The role of increase hBCATm in the endothelial cells of patients with Alzheimer’s Disease.

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Dedicated to the memory of Thomas William Joynes.
Rest in peace, granddad.
ABSTRACT.

Introduction and aims: The branched-chain aminotransferase (BCAT) enzymes are important in the regulation of brain L-glutamate. A unique function of these aminotransferases is their regulation by the redox environment, where our group have shown their function as oxidoreductases, and their ability to refold misfolded proteins in particular when S-glutathionylated. Our group recently showed a significant increase in the level of these proteins in Alzheimer’s disease brain. An increase in BCAT activity could generate excess L-glutamate, contributing to the excitotoxic environment observed under pathogenic conditions. Alternatively, we hypothesize that if this protein is modified in response to cellular stress, it may play a more prominent role in regulating redox status or protein folding. To address these questions this project focussed on the design of chemical inhibitors and knock-down models together with a co-culture model of the human cerebrovasculature and specifically targeted key metabolic and redox pathways.

Methods: Several chemical inhibitors based on 4-Benzoyloxyphenylacetic acid were identified using the DockBlaster and Schrödinger software suites, synthesized and structurally verified using proton nuclear magnetic resonance (NMR) spectroscopy. The functional impact and specificity of these inhibitors was assessed using the BCAT radiolabelled assay and a coupled-enzyme assay. In tandem, siRNA was used to knock-down both isoforms in SH-SY5Y cells, and validated using western blot analysis and RT-PCR. The impact of BCATm knock-down on the expression of redox proteins, in addition to selected metabolic proteins, was assessed by western blot analysis. Functional redox assays, including glutathione concentration and metabolic activity, were also used to investigate the impact of human BCAT (hBCAT) expression in neuronal cells. Finally, a model of the blood-brain barrier (BBB) was developed and validated for studies into the role of mitochondrial hBCAT (hBCATm) in brain microvasculature.

Results: For the first time we have identified a family of chemical inhibitors of hBCATm. In particular, benzofenac has a two-fold greater enzyme affinity for hBCATm (K_i=43 µM) than cytosolic hBCAT (hBCATc) (K_i=93 µM) and a four-fold greater inhibition relative to alanine transaminase (ALT; K_i=167 µM). These inhibitors will require further optimisation but have potential as tools to assess the cellular function of hBCAT. In separate studies, knock-down of hBCATm in SH-SY5Y neuronal cells demonstrated that hBCATm expression has an impact on the metabolic and redox status of the cell. In particular, knock-down caused a >70% decrease in glutaredoxin (GRx), thioredoxin (TRx), branched-chain α-keto acid dehydrogenase α-subunit (BCKDHA), and AU-rich binding homolog of enoyl-CoA hydratase (AUH) expression. Interestingly, this effect was attenuated when cells were treated with L-leucine, indicating that these mechanisms may be regulated by a metabolic signal. L-glutamate treatment was also found to significantly increase hBCATm expression, but decreased glutamate dehydrogenase (GDH) expression, except in cells overexpressing hBCATm, suggesting a metabolic synergy between the two enzymes. Finally, total glutathione concentration was significantly decreased on hBCATm knock-down, while sensitivity to L-glutamate toxicity was significantly increased.

Discussion: Results from this work have significantly contributed to the design of cellular models, which can be used to further investigate the role of BCAT in metabolic and redox metabolism. A model of the human blood-brain barrier, developed in this thesis, will also contribute to evaluating the endothelial role of hBCATm. The initial impact of limiting BCAT expression is both a decrease in the expression of key metabolic proteins and also the cellular reductants. Together this had an impact on cell survival. Knowledge of these pathways and their regulation will be important to our understanding, of not only the regulation of brain L-glutamate, but also the role of BCAT in protein folding and cellular redox status. These factors are fundamentally important to development of neurodegenerative conditions but also in tumour development such as gliomas.
ABBREVIATIONS.

3MeBOPAA – 3-methyl-4-[(4-methylbenzyl)oxy]-phenylacetic acid

3OMeBOPAA – 4-(benzyloxy)-3-hydroxy-phenylacetic acid

4EBP1 – Eukaryotic translation initiation factor 4E-binding protein 1

AAOA – Aminoxyacetic acid

Aβ – Amyloid β peptide

Aβ40 – Amyloid β peptide 1-40

Aβ42 – Amyloid β peptide 1-42

AD – Alzheimer’s disease

AJ – Adherens junctions

Akt – RAC-alpha serine/threonine-protein kinase

ALT – Alanine transaminase

AMPA – α-amino-3-hydroxy-5-methyl-4-isoxazolepropionic acid

AMPK – 5’ adenosine monophosphate-activated protein kinase

APP – Amyloid precursor protein

APS – Ammonium persulphate

ARE – Antioxidant-response element

AST – Aspartate aminotransferase

ATF4 – Activating transcription factor 4

ATP – Adenosine triphosphate

AUH – AU RNA binding protein/enoyl-CoA hydratase

BBB – Blood brain barrier

BCAA – Branched-chain amino acids

BCAT – Branched-chain aminotransferase

BCAT1 – Branched-chain aminotransferase gene (cytosolic isoform)

BCAT2 – Branched-chain aminotransferase gene (mitochondrial isoform)
BCKA – Branched-chain α-keto acid
BCKD – Branched-chain α-keto acid dehydrogenase
BCKDHA – Branched-chain α-keto acid dehydrogenase, E1α subunit
BiP – Binding immunoglobulin protein
Benzofenac – 4-(benzyloxy)-3-chloro-phenylacetic acid
BOPAA – 4-(benzyloxy)phenylacetic acid
BSA – Bovine serum albumin
CHOP – C/EBP [CCAAT/enhancer binding protein] homologous protein
cDNA – Complementary DNA
CSF – Cerebrospinal fluid
Cys – Cysteine
DCC – N,N-dicyclohexylcarbodiimide
DCM – Dichloromethane
ddDNA – Double stranded DNA
DMAP – 4-dimethylaminopyridine
DMEM – Dulbecco’s Modified Eagle’s medium
DMF – Dimethylformamide
DPM – Disintegrations per minute
DTNB – 5,5’-dithiobis-2-nitrobenzoic acid
DTT – Dithiothreitol
E-PLP – Enzyme-Pyridoxal L-phosphate complex
E-PMP – Enzyme-Pyridoxal monophosphate complex
FBS – Fetal bovine serum
EAAT1 – Excitatory amino acid transporter 1
EDTA – Ethylenediaminetetraacetic acid
eGFP – Enhanced green fluorescent protein
EGTA – Ethyleneglycoltetraacetic acid

eIF2α – Eukaryotic translation initiation factor 2 α

FT-IR – Fourier transform infrared

GABA – γ-aminobutyric acid

GAD – L-glutamate decarboxylase

GAPDH – Glyceraldehyde 3-phosphate dehydrogenase

GC – Gas chromatography

GC-MS – Gas chromatography coupled mass spectrometry

GDH – Glutamate dehydrogenase

GLUT1 – Glucose transporter 1

GPx – Glutathione peroxidase

GRx – Glutaredoxin

GS – Glutamine synthase

GSH – Glutathione (reduced)

GSK3β – Glycogen synthase kinase 3 β

GSSG – Glutathione (oxidized)

HAPAA – 2-hydroxy-4-(4-methylbenzoyl)aminophenylacetic acid

hBCAT – Human branched-chain aminotransferase

hBCATc – Human branched chain aminotransferase (cytosolic isoform)

hBCATm – Human branched-chain aminotransferase (mitochondrial isoform)

HBSS – Hank’s buffered salt solution

HEPES – 4-(2-hydroxyethyl)-1-piperazineethanesulfonic acid

HMG-CoA - 3-hydroxy-3-methylglutaryl-coenzyme A

HSPAA – 2-hydroxy-4-[[4-(methylphenyl)sulfanyl]carbonyl]phenylacetic acid

HTS – High-throughput screening

IC50 – Half maximal inhibitory concentration
IPTG – Isopropyl β-D-1-thiogalactopyranoside
IRS – Insulin receptor substrate
ISR – Integrated stress response
KA – Kainic acid
$K_{\text{cat}}$ – Substrate molecules turned over per enzyme molecule, per second
KEAP1 – Kelch-like ECH [erythroid cell-derived protein with CNC homology]-associated protein 1
$K_{\text{eq}}$ – Equilibrium constant
KG – α-Ketoglutarate
KIC – α-ketoisocaproate
KIV – α-ketoisovalerate
Klf15 – Kruppel-Like Factor 15
KMV – α-keto-β-methylvalerate
KMV - Keto-β-methylvalerate
LC-MS – Liquid chromatography coupled mass spectrometry
LDS – Lithium dodecyl sulphate
LNAAT – Large neutral amino acid transporter
MAP – Microtubule-associated protein
MAPT – Tau microtubule-associated protein gene
Mct – Monocarboxylic acid transporter
MeBenzofenac – 3-chloro-4-[(4-methylbenzyl)oxy]-phenylacetic acid
MG-CoA – 3-methylglutaconyl-CoA
MOPS – 3-[(N-morpholino)propanesulfonic acid
MS – Mass spectrometry
MSUD – Maple syrup urine disease
mTOR – Mammalian target of Rapamycin
mTOR1 – Mammalian target of Rapamycin complex 1
MTS – 3-(4,5-dimethylthiazol-2-yl)-5-(3-carboxymethoxyphenyl)-2-(4-sulfophenyl)-2H-tetrazolium

NADH – Nicotinamide adenine dinucleotide

NADPH – Nicotinamide adenine dinucleotide phosphate

Na-Flu – Sodium fluorescein

NEM – N-ethylmaleimide

NFT – Neurofibrillary tangles

NMDA – N-methyl-D-aspartic acid

NMDAR – N-methyl-D-aspartic acid receptor

NMR – Nuclear magnetic resonance (spectroscopy)

NO – Nitric oxide

Nrf2 – Nuclear factor (erythroid-derived 2)-like 2

P70S6K – Ribosomal protein S6 kinase beta-1

p-eIF2α – Phosphorylated eukaryotic translation initiation factor 2 α

P-gp – P-glycoprotein

PAG – Phosphate- activated glutaminase

PDI – Protein disulphide isomerase

Pe – Permeability coefficient

PERK – Protein kinase RNA-like endoplasmic reticulum kinase

PI3K – Phosphoinositide 3-kinase

pIC50 – negative logarithm of IC50 [Half maximal inhibitory concentration]

PLP – Pyridoxal L-phosphate

PMP – Pyridoxine monophosphate

PP18b – Placental protein 18b

PP2A – Protein phosphatase 2A

Rheb – Ras homolog enriched in brain

RNAi – RNA interference
ROS – Reactive oxygen species
RT – Retention time
SDS – Sodium dodecyl sulphate
shRNA – Short hairpin RNA
siRNA – Small interfering RNA
ssRNA – Single stranded RNA
SOD – Superoxide dismutase
TBS – Tris-buffered saline
TBST – Tris-buffered saline with Tween-20
TCA – Trichloroacetic acid
TEMED – Tetramethylethylenediamine
Tris – Hydroxymethylaminomethane
TJ – Tight junction
TOR – Target of Rapamycin
TR – Thyroid receptor
TRx – Thioredoxin
TSC1 – Tuberous sclerosis 1
U – Enzyme units
VDCC – Voltage dependent calcium channels
ZO – Zonula occludens
PRESENTATIONS AND PUBLISHED WORKS.

Oral presentation

“BCAT and the brain” Postgraduate research forum 2015, University of the West of England.

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