Transferrin receptor monoclonal antibody was purchased from (Thermo Fisher Scientific, CAT#: 13-6800 RRID: AB_2533029). Table 1 details the dilutions used. All consumables were purchased within the years 2012 to 2018 except the transferrin receptor antibody which was purchased in 2019.

In vitro model of the BBB

Cell culture

The hCMEC/D3 (human) and bEnd.3 (mouse) are well-established models of the BBB [21, 30–32]. They are not listed on the misidentified cell line register (version 9 released 14th October 2018). Both lines require different mediums to grow to confluence and their BBB phenotype has been confirmed using Western blots, confocal and transmission electron microscopy [32] (“Western Blot procedure” section). Functional expression of several transporters has been demonstrated by our group [32–34].

a. hCMEC/D3 cells (passages 27–35) were provided under a MTA and maintained in Clonetics® endothelial cell growth medium-2 MV Bullet Kit (Cat# CC-3162 Lonza, UK) containing the endothelial basal medium, the SingleQuotsTM growth factor kit, foe-

Table 1 Primary and secondary antibodies used for protein expression studies

| Protein (Gene) | Primary antibody | Secondary antibody Western blot (WB) |
|---------------|------------------|---------------------------------------|
| OCT-1 (SLC22A1) | Rabbit polyclonal anti-human and mouse (Abcam, Cat#ab55916; RRID:AB_882579), WB dilution—1:250 | Goat anti-rabbit HRP (Abcam, cat#ab6721: RRID:AB_955447) dilution—1:1000 |
| OCT-2 (SLC22A2) | Rabbit monoclonal to human and mouse (Abcam, Cat#ab170871: RRID:AB_2751021), WB dilution—1:2000 | Goat anti-rabbit HRP (Abcam, cat#ab6721: RRID:AB_955447) dilution—1:1000 |
| OCT-3 (SLC22A3) | Rabbit polyclonal to human and mouse (Abcam, Cat#ab183071; RRID:AB_2751016), WB dilution—1:600 | Goat anti-rabbit HRP (Abcam, cat#ab6721: RRID:AB_955447) dilution—1:1000 |
| OCTN1 (SLC22A4) | Rabbit polyclonal to human and mouse (Abcam, Cat#ab200641; RRID:AB_2751017), WB dilution—1:1000 | Goat anti-rabbit IgG-HRP (Cell Signalling. Cat#7074S:AB_2099233) dilution—1:1000 |
| OCTN2 (SLC22A5) | Rabbit polyclonal to human and mouse (Abcam, Cat#ab180757; RRID:AB_2751018), WB dilution—1:1000 | Goat anti-rabbit IgG-HRP (Cell Signalling. Cat#7074S:AB_2099233) dilution—1:1000 |
| MATE1 (SLC47A1) | Rabbit polyclonal to human from Abcam (Cat#104016: RRID AB_10711136), WB dilution—1:500 | Goat anti-rabbit HRP (Abcam, cat#ab6721: RRID:AB_955447) dilution—1:1000 |
| MATE1 (SLC47A1) | Rabbit polyclonal to mouse from Alomone labs (Cat#ANT-131: RRID:AB_2751020), WB dilution—1:800 | Goat anti-rabbit HRP (Abcam, cat#ab6721: RRID:AB_955447) dilution—1:1000 |
| MATE2 (SLC47A2) | Rabbit polyclonal to human and mouse (Abcam, Cat#ab174344: RRID:AB_2751019), WB dilution—1:500 | Goat anti-rabbit HRP (Abcam, cat#ab6721: RRID:AB_955447) dilution—1:1000 |
| PMAT (SLC29A4) | Mouse monoclonal to human and rat (Abcam, Cat#ab56554: RRID:AB_2190909), WB dilution—1:500 | Rabbit anti-mouse HRP (Abcam, Cat#ab6728: RRID:AB_955440) dilution 1:2000 |
| PMAT (SLC29A4) | Rabbit polyclonal to human, mouse and rat (Bioss Antibodies; Cat#:bs-4176R: RRID:AB_11108960), WB Dilution—1:800 for hCMEC/D3—1:650 for b.End3, 1:600 mouse capillaries | Goat anti-rabbit (IgG)-HRP (Cell Signalling. Cat#7074S:AB_2099233) dilution—1:1000 |
| TFR | Transferrin receptor monoclonal antibody (Thermo Fisher Scientific, Cat# 13-6800: RRID: AB_2533029), WB Dilution—1:1000 | Rabbit polyclonal Secondary to Mouse IgG—HRP (Abcam, Cat#ab6728 RRID: 955_440) WB dilution—1:1000 |
| GAPDH | Rabbit polyclonal to GAPDH (Abcam, Cat#ab5485: RRID:AB_302725), WB dilution 1:2500 or 1:10,000 | |
b. bEnd.3 cells (passages 17–25), isolated from the SV129 strain of mice and transformed with the Polyoma virus middle T-antigen, were purchased from ATCC® and underwent authentication tests during the accessioning process (CRL-2299: RRID:CVCL0170) [36]. They were grown in T-75 flasks (Fisher Scientific, cat# 15350591) using high glucose Dulbeccos Modified Eagles Medium (Sigma-Aldrich, UK, cat# D6429) supplemented with 10% FBS (vol/vol) and 1% penicillin-streptomycin (vol/vol: Fisher Scientific cat#10003927).

Both lines were maintained at 37 °C/5% CO2 in an incubator with saturated humidity. Medium was changed every 2–3 days. Cells were split when they reached 80–90% confluency and were seeded onto 96-well plates (ThermoScientific, UK) at 20,000 cells/cm2 for b.End3 cells and 25,000 cells/cm2 for hCMEC/D3 cells. The hCMEC/D3 96-well plates were pre-coated with 0.1 mg/ml rat tail collagen type1 (Gibco cat# A1048301).

Both lines were confluent in 4–5 days and were left for another 4 days to allow for further differentiation before experimentation. The medium was changed every 2–3 days.

**Drug accumulation assay**

Accumulation assays were performed on confluent cell monolayers grown in the centre 60 wells of 96-well plates. Each passage was regarded as one ‘n’. No sample calculation was performed. The accumulation buffer (pH7.4) composition was 135 mM NaCl, 10 mM HEPES, 5.4 mM KCl, 1.5 mM CaCl$_2$, 1.2 mM MgCl$_2$, and 1.1 mM d-glucose and water. It also contained 3.7–7.7 nM [$^{3}$H]haloperidol and 3.8 μM [$^{14}$C]sucrose. [$^{14}$C]sucrose is a similar size to the test molecules and was used as an inert marker of extracellular space and membrane integrity. After the exposure period, buffer was aspirated and the wells washed with ice-cold PBS$^+$ (Sigma-Aldrich, UK) to remove drug that was not taken up by cells and to stop further transport. 1% Triton X-100 (Sigma-Aldrich) was added and the plate was incubated for an hour at 37 °C to lyse the cells and to release accumulated [$^{3}$H]drug. 100 μl from each of the wells was transferred to a vial and scintillation fluid (4 ml) added (Optiphase Hisafe 2, PerkinElmer, UK). Radioactivity was measured using a Packard Tri-Carb 2900TR liquid scintillation counter (PerkinElmer, UK) and corrected for background. The remaining 100 μl in each well were used to perform a bicinchoninic acid (BCA) protein assay. A range of 2–30 μl mg$^{-1}$ of protein was acceptable. All data for [$^{3}$H]amisulpride or [$^{3}$H]haloperidol were expressed as a volume of distribution (V$_d$) after correction for [$^{14}$C]sucrose. The V$_d$ was calculated from the sum of accumulated radioactivity (a sum of efflux and influx of the molecule) (disintegrations per minute (dpm)/mg protein) over the ratio of dpm/μl of accumulation buffer. Outliers were identified by examining the [$^{14}$C]sucrose values. [$^{14}$C]sucrose values are presented in the figures and tables.

**Transporter inhibition assay**

Transporter interaction was examined by incubating [$^{3}$H]amisulpride and [$^{14}$C]sucrose with potential inhibitors. These included unlabelled amisulpride and inhibitors/substrates of OCT, OCTN, MATE, P-gp, BCRP, and MRP at established concentrations (Additional file 1: Table S1). The MATE1 inhibitor, fomitidine, was also utilized at a higher concentration of 2 μM. ATP depletion (Promega Enliten assay) was carried out by pre-incubation with the glycolysis inhibitor, 10 mM 2-deoxy-d-glucose, for an hour before incubating the cells with [$^{3}$H]amisulpride and [$^{14}$C]sucrose. This assay had previously been shown to inhibit drug efflux by our group [32, 34].

**Cytotoxicity assay**

Cytotoxicity of amisulpride (0.1–20 μM) and eflornithine (250–500 μM) on both cell lines was assessed using 3-(4,5-dimethylthiazol-2-yl)-2,5-diphenyltetrazolium bromide (MTT) assay [32]. The results were expressed as a percentage of cell viability. This assay was also utilized on hCMEC/D3 cells (passage 32 and/or 33) for 2 h to assess toxicity of 1 μM fomitidine, 2 μM fomitidine, 2 μM lopinavir, 20 μM ergothioneine, 5 μM l-carnitine, 3 μM nifekalant hydrochloride and 1.5 μM [$^{14}$C]sucrose.

We have already published MTT assay results for eflornithine (250 μM) in hCMEC/D3 cells and haloperidol (40 μM), dexamethasone (200 μM), pentamidine (10 μM), ko143 (1 μM), MK571 (10 μM), amantadine (500 μM), corticosterone (50 μM), phophorhod A (1 μM) and prazosin (100 μM) in both cell lines [32, 34]. No significant effect was observed except with prazosin on b.End3 cells.

**Lipophilicity**

Lipophilicity is a standard physicochemical measure which is very important in terms of understanding drug distribution across membranes including the BBB and can be expressed in the form of an octanol-saline partition coefficient. The higher the lipophilicity the greater the ability of the molecule to cross the plasma membrane by passive diffusion. An octanol-saline partition coefficient for [$^{3}$H]amisulpride and [$^{3}$H]haloperidol was determined [37].
In silico computational study

Using in silico molecular docking, we tested the molecular level interactions of amisulpride, haloperidol and prazosin (OCT1 and OCT3 substrate) with the transporters OCT1, PMAT and MATE1. Due to the unavailability of the crystal structures of these transporters, molecular models of OCT1, PMAT and MATE1 were developed using homology modelling with Swiss-model webserver using PDB codes 4PPY, 5Y50 and 4ZOW, respectively, as templates (Additional file 1). A further molecular docking study was performed to explore the interaction of amisulpride, dexamethasone (P-gp substrate) and colchicine (P-gp substrate) in the binding site of the multidrug transporter ABCB1 (P-glycoprotein), and the pdb code 6FN1 [38] was used as the template. Molecular docking was performed using Dock Ligands (CDOCKER) protocol from Discovery studio version 4.0. CDOCKER is an implementation of a CHARMM based docking tool where each orientation is subjected to simulated annealing molecular dynamics. The binding sites were chosen after comparing the docking results of the ligands in all possible binding cavities within the transporters.

Animal model of AD

All experiments were performed in accordance with the Animal Scientific Procedures Act (1986) and Amendment Regulations 2012 and with consideration to the ARRIVE guidelines. The study was approved by the King’s College London Animal Welfare and Ethical Review Body and performed under license: 70/7755. A total of 40 mice were used for this study. Mice were housed at King’s College London. C57BL6/129 mice (wild-type) and the triple transgenic AD model (3×TgAD) of C57 mice which harbour the human APP-Swedish mutation (KM670/671NL), tau mutation (P301L), and presenilin-1 mutation (M146V) were utilized at 12–13 (mid-age), 16 (old) or 24 (elderly) months old. The 3×TgAD is an established AD model displaying the temporal and spatial progression and mirroring the neuropathological development seen in AD [39]. These mice were bred and genotyped at King’s College London and were a gift. They had originally been supplied by the Jackson Laboratory (RRID:IMSR_JAX:008880). No sample size calculation was performed and the study was not pre-registered. Previous experience suggested that even with the limited number of animals available there would still be sufficient power (95%) to detect significant differences (p < 0.05) [40]. Welfare was assessed daily by animal technologists. Animals were identified by earmarks and housed together by age and genotype in guideline compliant cages. All animals were maintained under standard temperature/lighting conditions and given food and water ad libitum. The experiment on each animal had to be performed within set time frames to allow the three age groups to be achieved. Experiments were performed between 9am and 5 pm. The experimenter was not blinded.

The median life span of 3×TgAD mice has been reported to be 673 days (22 months) which is shorter than the 907 day (30 months) lifespan of C57BL/6J [41]. Adult male BALB/c mice (inbred strain) were purchased from Harlan UK Limited (Oxon, UK). All mice were anaesthetised (2 mg/kg i.p. medetomidine hydrochloride and 150 mg/kg i.p. ketamine) and heparinized (100 U i.p.) in a procedure room separate to the laboratory where the brain perfusion was performed. Advice was sought from the named veterinary surgeon regarding the anaesthetic.

In situ brain perfusion

To assess differences in [3H]amisulpride and [14C]sucrose transport into the brain in ageing and in AD, wildtype (C57BL6/129 n = 17) and transgenic AD (3×Tg n = 6) mice were used. Perfusion (10 min; 5 ml/min) with a warmed (37 °C) and oxygenated (95% O2; 5% CO2) artificial plasma was via a cannula in the left ventricle of the heart as previously described [42]. The artificial plasma consisted of a modified Krebs–Henseleit mammalian Ringer solution with the following constituents: 117 mM NaCl, 4.7 mM KCl, 2.5 mM CaCl2, 1.2 mM MgSO4, 24.8 mM NaHCO3, 1.2 mM KH2PO4, 10 mM glucose, and 1 g/l bovine serum albumin. With the start of perfusion the right atrium of the heart was sectioned to prevent the recirculation of the artificial plasma. At the end of perfusion the animal was decapitated and the brain removed. To determine the brain concentration of [3H] amisulpride, frontal and occipital cortex samples were taken and weighed using a Leica S4E microscope (Purchased 2005). Brain samples were then solubilized with 0.5 ml of Solvable (PerkinElmer, USA) and liquid scintillation fluid (3.5 ml; Lumasar; PerkinElmer) added. Radioactivity in the samples were determined using the Tri-Carb2900TR scintillation counter (Purchased 2007).

Expression of results

Radioactive concentrations in the brain samples (dpm/g) were expressed as a percentage of that in the artificial plasma (dpm/ml) and termed RTISSUE (millilitres/100 g). All RTISSUE values for [3H]amisulpride were corrected for vascular/extracellular space by subtracting the [14C] sucrose RTISSUE value. Examination of the [14C]sucrose values (i.e. vascular space) and comparison to previously published values determines if the result is an outlier. This is acceptable for the WT animals, however, it is noted that [14C]sucrose (vascular space) may be affected in AD. No outliers were detected in this study.
hCMEC/D3 and b.End3 monolayers isolation

In order to perform Western blots, cell lines were grown to confluence in T-75 flasks (Thermo Scientific, UK) and left for 3–4 days, as previously described. The flask was then transferred to ice and the medium removed, before the cells were washed twice using ice-cold PBS+. Then, 1 ml of ice-cold Radio-Immunoprecipitation Assay (RIPA) buffer (Sigma-Aldrich, Dorset, UK) with added protease inhibitors (10% v/v) (Thermo Scientific, Loughborough, UK) was added to the flask to lyse the cells. A plastic cell scraper (Greiner Bio-One Ltd, Gloucestershire, UK) was used to scrape the cells off the bottom of the flask and the cell lysate was transferred to a pre-cooled 1.5 ml Eppendorf tube which was left on ice for 20 min. The tubes were then centrifuged at 10,000 rpm for 10 min at 4 °C using a Thermo Electron Corporation Heraeus Fresco17 bench-top micro-centrifuge. After centrifugation, the supernatant was transferred to another pre-cooled 1.5 ml Eppendorf and the pellet discarded. The resulting supernatant was taken for Western blot analysis.

Mouse brain capillary isolation

Brain capillaries from old-age (16 months) wild-type (3 females) and 3×TgAD (3 males, 1 female) mice were used to explore MATE1 expression. Brain capillaries were also isolated from elderly (24 month) age-matched wild-type C57BL6/129 mice (3 males, 2 females) and 3×TgAD mice (3 males, 2 females) for all other transporter studies. The left ventricle of the heart was cannulated and perfused (5 ml/min) with an oxygenated artificial plasma (modified Krebs–Henseleit mammalian Ringer) for up to 2 min. The right atrium was sectioned before perfusion was started. The mice were then decapitated and the perfused brain removed. The brain was homogenized in physiological buffer (brain weight × 3) and 26% dextran (brain weight × 4). The homogenate was subjected to density gradient centrifugation (5400×g for 15 min at 4 °C) to give an endothelial cell-enriched pellet and the supernatant was discarded [42]. 300 µl of ice-cold RIPA: ThermoFisher Scientific cat#89900) buffer with added protease inhibitors was added to the pellet at 4 °C to lyse the tissue and then centrifuged at 8000×g for 15 min at 4 °C. The resulting supernatant was taken for Western blot analysis.

Human tissue

Human tissue was provided with informed consent via the brains for dementia research (BDR) and were anonymized. BDR has ethical approval granted by the national health service (NHS) health research authority (NRES Committee London-City & East, UK: REC reference: 08/H0704/128+5. IRAS project ID:120436).

Tissue was received on the basis that it will be handled, stored, used and disposed of within the terms of the Human Tissue Act 2004. Post-mortem brain capillaries from healthy individuals (Braak stage 0–II; 86.8±1.5 years; 2 females, 3 males) and AD cases (Braak stage V–VI; 79.4±3.7 years; 2 females, 3 males) were used to investigate the expression of transporters (Case details—Additional file 1: Table S2). Medication history of the cases was supplied by the Manchester Brain Bank (Additional file 1: Table S3). In this study we identified those drugs prescribed as sedatives, antidepressants and antipsychotics.

Human brain microvasculature isolation

Brain capillaries from frontal cortex, caudate nucleus, and putamen samples were isolated after homogenising 300 mg tissue and carrying out a dextran-based density-gradient centrifugation to produce a capillary-enriched pellet.

The pellet was further lysed with 500 µl of ice-cold RIPA buffer with added protease inhibitors at 4 °C and then centrifuged at 8000×g for 15 min at 4 °C. The resulting supernatant was taken for Western blot analysis to examine transporter expression. The presence of transferrin receptor in the supernatant indicated that the method generated samples containing capillary endothelial cells.

Western Blot procedure

The supernatant protein concentration was determined using a BCA assay (Albumin standard, ThermoScientific). The supernatants were diluted and boiled for 5 min at 95 °C in 5× Laemmli sample buffer. Cell lines (30 µg except for MATE 1 antibody in Bend.3 cells where 15 µg was utilized and PMAT antibody in hCMEC/D3 and bEnd.3 cells where 20 µg and 10 µg was utilized respectively), mouse samples (15 µg for MATE1, OCTN1 and 2) and (30 µg for MATE2, PMAT and OCT1), human samples (10 µg for OCNT1 and 2) or 15–20 µg (for MATE1, MATE2, PMAT and OCT1) were loaded equally on 4–20% Mini-PROTEAN® TGX™ gels (Bio-Rad) alongside a molecular weight marker (Precision plus protein, Bio-Rad). Samples underwent SDS-PAGE at 160 V for 1 h. Proteins were transferred onto 0.45 µm polyvinylidene fluoride membranes (GE Healthcare, UK) after methanol activation at 100 V for 1 h. Membranes were blocked to reduce nonspecific binding using 5% milk with PBS-TWEEN® tablets (PBS-T) (Calbiochem, USA) at room temperature (RT) for 1 h. Membranes were incubated overnight at 4 °C with primary antibodies in PBS-T (Table 1). Membranes were washed in PBS-T (3 × 10 min) and incubated with the secondary antibody in PBS-T at RT for 1 h. Further washing in PBS-T (3 × 10...
membranes were then incubated with enhanced chemiluminescent reagent (ThermoScientific) for 30 s at RT.

Quantification of protein expression was determined by calculating the intensity ratio of the band of interest and the band of the loading control (tubulin or GAPDH). Band intensity ratio analysis was conducted using ImageJ software (NIH). Our group have previously published results from Pgp, BCRP, OCT1, OCT2 and OCT3 protein expression studies of bEnd.3 and hCMEC/D3 cells [32, 34].

Data analysis
Data are expressed as mean ± SEM. The data was analysed by two-way ANOVA with Holm-Sidak post hoc test for the accumulation studies and in situ perfusion studies, one-way ANOVA with Tukey’s post hoc test for MTT assay and Student’s t-test or two-way ANOVA for Western blot data using Sigmaplot version 13 (Systat, USA) or GraphPad Prism 7.03. p < 0.05 were considered as statistically significant. Exact p-values are provided in figure legends/"Results" section.

Results
Amisulpride accumulation and saturable transport
[^3]H]Amisulpride was able to accumulate in both hCMEC/D3 and bEnd3 cell lines to a greater extent than the baseline marker, [^14]C]sucrose (Fig. 2 and Additional file 1: Figure S1). Incubation of hCMEC/D3 and bEnd3 cell lines with 20 µM amisulpride significantly decreased the accumulation of[^3]H]amisulpride (6.5 nM) at 5 min—with a significant decrease of 37% in hCMEC/D3 cells and a decrease of 50% in bEnd.3 cells observed after 2 h (Fig. 2). No significant differences were observed for[^14]C]sucrose between the treatments, except in the bEnd3 cells at 120 min where the presence of 20 µM amisulpride decreased the accumulation of[^14]C]sucrose (Additional file 1: Figure S1). However, all[^14]C]sucrose values were within the expected range for these in vitro models. Studies also revealed that lower concentrations of unlabelled amisulpride (0.1 µM) did not affect accumulation of[^3]H]amisulpride in hCMEC/D3 (n = 5 passages) or bEnd.3 (n = 3 passages) cells at all time points (data not shown).

ABC transporter involvement
ATP depletion did not affect the accumulation of[^3]H]amisulpride or[^14]C]sucrose in either cell line (Additional file 1: Figure S2). The P-gp substrate, dexamethasone, BCRP substrate, ko143, and inhibitor, phoebophorbide A, and MRP family inhibitor, MK571, did not affect the accumulation of[^3]H]amisulpride or[^14]C]sucrose in either cell line (Additional file 1: Figure S3).

OCT, OCTN, PMAT and MATE involvement
Involvement of OCTs in[^3]H]amisulpride uptake was investigated by incubating hCMEC/D3 and bEnd.3 cells with the, OCT1 and 2 substrate, amantadine, the OCT1 and 3 substrate, prazosin, and the OCT3 substrate, corticosterone (Additional file 1: Figure S4).[^3]H]amisulpride accumulation did not change in the presence of amantadine in hCMEC/D3 cells, but significantly increased by 84% in bEnd.3 cells. In the presence of prazosin, there was a significantly reduced accumulation of[^3]H]amisulpride in hCMEC/D3 cells, but not in bEnd.3 cells.
cells. Corticosterone did not affect the accumulation of \([^{3}H]\)amisulpride in either cell line. No differences were found for \([^{14}C]\)sucrose between the treatments (Additional file 1: Figure S4).

\([^{3}H]\)amisulpride accumulation in hCMEC/D3 and b.End3 was unaffected by the presence of ergothioneine (OCTN1) and \(L\)-carnitine (OCTN2) respectively (Additional file 1: Figures S5 and S6). \([^{14}C]\)Sucrose \(V_d\) was not significantly different between the treatments, except significant differences were observed between \([^{14}C]\)sucrose and \(L\)-carnitine at 2 h suggestive that this time point for \([^{3}H]\)amisulpride should be ignored.

Incubation with the PMAT inhibitor, lopinavir, resulted in a significant increase in the \(V_d\) of \([^{3}H]\)amisulpride by 63.9% at 20 min, 83.2% at 30 min, 85.1% at 60 min and by 68.6% at 120 min (Fig. 3). The \(V_d\) of \([^{14}C]\)sucrose was not significantly different between control and test groups (Fig. 3). No affect was observed with this inhibitor in b.End3 cells (Additional file 1: Figure S7).

Incubation with the MATE1 inhibitor, famotidine (1 μM) resulted in no significant effect on the \(V_d\) of \([^{3}H]\)amisulpride in hCMEC/D3 cells (Fig. 4). The \(V_d\) of \([^{14}C]\)sucrose was not significantly different between control and test groups with (1 μM) famotidine during the standard 2 h incubation period. An assessment of the effect of 2 μM famotidine on \([^{14}C]\)sucrose alone revealed a loss of hCMEC/D3 integrity at 2 h. Further assessment of famotidine (2 μM) did not affect either \([^{3}H]\)amisulpride or \([^{14}C]\)sucrose accumulation in hCMEC/D3 cells over a 1 h period (Fig. 4). No affect was observed with famotidine (1 μM) in b.End3 with either \([^{3}H]\)amisulpride or \([^{14}C]\)sucrose (Additional file 1: Figure S7).

Interaction with the positively charged anti-psychotic drug, haloperidol

The effect of the cationic drug, haloperidol (40 μM), on radiolabelled amisulpride accumulation was also investigated. Incubation of unlabelled haloperidol with \([^{3}H]\)amisulpride did not yield any significant effects in either cell line (Additional file 1: Figure S9). No significant differences were found for \([^{14}C]\)sucrose between the treatments (Additional file 1: Figure S9).

Characteristics of haloperidol accumulation in hCMEC/D3 and b.End3 cell lines

hCMEC/D3 and b.End3 cells were incubated with unlabelled haloperidol (40 μM) along with \([^{3}H]\)haloperidol (10 nM). Incubation with unlabeled haloperidol significantly decreased the accumulation of radiolabelled haloperidol by approximately 93% in hCMEC/D3 cell line and by 94% in bEnd.3 cell line at all times (**p < 0.001) (Additional file 1: Figure S10). No significant differences were

![Graph](image-url)
found for $^{[14]}$C$sucrose$ between the treatments (Additional file 1: Figure S10).

ATP was depleted from both cell lines to determine the role of ABC transporters in the efflux of haloperidol. ATP depletion did not affect the accumulation of haloperidol in either cell line (Additional file 1: Figure S11). No significant differences were observed for $^{[14]}$C$sucrose$ between the treatments (Additional file 1: Figure S11). The hypothesis that haloperidol uptake is by OCT transporters was investigated by incubating the cells with OCT1 and 2 substrate amantadine (500 μM) and OCT1 and 3 substrate prazosin (100 μM). $^{[3]}$H$haloperidol$ accumulation significantly decreased in the presence of amantadine in both cell lines compared to control—by 89% in hCMEC/D3 cells and by 82% in bEnd.3 cells and in the presence of prazosin—by 85% in hCMEC/D3 cells and by 82% in bEnd.3 cells (**p < 0.001) (Additional file 1: Figure S12). No significant differences were found for $^{[14]}$C$sucrose$ between the treatments (Additional file 1: Figure S12).

The effects of other cationic drugs—unlabeled pentamidine (100 μM), unlabeled efornithine (250 μM) and unlabeled amisulpride (20 μM) on radiolabelled haloperidol accumulation in hCMEC/D3 was investigated. Unlabelled pentamidine significantly reduced the accumulation of radiolabelled haloperidol in the cell lines after 2 h—by 31% in hCMEC/D3 cells (**p < 0.01, and *p < 0.05). Unlabelled efornithine significantly
decreased the accumulation of radiolabelled haloperidol by 11% in hCMEC/D3 cells (**p < 0.001). Unlabelled amisulpride (20 μM) significantly decreased the accumulation of radiolabelled haloperidol in hCMEC/D3 cells—by 27% after 2 h (**p < 0.001) (Additional file 1: Figure S13). No significant differences were found for [14C]sucrose between the treatments (Additional file 1: Figure S13).

Cytotoxicity
No cytotoxic effects of amisulpride (0.1–20 μM), 1 μM famotidine, 2 μM famotidine, 2 μM lopinavir, 20 μM ergothioneine, 5 μM L-carnitine, 3 μM nifekalant hydrochloride and 1.5 μM [14C]sucrose (Additional file 1: Figure S14A and B) and efllornithine (250–500 μM) were detected using the MTT assay (data not shown). A marker molecule ([14C]sucrose) of extracellular/vascular space was included in all [3H]amisulpride accumulation experiments and ensured that any measured effect on [3H]amisulpride values could be interpreted correctly and was not simply due to loss of membrane integrity caused by the cytotoxic nature of the drugs/inhibitors utilized.

Lipophilicity
The octanol-saline partition coefficient for [3H]amisulpride was determined to be 0.0422 ± 0.0045 and for [3H]haloperidol was determined to be 0.6678 ± 0.1278.

Molecular docking studies with the SLC transporters—OCT1, MATE1 and PMAT
Amisulpride showed molecular interactions inside the binding site of OCT1 in the form of hydrogen bonds with amino acids Gln283, Asn288, Glu380 and Asn415 as well as hydrophobic interactions with amino acid residues Phe26 and Pro385 (Fig. 5a), while haloperidol showed hydrogen bonds with amino acids Gln 283, Gly384,

![Fig. 5](image-url) Molecular-level interactions of amisulpride within the binding site of OCT1 (a), MATE1 (b) and PMAT (c). Amisulpride is represented in stick-representation and amino acid residues in line-representations. Hydrogen bonds are represented in green dotted lines, and hydrophobic interactions are represented in pink dotted lines.
Trp388 and hydrophobic interactions with Gly384 and Pro385 while prazosin interacted with Asn288, Thr321, Ser324, Glu380, Asn411 through hydrogen bonds and hydrophobic interactions with Leu325, Val328, Gly384 and Pro385 (Additional file 1: Figure S15A). The best pose of amisulpride interacted with the binding pocket of OCT1 with a free energy of binding of $-14.28 \text{ kcal/mol}$ while the free energy of binding for haloperidol and prazosin were $-29.97 \text{ kcal/mol}$ and $-27.57 \text{ kcal/mol}$.

Amisulpride showed a similar level of interaction with MATE1 transporter with a free energy of binding of $-14.32 \text{ kcal/mol}$. It fit snugly within the binding pocket (Fig. 5b) and formed hydrogen bonds with amino acids Tyr45, Ser294 and Gln301 as well as hydrophobic interactions with amino acid residues Met44, Phe78 and Tyr160. The hydrophobic interactions appeared to play an important role in its interaction with MATE1 compared to its interaction with OCT1. The interaction of amisulpride was relatively weaker with PMAT compared to both OCT1 and MATE1 with free energy of binding $-11.4 \text{ kcal/mol}$. It formed a single hydrogen bond with Asn331 and interacted with hydrophobic interactions with amino acid residues Leu62, Leu236 and Ile239 through hydrophobic interactions (Fig. 5c).

A similar molecular docking study suggested haloperidol is a better substrate of both MATE1 and PMAT compared to amisulpride as it showed binding affinity of binding $-22.27 \text{ kcal/mol}$ and $-21.72 \text{ kcal/mol}$ for MATE1 and PMAT, respectively, which are notably higher than amisulpride (Fig. 5 and Additional file 1: Figure S16). It showed good interaction with MATE1 with hydrogen bonds with Tyr45 and Ser74 and hydrophobic interaction with Phe82 and Ala67 (Additional file 1: Figure S16A). However, the interaction of haloperidol with PMAT was limited to single hydrogen bond with Asp34 and hydrophobic interaction with Leu62 (Additional file 1: Figure S16B).

Molecular docking studies with the ABC transporter—P-gp

The molecular docking study revealed that amisulpride was not a substrate for P-gp with a free energy binding of $-1.81 \text{ kcal/mol}$ and the molecule was not able to interact favourably with the binding pocket of P-gp. P-gp substrates, dexamethasone and colchicine, showed notably superior interaction with P-gp with free energy of binding values of $-31.83 \text{ kcal/mol}$ and $-16.07 \text{ kcal/mol}$. Both dexamethasone and colchicine interacted with the binding pocket employing hydrogen bonds and hydrophobic interactions (Additional file 1: Figure S17 and Table S4).

CNS amisulpride delivery in vivo

The $RTISSUE$ values for $[^3H]$amisulpride did not differ from the $RTISSUE$ values for $[^{14}C]$sucrose in wildtype mice in all the age groups tested (Additional file 1: Table S5). There was also no effect of ageing on the brain distribution of $[^3H]$amisulpride or $[^{14}C]$sucrose. No differences were observed in the $[^{14}C]$sucrose $RTISSUE$ values in the frontal and occipital cortex between the wildtype and transgenic mice (Fig. 6). However, in transgenic mice the $RTISSUE$ value for $[^3H]$amisulpride was significantly higher than $[^{14}C]$sucrose in the frontal cortex, but not the occipital cortex. Importantly, in transgenic mice the sucrose-corrected $RTISSUE$ value for $[^3H]$amisulpride was significantly higher than that of wildtype mice in the frontal cortex, but not the occipital cortex (Fig. 6).

Fig. 6  The uptake of $[^3H]$amisulpride was determined in wildtype and 3× transgenic AD mice. Significant differences were observed for $[^3H]$ amisulpride between wildtype ($n=5$ frontal cortex and $n=6$ occipital cortex) and transgenic mice ($n=4$ each region)—**$p < 0.005$. $[^3H]$ Amisulpride data have been corrected for $[^{14}C]$sucrose. $[^{14}C]$Sucrose uptake is shown. No differences in paracellular permeability and membrane integrity were observed. All data are expressed as mean ± S.E.M, $n=4$–6 mice, 2 years old. Perfusion time was 10 min. 6 C57BL6/129 mice (3 males and 3 females: weight 37.0 ± 1.8 g) and 4 transgenic (2 males, 2 females: weight 29.1 ± 1.0 g) were used. Also see Additional file 1: Table S5.
Endothelial transporter expression

**Cell lines**

OCTN1, OCTN2, MATE1 and MATE2 expression was confirmed in hCMEC/D3 (passages 28 and 33) and bEnd.3 (passages 18, 19 and 23) cells (Additional file 1: Figures S18 and S19A). PMAT was expressed in hCMEC/D3 cells (passages 28, 31 and 32) and bEnd.3 (passages 17, 18, 20 and 24) (Additional file 1: Figure S19B and C).

**Wildtype and transgenic AD mice**

The total protein concentration measured in wildtype mice (188.2 ± 12.8 μg/100 μl) was not significantly different to that measured in 3×Tg AD mice (195.6 ± 13.3 μg/100 μl) brains, but this may be attributed to the fact that it was not possible to assess regional differences in these small samples. Although slight variability was observed, there was no significant differences in individual transporter (P-gp, OCT1, OCT2, OCT3, OCTN1, OCTN2, MATE1, MATE2 and PMAT) expression between the wildtype and 3×TgAD mice (Additional file 1: Figures S20 and S21). Note mice were 24 months old except for MATE1 and PMAT studies where the mice were 16 and 12 months old. Data not shown for P-gp, OCT1, OCT2, OCT3 and PMAT. The expression of all these BBB transporters suggests that a capillary enriched sample had been assessed.

**Human brain**

Human brain capillaries were isolated from the frontal cortex (for comparison with in situ perfusion experiments), caudate nucleus and the putamen (forming the striatum where high D2 and D3 receptor occupancy is observed in individuals with AD with amisulpride usage) of healthy controls and age-matched AD affected individuals (Additional file 1: Tables S2 and S3). The total protein concentration in the capillaries was found to be significantly lower in the caudate nucleus (by 37.4%) and putamen (by 32.5%), but not the frontal cortex samples, from individuals with AD compared to healthy controls (Additional file 1: Table S6).

Individual transporter expression in each brain region between AD and healthy cases is comparable as the same amount of protein has been loaded into each well. Note this amount was dependent on the antibody utilized so was variable. Transporter expression in the frontal cortex was less variable between healthy and AD cases than in the other regions studied with no significant differences in transporter expression being observed (Additional file 1: Figures S22–S28; Fig. 7). Expression of OCT1 did not change between control and individuals with AD in all the regions tested (Additional file 1: Figures S22, S23). PMAT and MATE1 expression was significantly lower in individuals with AD in the caudate nucleus (56.2%; p < 0.05) and putamen (74.8%; p < 0.05) samples, respectively, compared to control (Fig. 7; Student’s t-test). No other significant differences were observed, however, further cases are required to explore this more fully. Please note it is also likely that transporter expression in caudate nucleus and putamen AD samples is even lower than the healthy controls shown here (Fig. 7; Additional file 1: Figures S22–S28) as total protein expression is significantly reduced (Additional file 1: Table S6). The expression of all these BBB transporters would also indicate that a capillary enriched sample had been assessed.

**Medication history of the cases**

Additional file 1: Table S3 shows the medication history of the cases that have been used in this study. Sedatives, antidepressants and antipsychotic drugs (including haloperidol) were identified and listed together. A separate column lists all other medications.

**Discussion**

This study aimed to answer an important clinical question using an integrative approach to investigate the interaction between two drugs and BBB transporters and their potential pharmacodynamic relevance in AD. The in vitro cell culture models and in silico computational model allowed us to [1] identify the transporters, [2] assess the potential mechanisms of amisulpride and haloperidol transport [3] perform studies on human and mouse brain endothelium and [4] minimize animal studies in line with the 3Rs (replacement, refinement and reduction principles). The in situ brain perfusion technique allowed us to [1] study the whole animal [2] and utilize a mouse model of AD. Importantly this is the only model to exhibit both amyloid-β40 and 42 and tau pathology, mimicking human AD [39, 43]. Both plaque and tangle pathology are mainly restricted to the hippocampus, amygdala and cerebral cortex. Transporter expression in capillaries isolated from AD and age-matched human cases and mouse brain were also assessed. Medication history of the cases was evaluated.

Cell culture studies revealed a slow accumulation of [3H]amisulpride indicating a low BBB permeability. This can be linked to its low lipophilicity, as measured by the octanol-saline partition coefficient, and its inability to interact strongly with neutral and negatively charged lipid model systems thus its limited ability to passively diffuse across the lipid bilayer [22, 44]. Please note the plasma half-life of a single oral dose of amisulpride (50 or 200 mg) is ~12 h, which suggests that amisulpride will not have significantly degraded within the 2 h incubation period used in our study [45]. Furthermore the accumulation buffer does not contain plasma enzymes which would further increase the
half-life of amisulpride in our cell culture models. Unlabelled amisulpride (20 µM) significantly reduced 
\(^{3}H\)amisulpride accumulation in both cell lines, suggesting that there is a low affinity influx transporter at 
the plasma membrane or possibly at an intracellular 
organelle membrane (e.g. lysozyme) [46]. OCT1, OCT2 
and 3 have been shown to be expressed in hCMEC/
D3 and b.End3 cells in an earlier study by our group 
[32] and OCTN1 and OCTN2 were shown to be 
expressed in both cell lines in this present study. How-
ever, a study by Ohtsuki and colleagues found that the

Fig. 7 Individual values have been plotted for the transporter expression in the capillaries of frontal cortex, caudate nucleus and caudate putamen samples from healthy and AD affected individuals. The numbers in the key indicate the MRC ID designated to each sample. Details of the samples can be found in Additional file 1: Tables S2 and S6
protein expression of OCT1, OCT2, OCT3, OCTN1 and OCTN2 were below detectable limits in hCMEC/D3 cells [47].

In line with the findings of [14] and our molecular docking studies, our accumulation studies suggested that OCT1 may be the influx transporter involved, as prazosin (substrate for OCT1 and 3) reduced [3H]amisulpride uptake in hCMEC/D3 cells after 2 h, whereas corticosterone (OCT3 substrate) had no impact on [3H]amisulpride accumulation in either cell line. In addition, OCTN1 and 2 inhibitors did not affect accumulation of [3H]amisulpride into hCMEC/D3 or bEnd.3 cells. However, in the presence of amantadine, a substrate for several transporters of organic cations (OCT1, OCT2 [48]), MATE 1, MATE 2 [49, 50] and PMAT [51]) there was no effect on hCMEC/D3 cells, and an increase in [3H]amisulpride accumulation in bEnd.3 cells after 2 h incubation, which suggests the involvement of an efflux transporter. This is consistent with previous observations [22], which found higher amisulpride transport in the basolateral to apical direction (Pe 5.2 ± 3.6 × 10⁻⁶ cm/s) compared to the apical to basolateral direction (Pe < 10⁻⁶ cm/s) in porcine brain microvessel endothelium. The differences in the effect of inhibitors on the two cell lines may be explained by amisulpride being a substrate for multiple transporters and variations in the function/expressions of OCT1, MATE and PMAT transporters possibly related to species differences [52, 53]. The absence of any effect of prazosin on [3H]amisulpride uptake in bEnd.3 cells can be explained by prazosin-associated toxicity in this cell line, which causes protein values to decrease over the course of the experiment, resulting in no net effect on V_D values [32]. It is unlikely that the efflux transporter identified in our in vitro study is an ABC transporter, as neither ATP-depletion or substrates for P-gp, BCRP, or the MRP family had an effect on [3H]amisulpride accumulation in either cell line. Another study also revealed that amisulpride did not inhibit P-gp in an in vitro efflux assay [23], although they also utilised P-gp knockout and WT mice and these in vivo distribution studies suggested it was a P-gp substrate. Interestingly they considered that this may have little therapeutic impact due to its unusual receptor profile. Our single transporter computational studies revealed that amisulpride was not a P-gp substrate, but was a substrate for the SLC efflux transporters, PMAT and MATE1. Our in vitro hCMEC/D3 inhibitor studies with lopinavir, also indicated that amisulpride could be effluxed by PMAT. PMAT is thought to use a proton gradient to drive organic cation efflux from cells [53] and we (and others) have found that PMAT protein is expressed on human and mouse brain capillaries (Fig. 7, Additional file 1: Figures S19B, C and S28) [52–55]. PMAT mRNA has also been identified on the luminal and abluminal membrane of human, mouse and rat brain endothelial cells [53]. Other studies could not detect PMAT protein in isolated human capillaries and hCMEC/D3 cells [47, 56].

Interestingly our in vitro studies with the MATE1 inhibitor, famotidine, did not support the in silico data set which indicated that amisulpride was a MATE1 substrate. It is important to highlight that the lack of MATE1 inhibitor effect may not be conclusive proof of a lack of substrate interaction with the MATE1 transporter. It may be that the transporter was not sufficiently expressed [47, 52, 55, 56], although we could detect MATE1 protein in both our two cell lines, mice and human brain capillary samples. Other considerations are that the inhibitor may need to reach a therapeutic concentration within the cell to elicit a response as has been previously observed with MATE1 inhibitors [50], the substrate and the inhibitor may bind to different binding sites, the non-specificity of the inhibitor, and that amisulpride interacts with both influx (OCT) and other efflux (PMAT) transporters making a response difficult to detect [26, 53, 57].

Haloperidol has been observed to have a high degree of dopamine receptor (D2) occupancy within the brain at very low doses suggesting that haloperidol is very efficient at crossing the BBB. This is in agreement with our in vitro BBB data. Please note that the half-life of haloperidol has been reported to range 14.5–36.7 h (or up to 1.5 days) after a single oral dose so will not have been metabolized significantly over our 2 h incubation period [58]. Haloperidol may cross the BBB by passive diffusion, as supported by the relatively high octanol-saline partition coefficient of haloperidol (0.6678 ± 0.1278) compared to amisulpride (0.0422 ± 0.0045), or may involve transporters. The use of transporters by haloperidol is confirmed by the self-inhibition studies in both hCMEC/D3 and b.End3 cell lines. As haloperidol exists predominately (94.8%) as a positively charged drug at physiological pH (pKa is 8.66) the transporter is likely to be OCT, which is expressed at the BBB. This was confirmed in vitro using the OCT substrates, amantadine (OCT1 and 2) and prazosin (OCT1 and 3). We also investigated the involvement of ABC transporters in the transport of haloperidol. For this, ATP was depleted from the cells by incubating them with 10 mM 2-deoxy-D-glucose. No effects of ATP depletion were observed compared to control in either cell line suggesting that haloperidol is not a substrate for the ABC transporters P-gp, BCRP, or the MRP family at the BBB as previously observed [59, 60].

Radiolabelled amisulpride (6.5 nM) was also incubated with haloperidol (OCT1 substrate and P-gp inhibitor) [25, 61]. Radiolabeled amisulpride accumulation was not affected by haloperidol (40 μM) in either cell line. This may be the result of interactions with both OCT1 and
P-gp, although our inhibitor and in silico studies do not suggest amisulpride is a substrate for P-gp. Conversely when radiolabeled haloperidol (10 nM) was incubated with unlabeled amisulpride (20 μM) there was a significant decrease in accumulation. Overall these results may slightly reflect differences in the interaction of amisulpride and haloperidol with different transporter binding sites.

Further insight into the interaction of amisulpride and haloperidol with the transporters was obtained through the in silico computational studies, where amisulpride showed binding affinities towards the binding sites of the influx transporter, OCT1, as well as the efflux transporters, MATE1 and PMAT. The binding affinities of amisulpride towards the binding sites of these transporters showed comparable energies for OCT1 and MATE1 with molecular level interactions through hydrogen bonds and hydrophobic interactions with a number of amino acids within the binding pocket. The nature of the interaction of amisulpride with OCT1 is similar to that observed for haloperidol and prazosin which are known substrates for this transporter. The binding affinity and level of interaction were slightly weaker with PMAT compared to OCT1 and MATE1, but still considerable level of interactions were observed suggesting amisulpride is a weak substrate of PMAT. Haloperidol showed comparable affinity for both MATE1 and PMAT suggesting it is a substrate for both transporters. MATE1 and PMAT are known to use a proton gradient to drive organic cation efflux from cells [53, 54, 62, 63]. MATE1, like P-gp, is thought to be expressed on the luminal membrane of the BBB, although this remains to be confirmed [64].

The in silico study supports the experimental observations and provides further evidence that amisulpride might be influxed through OCT1 and effluxed through PMAT and MATE1 but not P-gp. When amisulpride transport was investigated in vivo, a low BBB permeability to [3H]amisulpride was also observed in the wild-type mice and this did not change with age. Further studies in the 3×TG AD mice revealed an increased CNS uptake which was not accounted for by altered BBB integrity or changes in vascular space in the AD model mice, as there were no differences in [14C]sucrose uptake between the two groups; and neither was it explained by non-expression of the transporters studied (P-gp, OCT1, OCT2, OCT3, OCTN1, OCTN2, MATE1, MATE2 and PMAT). However, in post-mortem human brain expression of the efflux transporter MATE1 was lower in individuals with AD compared to age-matched healthy controls in the putamen; and PMAT showed a similar trend in the caudate nucleus, but there was no change in expression levels of these transporters in the frontal cortex. Importantly it has been reported that BBB impairments stem from AD abnormality instead of from vascular comorbidities [65]. Further evidence for regional tissue changes with AD came from our total protein measures (Additional file 1: Table S4). This may be linked to changes in the expression of other transporters such as P-gp [18] and GLUT1 [66] as well as the other SLC transporters measured in this present study. Interestingly regional differences are known to exist in the BBB transport and the intracellular distribution of antipsychotics [67]. The reduced MATE1 and PMAT expression observed in AD may therefore underpin the heightened sensitivity to amisulpride observed in the clinical population especially as together our in silico and in vitro studies suggested that amisulpride was a substrate for MATE 1 and PMAT. Several medications listed in Additional file 1: Table S3 (e.g. citalopram, metoclopramide and loratadine) have previously been identified as OCT1 inhibitors [25, 26]. It is not yet known if they also interact with MATE1 and PMAT. One of the medications (ranitidine) is an inhibitor of both OCT1 and MATE [64].

**Conclusions**

This study included a detailed evaluation of transporter expression and usage at the BBB using in silico computational approaches, in vitro models and an in vivo animal model of AD as well as patient material. The datasets have provided evidence of an interaction of amisulpride and haloperidol with both influx (OCT1) and efflux (MATE1 and PMAT) transporters, which may be expressed at the luminal or abluminal membranes of the BBB and/or at an intracellular membrane. In vitro and in silico studies indicated that amisulpride was not a substrate for ABC transporters including P-gp. Furthermore, the study is of key importance as the results suggest that the heightened sensitivity to amisulpride observed in older people with AD is possibly due to previously unreported changes in SLC transporter expression, which increase amisulpride entry into, or possibly reduce clearance from the brain. This study is also the first step in the process of characterising age and AD-specific changes in SLC transporters of organic cations. Overall our study has implications beyond amisulpride prescribing, as it suggests that dose adjustments may be required for other drugs (e.g. haloperidol) which are substrates for SLC transporters in particular MATE1 and PMAT.

**Supplementary information**

Supplementary information accompanies this paper at https://doi.org/10.1186/s12987-019-0158-1.
Abbreviations

OCT: organic cation transporters; OCTN: organic cation transporters novel; hCMEC/D3: immortalized human cerebral microvascular endothelial cell line; MATEs: multi-drug and toxic compound efflux proteins; PMAT: plasma membrane monoamine transporter; 3-TgAD: triple transgenic AD model; FBS: foetal bovine serum; BBB: blood–brain barrier; AD: Alzheimer’s disease; P-gp: P-glycoprotein; BCRP: breast cancer resistance protein; MRPs: multi-drug resistance protein; RIPA: radio-immunoprecipitation assay; SLC: solute carrier; eNOS: 3’S,5’dimethylythiazol-2-yl)-2,5-diphenyltetrazolium bromide; APP: amyloid precursor protein; PBS-T: PBS-TWEEN® tablets; RT: room temperature; BDR: brains for dementia research; Vd: volume of distribution; WB: Western blot.

Acknowledgements

We would like to thank Professor I. Romero, Professor B. Weksler, and Professor P. Couraud for the hCMEC/D3 cell line provided under MTA and Dr. Manasi Nandi for advice regarding the Western blots.

Authors’ contributions

GNS co-designed the study with SAT. Performed the brain perfusions (with AF). Performed MTT assay for amisulpride and effrinithine, Western blot analysis on human tissue and the bEnd.3 and D3 cell culture studies with unlabelled amisulpride, ATP depletion, ATP-transporter and OCT inhibitors. Analysed the resulting data. Contributed to the writing of the manuscript. ALF performed the brain perfusions (with GNS). Performed Western blots for MATE1 in bEnd.3, hCMEC/D3, wild-type and 3-Tg AD mice. Analysed the resulting data. STB and RL performed the cell culture studies with MATE1, MATE2, PMAT, OCTN1 and OCTN2 inhibitors with hCMEC/D3 and bEnd.3 respectively. Analysed the resulting data. STB also performed MTT assay with ergothionine, respectively. Analysed the resulting data. STB also performed MTT assay with hCMEC/D3 and bEnd.3; mouse brain endothelial cell lines; VLOSLP: very late-onset (> 60 years) schizophrenia-like psychosis; NHS: national health service; MTI: 3,4,5-dimethylythiazol-2-yl)-2,5-diphenyltetrazolium bromide; APP: amyloid precursor protein; PBS-T: PBS-TWEEN® tablets; RT: room temperature; BDR: brains for dementia research; Vd: volume of distribution; WB: Western blot.

Funding

This research was supported by the Medical Research Council (Ph.D. student for Ms. Sekhar Ph.D. studies MR/K500811/1), the King’s College London Alzheimer’s Research Unit (ARU) network centre seed fund (to Dr. Suzanne Reeves, Dr. Sarah Thomas and Ms. Gayathri Sekhar), The Edmond and Lily Safra Foundation (Dr. Broadstock) and a multi-user equipment grant from The Wellcome Trust (Dr. Thomas (PI), Professors Francis, McMahon, Malcangio, Reeves, Dr. Sarah Thomas and Ms. Gayathri Sekhar), The Maurice Wolfson Centre for Age-Related Diseases, King’s College London, Guy’s Campus, London SE1 1UL, UK. 4 Maurice Wolfson Clinical Neuroscience Institute, King’s College London, 125 Coldharbour Lane, Camberwell, London SE5 9N, UK. 5 Division of Psychiatry, Faculty of Brain Sciences, University College London, 149 Tottenham Court Road, London WIT 7NF, UK.

Availability of data and materials

See Additional File 1.

Ethics approval and consent to participate

Human tissue was provided with informed consent via the brains for dementia research (BDR) and were anonymized. BDR has ethical approval granted by the national health service (NHS) health research authority (NRES Committee London City-East, UK, REC reference 08/H0704/128+5. IRAS project ID:120436). Tissue was received on the basis that it will be handled, stored, used and disposed of within the terms of the Human Tissue Act 2004. Specifically, the human brain tissue samples were supplied by The Manchester Brain Bank, which is part of the Brains for Dementia Research programme jointly funded by Alzheimer’s Research UK and Alzheimer’s Society (BDR Reference: Amendment to study-BDR14 TRID 170 recipient PI: Dr. Sarah Thomas).

Consent for publication

Not applicable.

Competing interests

The authors declare that they have no competing interests.

Author details

1 Faculty of Life Sciences and Medicine, School of Cancer and Pharmaceuti
cal Sciences, King’s College London, Franklin-Wilkins Building, 150 Stamford
Street, Waterloo, London SE1 9HN, UK. 2 Faculty of Pharmacy, Msr Interna
tional University, Cairo 11431, Egypt. 3 Wolfson Centre for Age-Related Dis
eases, King’s College London, Guy’s Campus, London SE1 1UL, UK. 4 Maurice
Wolfson Clinical Neuroscience Institute, King’s College London, 125 Coldharbour
Lane, Camberwell, London SE5 9N, UK. 5 Division of Psychiatry, Faculty of Brain
Sciences, University College London, 149 Tottenham Court Road, London WIT
7NF, UK.

Received: 10 October 2019 Accepted: 2 December 2019

References

1. Ballard C, Howard R. Neuroleptic drugs in dementia: benefits and harm. Nat Rev Neurosci. 2006;7(6):492–500. https://doi.org/10.1038/nrn1926.
2. Schneider LS, Tariot PN, Dagerman KS, Davis SM, Hsiao JK, Ismail MS, et al. Effectiveness of atypical antipsychotic drugs in patients with Alzheimer’s disease. N Engl J Med. 2006;355(15):1525–38. https://doi.org/10.1056/NEJMoa061240.
3. Reeves S, McLachlan E, Bertrand J, Antonio FD, Brownings S, Nair A, et al. Therapeutic window of dopamine D2/3 receptor occupancy to treat psychosis in Alzheimer’s disease. Brain. 2017;140(4):1117–27. https://doi.org/10.1093/brain/aww359.
4. Cavagaggio G, Graff-Guerrero A. Is antipsychotic sensitivity in Alzheimer’s disease secondary to abnormal blood-brain barrier integrity? Brain. 2017;140(4):865–7.
5. Mauri MC, Paletta S, Maffini M, Colasanti A, Dragogna F, Di Pace C, et al. Clinical pharmacology of atypical antipsychotics. EXCLI J. 2014;13:1163.
6. Hiemke C, Baumann P, Bergemann N, Conca A, Dietmaier O, Egberts K, et al. AGNP consensus guidelines for therapeutic drug monitoring in psychiatry: update 2011. Pharmacopsychiatry. 2011;44(6):195–235.
7. Sparkhass A, Taylor D, Patel MX, Kapur S. Amisulpride—dose, plasma concentra
tion, occupancy and response: implications for therapeutic drug monitoring. Acta Psychiatr Scand. 2009;120(6):416–28. https://doi.org/10.1111/j.1600-0447.2009.01429.x.
8. Lako IM, van den Heuvel ER, Kneutgenring B, Bruggeman M, Taxis K. Estimat
ing dopamine D2 receptor occupancy for doses of 8 antipsychotics. J Clin Psychopharmacol. 2013;33(3):675–81.
9. Schoemaker H, Claustre Y, Fage D, Rouquier L, Chergui K, Curet O, et al. Neurochemical characteristics of amisulpride, an atypical dopamine D2/3
d receptor antagonist with both presynaptic and limbic selectivity. J Pharmacol Exp Ther. 1997;280(1):83–97.
10. Clark-Papavas C, Dunn JT, Greaves S, Mogg A, Gomes R, Brownings S, et al. Towards a therapeutic window of D2/3 occupancy for treatment of psychosis in Alzheimer’s disease, with [18F]fallypride positron emission tomography. Int J Geriatr Psychiatry. 2014;29(10):1001–9.
11. Reeves S, Bertrand J, D’Antonio F, McLachlan E, Nair A, Brownings S, et al. A population approach to characterise amisulpride pharmacokinetics in older people and Alzheimer’s disease. Psychopharmacology. 2016;233(18):3371–81. https://doi.org/10.1007/s00213-016-4379-6.
12. Reeves S, Eggleston K, Cott E, McLachlan E, Brownings S, Nair A, et al. Therapeutic D2/3 receptor occupancies and response with low amisulpride blood concentrations in very late-onset schizophrenia-like psychosis (VLOSLP). Int J Geriatr Psychiatry. 2018;33(2):396–404. https://doi.org/10.1002/gps.4758.
13. Saidjami M, Karimi Dermanci F, Sohrabi S, Patching SG. Efflux proteins at the blood–brain barrier: review and bioinformatics analysis. Xenobiotica. 2017;47:1–27. https://doi.org/10.1080/00498254.2017.1328148.
14.Dos Santos Pereira JN, Tadjerpisheh S, Abu Abed M, Saadatmand AR, Weks
er B, Romero IA, et al. The poorly permeable membrane antipsychotic
drugs amisulpride and sulpiride are substrates of the organic cation transporters from the SLC22 family. AAPS J. 2014;16(6):1247–58.

15. Natesan S, Reckless GE, Barlow KBL, Nobrega JN, Kapur S. Amisulpride the ‘atypical antipsychotic analgesic’—comparison to haloperidol, risperidone and clozapine. Schizophr Res. 2008;105(1):13–24.

16. Kania KD, Wijesuriya HC, Hardiky SB, Barrand MA. Beta amyloid effects on expression of multidrug efflux transporters in brain endothelial cells. Brain Res. 2011;1418:11–11.

17. van Assema DM, Lubbenink M, Boellaard R, Schuit RC, Windhorst AD, Scheltens P. Lammertse AA, van Berckel BN. P-glycoprotein function at the blood-brain barrier: effects of age and gender. Mol Imaging Biol. 2012;14(6):771–7.

18. Vogelgesang S, Cascorbi I, Schroeder E, Pahnke J, Kroemer HK, Siegmund P, et al. Differential substrate and inhibitory activities of ranitidine and famotidine toward human organic cation transporter 1 (hOCT1; OCT1) and multiple human multidrug efflux transporters: effect of trypanosome infection, combination therapy, P-glycoprotein, and multidrug resistance-associated protein. J Pharmacol Exp Ther. 2009;329(3):967–71.

19. Rae EA, Brown RE. The problem of genotype and sex differences in life expectancy in transgenic AD mice. Neurobiol Biobehav Rev. 2015;57:238–51.

20. Sanderson L, Khan A, Thomas S. Distribution of suramin, an antitrypanosomal drug, across the blood-brain and blood-cerebrospinal fluid interfaces in wild-type and P-glycoprotein transporter-deficient mice. Antimicrob Agents Chemother. 2007;51(9):3146–46.

21. Itagaki S, Ganapathy V, Ho HTB, Zhou M, Babu E, Wang J. Electrophysiological characterization of the polyspecific organic cation transporter OCT1 on expression of multidrug efflux transporters in brain endothelial cells expressing the middle T oncogene. Cell. 1990;62(3):435–45.

22. Montesano R, Pepper MS, Möhle-Steinlein U, Risau W, Wagner EF, Orli C. Increased proteolytic activity is responsible for the aberrant morpho-geometric behavior of endothelial cells expressing the middle T oncogene. Cell. 1990;62(3):435–45.

23. Fujiwara S, Fodor M, Njemisidou D, Schmitt EM, Liao SY. Drug distribution with influx and efflux transporters expressed at the blood-brain barrier. J Neurochem. 2004;57(1):1106–11.

24. Sekhar G, Reyes S, Thomas SA. The transport of nifurtimox, an anti-trypanosomal drug, in an in vitro model of the human blood–brain barrier: evidence for involvement of breast cancer resistance protein. Brain Res. 2012;1436:111–21.

25. Sekhar G, Reyes S, Thomas SA. The human brain endothelial cell line hCMEC/D3 as a human blood-brain barrier model for drug transport studies. J Neurochem. 2008;107(5):1358–68.

26. Sekhar G, Reyes S, Thomas SA. Exploring the interaction of amisulpride with PKC delta in mouse brain capillary endothelial cells. Eur J Pharm Biopharm. 2012;87(1):30–9.

27. Sekhar G, Reyes S, Thomas SA. The transport of nifurtimox, an anti-trypanosomal drug, in an in vitro model of the human blood–brain barrier: evidence for involvement of breast cancer resistance protein. Brain Res. 2012;1436:111–21.

28. Sekhar G, Reyes S, Thomas SA. Exploring the interaction of amisulpride with PKC delta in mouse brain capillary endothelial cells. Eur J Pharm Biopharm. 2012;87(1):30–9.

29. Sekhar G, Reyes S, Thomas SA. The transport of nifurtimox, an anti-trypanosomal drug, in an in vitro model of the human blood–brain barrier: evidence for involvement of breast cancer resistance protein. Brain Res. 2012;1436:111–21.

30. Sekhar G, Reyes S, Thomas SA. Exploring the interaction of amisulpride with PKC delta in mouse brain capillary endothelial cells. Eur J Pharm Biopharm. 2012;87(1):30–9.

31. Sekhar G, Reyes S, Thomas SA. Exploring the interaction of amisulpride with PKC delta in mouse brain capillary endothelial cells. Eur J Pharm Biopharm. 2012;87(1):30–9.
and its implications to the transport of neurotoxins. J Neurochem. 2015. https://doi.org/10.1111/jnc.13383.
54. Hiasa M, Matsumoto T, Kornatsu T, Moriyama Y. Wide variety of locations for rodent MATE1, a transporter protein that mediates the final excretion step for toxic organic cations. Am J Physiol Cell Physiol. 2006;291(4):C637–46.
55. Kurosawa K, Taka Y, Higuchi K, Yamaguchi T, Nakamura T, Mochizuki T, et al. Expression and functional characterization of drug transporters in brain microvascular endothelial cells derived from human induced pluripotent stem cells. Mol Pharm. 2018;15(12):5546–55. https://doi.org/10.1021/acs.molpharmaceut.8b00369.
56. Uchida Y, Ohtsuki S, Katsukura Y, Ikeda C, Suzuki T, Kamie J, et al. Quantitative targeted absolute proteomics of human blood-barrier transporters and receptors. J Neurochem. 2013;117(2):333–45.
57. Wittwer MB, Zur AA, Khuri N, Kido Y, Kosaka A, Zhang X, et al. Discovery of potent, selective multidrug and toxin extrusion transporter 1 (MATE1, SLC47A1) inhibitors through prescription drug profiling and computational modeling. J Med Chem. 2013;56(3):781–95. https://doi.org/10.1021/jm301302s.
58. de Leon J, Diaz Fi, Wedlund P, Jostassen RC, Cooper TB, Simpson GM. Haloperidol half-life after chronic dosing. J Clin Psychopharmacol. 2004;24(6):656–60.
59. Iwaki K, Sakaida K, Nakamura M, Komoto C, Okamura N, et al. Haloperidol is an inhibitor but not substrate for MDR1/P-glycoprotein. J Pharm Pharmacol. 2006;58(12):1617–22.
60. Schinkel AH, Wagaar E, Mol CA, van Deemter L. P-glycoprotein in the blood-brain barrier of mice influences the brain penetration and pharmacological activity of many drugs. J Clin Invest. 1996;97(12):2517–24.
61. Matsson P, Pedersen JM, Norinder U, Bengtsson CÅ, Artursson P. Identification of novel specific and general inhibitors of the three major human ATP-binding cassette transporters P-gp, BCRP and MRP2 among registered drugs. Pharm Res. 2009;26(8):1816–31.
62. André P, Saubaméa B, Cochois-Guégan V, Marie-Claire C, Cattelotte J, Smirnova M, et al. Transport of biogenic amine neurotransmitters at the mouse blood-retina and blood-brain barriers by uptake1 and uptake2. J Cereb Blood Flow Metab. 2012;32(11):1989–2001.
63. Ito S, Kusuhara H, Yokochi M, Toyoshima J, Inoue K, Yuasa H, et al. Competitive inhibition of the luminal efflux by multidrug and toxin extrusions, but not basalateral uptake by organic cation transporter 2, is the likely mechanism underlying the pharmacokinetic drug-drug interactions caused by cimetidine in the kidney. J Pharmacol Exp Ther. 2012;340(2):393–403.
64. Staud F, Cervey L, Ahmadi-Moghadam D, Cekova M. Multidrug and toxin extrusion proteins (MATE/SLC47); role in pharmacokinetics. Int J Biochem Cell Biol. 2013;45(9):2007–11.
65. van de Haar HJ, Burgmans S, Jansen JFA, van Osch MJP, van Buchem MA, Muller M, et al. Blood-brain barrier leakage in patients with early Alzheimer disease. Radiology. 2016;281(2):527–35. https://doi.org/10.1148/radiol.2016152244.
66. Horwood N, Davies DC. Immunolabelling of hippocampal microvessel glucose transporter protein is reduced in Alzheimer’s disease. Virchows Arch. 1994;425(1):69–72.
67. Loryan I, Melander E, Svensson M, Payan M, König F, Jansson B, et al. In-depth neuropharmacokinetic analysis of antipsychotics based on a novel approach to estimate unbound target-site concentration in CNS regions: link to spatial receptor occupancy. Mol Psychiatry. 2016;21(11):1527–36.

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