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Tau Monoclonal Antibody (TAU-5)

Catalog Number **AHB0042**

Product data sheet

Details	
Size	100 µg
Host/Isotope	Mouse / IgG1
Class	Monoclonal
Type	Antibody
Clone	TAU-5
Immunogen	Purified bovine microtubule-associated proteins.
Conjugate	Unconjugated
Form	Liquid
Concentration	0.5 mg/mL
Purification	purified
Storage buffer	PBS, pH 7.4
Contains	15mM sodium azide
Storage Conditions	-20°C

Species Reactivity	
Species reactivity	Bovine, Human, Mouse, Sheep, Rat
Published species	Rat, Non-human primate, Bovine, Mouse, Human, Not Applicable

Tested Applications	Dilution *
Immunohistochemistry (Frozen) (IHC (F))	1:500
Immunohistochemistry (Paraffin) (IHC (P))	1:20-1:200
Immunoprecipitation (IP)	Assay-dependent
Western Blot (WB)	1:500
Immunocytochemistry (ICC/IF)	Assay-dependent

Published Applications	
Western Blot (WB)	See 67 publications below
Dot blot (DB)	See 1 publications below
Immunoelectrophoresis (IE)	See 1 publications below
Immunoprecipitation (IP)	See 5 publications below
in situ PLA (PLA)	See 1 publications below
Immunohistochemistry (IHC)	See 8 publications below
Immunocytochemistry (ICC/IF)	See 5 publications below
Immunohistochemistry (Frozen) (IHC (F))	See 2 publications below
Miscellaneous PubMed (Misc)	See 5 publications below
ELISA (ELISA)	See 2 publications below

* Suggested working dilutions are given as a guide only. It is recommended that the user titrate the product for use in their own experiment using appropriate negative and positive controls.

Background/Target Information

Tau is a neuronal microtubule-associated protein found predominantly on axons. The function of Tau is to promote tubulin polymerization and stabilize microtubules. The C-terminus binds axonal microtubules while the N- terminus binds neural plasma membrane components, suggesting that tau functions as a linker protein between both. Axonal polarity is predetermined by TAU/MAPT localization (in the neuronal cell) in the domain of the cell body defined by the centrosome. The short isoforms allow plasticity of the cytoskeleton while the longer isoforms may preferentially play a role in its stabilization. In its hyper-phosphorylated form, Tau is the major component of paired helical filaments (PHF), the building block of neurofibrillary lesions in Alzheimer's diseases (AD) brain. Hyper-phosphorylation impairs the microtubule binding function of Tau, resulting in the destabilization of microtubules in AD brains, ultimately leading to the degeneration of the affected neurons. Numerous serine/threonine kinases phosphorylate Tau, including GSK-3beta, protein kinase A (PKA), cyclin-dependent kinase 5 (cdk5) and casein kinase II. Hyper-phosphorylated Tau is found in neurofibrillary lesions in a range and other central nervous system disorders such as Pick's disease, frontotemporal dementia, cortico-basal degeneration and progressive supranuclear palsy.

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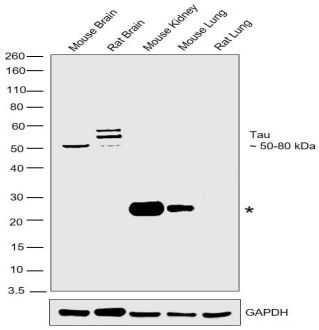
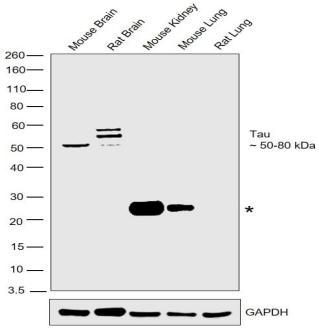
Product Images For Tau Monoclonal Antibody (TAU-5)



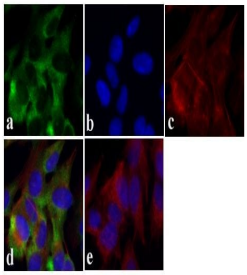
Tau Antibody (AHB0042)

Antibody specificity was demonstrated by detection of differential basal expression of the target across Mouse Brain, Rat Brain, Mouse Kidney, Mouse Lung, Rat Lung owing to their inherent genetic constitution. Relative expression of Tau was observed in Mouse and Rat Brain as compared to Kidney and Lung tissues using Anti-Tau Monoclonal Antibody (TAU-5) (Product # AHB0042) in Western Blot. {RE}



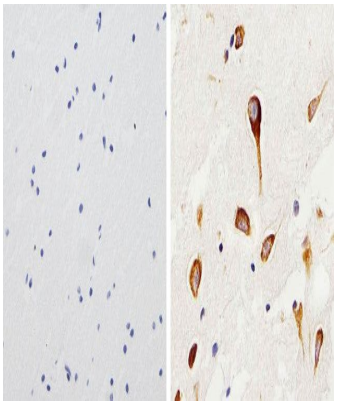
Tau Antibody (AHB0042) in WB

Western blot was performed using Anti-Tau Monoclonal Antibody (TAU-5) (Product # AHB0042) and 40-80 kDa bands corresponding to Tau were observed in Mouse and Rat Brain but not in Mouse Kidney or Mouse and Rat Lung. Tissue extracts (30 µg lysate) of Mouse Brain (Lane 1), Rat Brain (Lane 2), Mouse Kidney (Lane 3), Mouse Lung (Lane 4) or Rat Lung (Lane 5) were electrophoresed using NuPAGE™ 10% Bis-Tris Protein Gel (Product # NP0301BOX). Resolved proteins were then transferred onto a Nitrocellulose membrane (Product # IB23001) by iBlot® 2 Dry Blotting System (Product # IB21001). The blot was probed with the primary antibody (1:500) and detected by chemiluminescence with Goat anti-Mouse IgG (H+L) Superclonal™ Recombinant Secondary Antibody, HRP (Product # A28177,1:4000) using the iBright FL 1000 (Product # A32752). Chemiluminescent detection was performed using Novex® ECL Chemiluminescent Substrate Reagent Kit (Product # WP20005). Additional band at 25 kDa corresponds to tissue IgG and has been marked with an asterisk.



Tau Antibody (AHB0042) in ICC/IF

Immunofluorescent analysis of Tau (Tau-5) was done on 70% confluent log phase SHSY5Y cells. The cells were fixed with 4% paraformaldehyde for 15 minutes, permeabilized with 0.25% Triton™ X-100 for 10 minutes, and blocked with 5% BSA for 1 hour at room temperature. The cells were labeled with Tau (Tau-5) Mouse Monoclonal Antibody (Product # AHB0042) at 1 µg/mL in 1% BSA and incubated for 3 hours at room temperature and then labeled with Alexa Fluor 488 Rabbit Anti-Mouse IgG Secondary Antibody (Product # A-11059) at a dilution of 1:400 for 30 minutes at room temperature (Panel a: green). Nuclei (Panel b: blue) were stained with SlowFade Gold Antifade Mountant with DAPI (Product # S36938). F-actin (Panel c: red) was stained with Alexa Fluor 594 Phalloidin (Product # A12381). Panel d is a merged image showing cytoplasmic localization and panel e is a no primary antibody control. The images were captured at 20X magnification.



Tau Antibody (AHB0042) in IHC (P)

Immunohistochemistry analysis of Tau showing staining in the cytoplasm and weak nuclear staining of paraffin-embedded human brain tissue (right) compared to a negative control without primary antibody (left). To expose target proteins, antigen retrieval was performed using 10mM sodium citrate (pH 6.0), microwaved for 8-15 min. Following antigen retrieval, tissues were blocked in 3% H2O2-methanol for 15 min at room temperature, washed with ddH2O and PBS, and then probed with a Tau (Product # AHB0042) diluted in 3% BSA-PBS at a dilution of 1:200 overnight at 4°C in a humidified chamber. Tissues were washed extensively in PBST and detection was performed using an HRP-conjugated secondary antibody followed by colorimetric detection using a DAB kit. Tissues were counterstained with hematoxylin and dehydrated with ethanol and xylene to prep for mounting.

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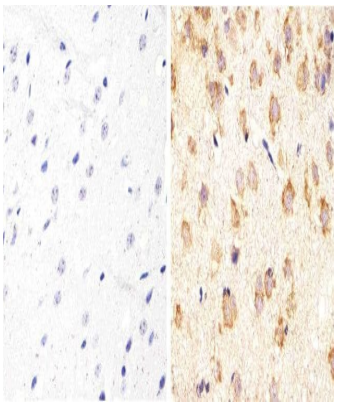
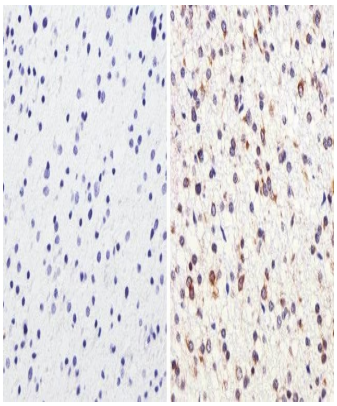


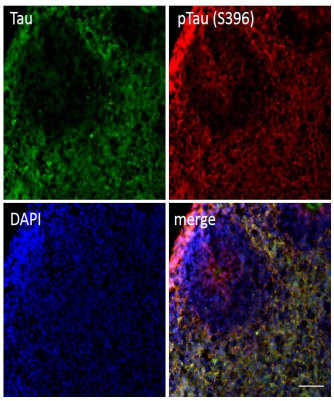
Tau Antibody (AHB0042) in IHC (P)

Immunohistochemistry analysis of Tau showing staining in the cytoplasm of paraffin-embedded mouse brain tissue (right) compared to a negative control without primary antibody (left). To expose target proteins, antigen retrieval was performed using 10mM sodium citrate (pH 6.0), microwaved for 8-15 min. Following antigen retrieval, tissues were blocked in 3% H2O2-methanol for 15 min at room temperature, washed with ddH2O and PBS, and then probed with a Tau (Product # AHB0042) diluted in 3% BSA-PBS at a dilution of 1:20 overnight at 4°C in a humidified chamber. Tissues were washed extensively in PBST and detection was performed using an HRP-conjugated secondary antibody followed by colorimetric detection using a DAB kit. Tissues were counterstained with hematoxylin and dehydrated with ethanol and xylene to prep for mounting.



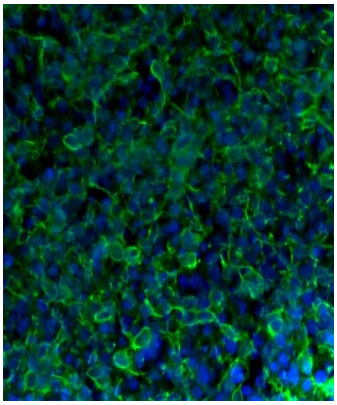
Tau Antibody (AHB0042) in IHC (P)

Immunohistochemistry analysis of Tau showing staining in the cytoplasm of paraffin-embedded human astroglioma (right) compared to a negative control without primary antibody (left). To expose target proteins, antigen retrieval was performed using 10mM sodium citrate (pH 6.0), microwaved for 8-15 min. Following antigen retrieval, tissues were blocked in 3% H2O2-methanol for 15 min at room temperature, washed with ddH2O and PBS, and then probed with a Tau (Product # AHB0042) diluted in 3% BSA-PBS at a dilution of 1:50 overnight at 4°C in a humidified chamber. Tissues were washed extensively in PBST and detection was performed using an HRP-conjugated secondary antibody followed by colorimetric detection using a DAB kit. Tissues were counterstained with hematoxylin and dehydrated with ethanol and xylene to prep for mounting.



Tau Antibody (AHB0042) in IHC (F)

Immunofluorescent analysis of p-Tau (S396) and Tau in human iPSC-derived forebrain organoids derived at Day 40. The organoids were fixed with 4% PFA for 1 hour at room temperature, followed by incubation with 30% sucrose solution overnight at 4°C. The organoids were then embedded in OCT and cryosectioned at 5 µm, permeabilized with 0.2% Triton X-100 for 20 min, and blocked with 10% donkey serum in PBS for 30 min at room temperature. Organoid slices were stained with a Mouse Tau (TAU-5) monoclonal antibody (green; Product # AHB0042) at a dilution of 1:500 and a Rabbit p-Tau (Ser396) polyclonal antibody (red; Product # 44-752G) at a dilution of 1:500 in blocking buffer overnight at 4°C, and then incubated with Donkey anti-Mouse Alexa Fluor 488 (Product # R37114), Donkey anti-Rabbit Alexa Fluor 568 (Product # A10042) at a dilution of 1:1000 as well as DAPI (blue; 1:25000) in blocking solution at room temperature for 1 hour. Images were taken at 20X magnification. Scale bar: 50 µm. Data courtesy of Dr. Zhexing Wen at Emory University.



Tau Antibody (AHB0042) in IHC (F)

Immunofluorescent analysis of Tau in human iPSC-derived forebrain organoids derived at Day 40. The organoids were fixed with 4% PFA for 1 hour at room temperature, followed by incubation with 30% sucrose solution overnight at 4°C. The organoids were then embedded in OCT and cryosectioned at 5 µm, permeabilized with 0.2% Triton X-100 for 20 min, and blocked with 10% donkey serum in PBS for 30 min at room temperature. Organoid slices were stained with a Mouse Tau monoclonal antibody (green; Product # AHB0042) at a dilution of 1:500 in blocking buffer overnight at 4°C, and then incubated with Donkey anti-Mouse Alexa Fluor 488 (Product # R37114) at a dilution of 1:1000 as well as DAPI (blue; 1:25000) in blocking solution at room temperature for 1 hour. Images were taken at 20X magnification. Data courtesy of Dr. Zhexing Wen at Emory University.

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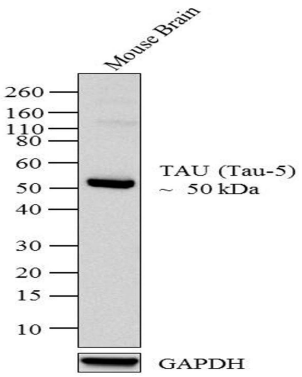

Tau Antibody (AHB0042) in WB

Western blot analysis of Tau-5 was performed by loading 20 µg of Mouse brain tissue lysates using Novex® NuPAGE® 4-12 % Bis-Tris gel (Product # NP0321BOX), XCell SureLock Electrophoresis System (Product # EI0002), Novex® Sharp Pre-Stained Protein Standard (Product # LC5800), and iBlot® Dry Blotting System (Product # IB21001). Proteins were transferred to a nitrocellulose membrane and blocked with 5 % skim milk for 1 hour at room temperature. Tau-5 was detected at ~50 kDa using Tau-5 Mouse Monoclonal Antibody (Product # AHB0042) at 0.5-1 µg/mL in 2.5 % skim milk at 4°C overnight on a rocking platform. Goat anti-Mouse HRP Secondary Antibody (Product # 62-6520) at 1:4000 dilution was used and chemiluminescent detection was performed using Pierce™ ECL Western blotting Substrate (Product # 32106).

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PubMed References For Tau Monoclonal Antibody (TAU-5)

67 Western Blot References

Species / Dilution	Summary
Not Applicable / 1:1000	AHB0042 was used in western blot to evaluate if OVX/stress affects levels of Alzheimer's disease-related molecules in the mouse hippocampus Neuroscience letters (2008; 433: 141) **"Enhanced activity of hippocampal BACE1 in a mouse model of postmenopausal memory deficits."** Author(s):Fukuzaki E,Takuma K,Himeno Y,Yoshida S,Funatsu Y,Kitahara Y,Mizoguchi H,Ibi D,Koike K,Inoue M,Yamada K PubMed Article URL:http://dx.doi.org/10.1016/j.neulet.2007.12.060
Not Applicable / 1:1000	AHB0042 was used in immunocytochemistry and western blot to determine modulation of cellular release of tau by FRMD4A-cytohesin signaling Journal of cell science (2016; 129: 2003) **"FRMD4A-cytohesin signaling modulates the cellular release of tau."** Author(s):Yan X,Nykänen NP,Brunello CA,Haapasalo A,Hiltunen M,Uronen RL,Huttunen HJ PubMed Article URL:http://dx.doi.org/10.1242/jcs.180745
Mouse / Not Cited	AHB0042 was used in Western Blotting to show reduced, behaviourally driven, gamma oscillations before the onset of plaque formation or cognitive decline in a mouse model of Alzheimer's disease. Nature (2016; 540: 230) **"Gamma frequency entrainment attenuates amyloid load and modifies microglia."** Author(s):Iaccarino HF,Singer AC,Martorell AJ,Rudenko A,Gao F,Gillingham TZ,Mathys H,Seo J,Kritskiy O,Abdurrob F,Adaikkan C,Canter RG,Rueda R,Brown EN,Boyden ES,Tsai LH PubMed Article URL:http://dx.doi.org/10.1038/nature20587
Mouse / Not Cited	AHB0042 was used in Western Blotting to show that GENUS can be achieved through multiple sensory modalities with wide-ranging effects across multiple brain areas to improve cognitive function. Cell (2019; 177: 256) **"Multi-sensory Gamma Stimulation Ameliorates Alzheimer's-Associated Pathology and Improves Cognition."** Author(s):Martorell AJ,Paulson AL,Suk HJ,Abdurrob F,Drummond GT,Guan W,Young JZ,Kim DN,Kritskiy O,Barker SJ,Mangena V,Prince SM,Brown EN,Chung K,Boyden ES,Singer AC,Tsai LH PubMed Article URL:http://dx.doi.org/10.1016/j.cell.2019.02.014
Bovine / 1:200	AHB0042 was used in Western Blotting to investigate the role of glucocorticoids in mitophagy inhibition and subsequent synaptic defects in hippocampal neurons, SH-SY5Y cells, and ICR mice. Nature communications (2021; 12:) **"BNIP3L/NIX-mediated mitophagy protects against glucocorticoid-induced synapse defects."** Author(s):Choi GE,Lee HJ,Chae CW,Cho JH,Jung YH,Kim JS,Kim SY,Lim JR,Han HJ PubMed Article URL:http://dx.doi.org/10.1038/s41467-020-20679-y
Mouse / Not Cited	AHB0042 was used in western blot identify and investigate Reelin-regulated signaling pathways in the brain Molecular and cellular biology (2007; 27: 7113) **"Reelin signals through phosphatidylinositol 3-kinase and Akt to control cortical development and through mTor to regulate dendritic growth."** Author(s):Jossin Y,Goffinet AM PubMed Article URL:http://dx.doi.org/10.1128/MCB.00928-07
Rat / 1:200	AHB0042 was used in Western Blotting to provide a novel mechanism of the hyperphosphorylation of tau and identify both PKA and GSK-3 as promising therapeutic targets for AD and other tauopathies. The Journal of biological chemistry (2004; 279: 50078) **"Tau becomes a more favorable substrate for GSK-3 when it is prephosphorylated by PKA in rat brain."** Author(s):Liu SJ,Zhang JY,Li HL,Fang ZY,Wang Q,Deng HM,Gong CX,Grundke-Iqbal I,Iqbal K,Wang JZ PubMed Article URL:http://dx.doi.org/10.1074/jbc.M406109200
Mouse / Not Cited	AHB0042 was used in western blot to test the effects of high fat diet using a mouse model of familial Alzheimer disease. Biochimica et biophysica acta (2015; 1852: 1687) **"High-fat diet-induced deregulation of hippocampal insulin signaling and mitochondrial homeostasis deficiences contribute to Alzheimer disease pathology in rodents."** Author(s):Petrov D,Pedrós I,Artiach G,Sureda FX,Barroso E,Pallàs M,Casadesús G,Beas-Zarate C,Carro E,Ferrer I,Vazquez-Carrera M,Folch J,Camins A PubMed Article URL:http://dx.doi.org/10.1016/j.bbadis.2015.05.004

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	AHB0042 was used in Western Blotting to evaluate the effects of aerobic exercise on the phosphorylation and acetylation of tau and the expressions of tau-related proteins, after middle cerebral artery occlusion (MCAO) stroke.
Rat / 1:1000	International journal of molecular sciences (2020; 21:) **"Effects of Aerobic Exercise on Tau and Related Proteins in Rats with the Middle Cerebral Artery Occlusion."** Author(s):Mankhong S,Kim S,Moon S,Lee KH,Jeon HE,Hwang BH,Beak JW,Joa KL,Kang JH PubMed Article URL:http://dx.doi.org/10.3390/ijms21165842
	AHB0042 was used in western blot to study reduction of hyperphosphorilated Tau by beta-secretase 1's targeting implying autophagy actors in 3xTg-AD mice
Not Applicable / 1:1000	Frontiers in cellular neuroscience (2016; 9:) **"-Secretase 1's Targeting Reduces Hyperphosphorilated Tau, Implying Autophagy Actors in 3xTg-AD Mice."** Author(s):Piedrahita D,Castro-Alvarez JF,Boudreau RL,Villegas-Lanau A,Kosik KS,Gallego-Gomez JC,Cardona-Gómez GP PubMed Article URL:http://dx.doi.org/10.3389/fncel.2015.00498
	AHB0042 was used in Western Blot to pave the way to evaluate whether similar results occur in vivo and through what mechanisms.
Rat / Not Cited	International journal of molecular sciences (2022; 23:) **"Extracellular Vesicles Derived from Young Neural Cultures Attenuate Astrocytic Reactivity In Vitro."** Author(s):Almansa D,Peinado H,García-Rodríguez R,Casadomé-Perales Á,Dotti CG,Guix FX PubMed Article URL:http://dx.doi.org/10.3390/ijms23031371
	AHB0042 was used in Western Blotting to study whether HS3ST2 and its 3-O-sulphated heparan sulphate products are involved in the mechanisms leading to the abnormal phosphorylation of tau in Alzheimer's disease and related tauopathies.
Human / Not Cited	Brain : a journal of neurology (2015; 138: 1339) **"HS3ST2 expression is critical for the abnormal phosphorylation of tau in Alzheimer's disease-related tau pathology."** Author(s):Sepulveda-Diaz JE,Alavi Naini SM,Huynh MB,Ouidja MO,Yanicostas C,Chantepie S,Villares J,Lamari F,Jospin E,van Kuppevelt TH,Mensah-Nyagan AG,Raisman-Vozari R,Soussi-Yanicostas N,Papy-Garcia D PubMed Article URL:http://dx.doi.org/10.1093/brain/awv056
	AHB0042 was used in western blot to elucidate how the removal of assembled neurofilaments from axons or misaccumulating neurofilaments in motor neuron cell bodies slows disease in a mouse model of amyotrophic lateral sclerosis
Not Applicable / 1:1000	Proceedings of the National Academy of Sciences of the United States of America (2005; 102: 10351) **"Altered axonal architecture by removal of the heavily phosphorylated neurofilament tail domains strongly slows superoxide dismutase 1 mutant-mediated ALS."** Author(s):Lobsiger CS,Garcia ML,Ward CM,Cleveland DW PubMed Article URL:http://dx.doi.org/10.1073/pnas.0503862102
	AHB0042 was used in Western Blot to investigate the potential role of miR-124/PTPN1 in the tau pathology of AD.
Mouse / Not Cited	Journal of neurochemistry (2020; 154: 441) **"Correcting abnormalities in miR-124/PTPN1 signaling rescues tau pathology in Alzheimer's disease."** Author(s):Hou TY,Zhou Y,Zhu LS,Wang X,Pang P,Wang DQ,Liuyang ZY,Man H,Lu Y,Zhu LQ,Liu D PubMed Article URL:http://dx.doi.org/10.1111/jnc.14961
	AHB0042 was used in western blot to determine that Abeta deposition or oestrogen deficiency increases PP2A phosphorylation to compromise tau dephosphorylation and cause neurofibrillary tangle formation in Alzheimer's disease
Not Applicable / 1:1000	Journal of cellular and molecular medicine (2008; 12: 241) **"Phosphorylated PP2A (tyrosine 307) is associated with Alzheimer neurofibrillary pathology."** Author(s):Liu R,Zhou XW,Tanila H,Bjorkdahl C,Wang JZ,Guan ZZ,Cao Y,Gustafsson JA,Winblad B,Pei JJ PubMed Article URL:http://dx.doi.org/10.1111/j.1582-4934.2008.00249.x
	AHB0042 was used in Western Blotting to study the synaptic role of Tau.
Mouse / 1:500	Philosophical transactions of the Royal Society of London. Series B, Biological sciences (2014; 369:) **"Microtubule-associated protein tau is essential for long-term depression in the hippocampus."** Author(s):Kimura T,Whitcomb DJ,Jo J,Regan P,Piers T,Heo S,Brown C,Hashikawa T,Murayama M,Seok H,Sotiropoulos I,Kim E,Collingridge GL,Takashima A,Cho K PubMed Article URL:http://dx.doi.org/10.1098/rstb.2013.0144
	AHB0042 was used in Western Blotting to investigate the function of miR-128 as a tumour suppressor in glioma.
Mouse / Not Cited	Oncogene (2012; 31: 1884) **"Pro-neural miR-128 is a glioma tumor suppressor that targets mitogenic kinases."** Author(s):Papagiannakopoulos T,Friedmann-Morvinski D,Neveu P,Dugas JC,Gill RM,Huillard E,Liu C,Zong H,Rowitch DH,Barres BA,Verma IM,Kosik KS PubMed Article URL:http://dx.doi.org/10.1038/onc.2011.380

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Rat / Not Cited	The EMBO journal (2005; 24: 209) **"A Cdk5 inhibitory peptide reduces tau hyperphosphorylation and apoptosis in neurons."** Author(s):Zheng YL,Kesavapany S,Gravell M,Hamilton RS,Schubert M,Amin N,Albers W,Grant P,Pant HC PubMed Article URL:http://dx.doi.org/10.1038/sj.emboj.7600441
Mouse / 1:2,500	AHB0042 was used in Western Blotting to conclude that the role of ER stress in AD pathogenesis needs to be carefully addressed in future studies. The Journal of biological chemistry (2018; 293: 3118) **"Endoplasmic reticulum stress responses in mouse models of Alzheimer's disease: Overexpression paradigm versus knockin paradigm."** Author(s):Hashimoto S,Ishii A,Kamano N,Watamura N,Saito T,Ohshima T,Yokosuka M,Saido TC PubMed Article URL:http://dx.doi.org/10.1074/jbc.M117.811315
Not Applicable / 1:1000	AHB0042 was used in western blot to study Alzheimer's disease by use of a domestic cat animal model Acta neuropathologica communications (2015; 3:) **"The domestic cat as a natural animal model of Alzheimer's disease."** Author(s):Chambers JK,Tokuda T,Uchida K,Ishii R,Tatebe H,Takahashi E,Tomiyama T,Une Y,Nakayama H PubMed Article URL:http://dx.doi.org/10.1186/s40478-015-0258-3
Not Applicable / 1:10,000	AHB0042 was used in immunohistochemistry - paraffin section and western blot to characterize a new mouse model of Alzheimer's disease The American journal of pathology (2006; 169: 599) **"Alzheimer's disease-like tau neuropathology leads to memory deficits and loss of functional synapses in a novel mutated tau transgenic mouse without any motor deficits."** Author(s):Schindowski K,Bretteville A,Leroy K,Bégard S,Brion JP,Hamdane M,Buée L PubMed Article URL:http://dx.doi.org/10.2353/ajpath.2006.060002
Not Applicable / Not Cited	AHB0042 was used in Western Blotting to show that non-lethal reductions in axonal transport, and perhaps other types of minor axonal stress, are sufficient to induce and/or accelerate abnormal tau behaviour characteristic of Alzheimer's disease and other neurodegenerative tauopathies. Human molecular genetics (2010; 19: 4399) **"Kinesin-1 transport reductions enhance human tau hyperphosphorylation, aggregation and neurodegeneration in animal models of tauopathies."** Author(s):Falzone TL,Gunawardena S,McCleary D,Reis GF,Goldstein LS PubMed Article URL:http://dx.doi.org/10.1093/hmg/ddq363
Human / 1:2000	AHB0042 was used in western blot to see that the granulin Alzheimer's disease risk variant has no significant effects on florbetapir positron emission tomographic amyloid imaging and cerebrospinal fluid Abeta levels Acta neuropathologica (2017; 133: 785) **"Opposing effects of progranulin deficiency on amyloid and tau pathologies via microglial TYROBP network."** Author(s):Takahashi H,Klein ZA,Bhagat SM,Kaufman AC,Kostylev MA,Ikezu T,Strittmatter SM PubMed Article URL:http://dx.doi.org/10.1007/s00401-017-1668-z
Not Applicable / 1:500	AHB0042 was used in western blot to study how induction of Calyculin A in N2a cells can attenuate axonal transport impairment and axonopathy by Berberine PloS one (2015; 9:) **"Berberine attenuates axonal transport impairment and axonopathy induced by Calyculin A in N2a cells."** Author(s):Liu X,Zhou J,Abid MD,Yan H,Huang H,Wan L,Feng Z,Chen J PubMed Article URL:http://dx.doi.org/10.1371/journal.pone.0093974
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Mouse / Not Cited	AHB0042 was used in Western Blotting to find that DHCR24 knockdown can lead to tau hyperphosphorylation in the astrocyte itself by activating lipid raft-dependent Ras/MEK/ERK signaling, which might contribute to the pathogenesis of AD and other degenerative tauopathies. Molecular neurobiology (2022; 59: 5856) **"DHCR24 Knockdown Induces Tau Hyperphosphorylation at Thr181, Ser199, Ser262, and Ser396 Sites via Activation of the Lipid Raft-Dependent Ras/MEK/ERK Signaling Pathway in C8D1A Astrocytes."** Author(s):Mai M,Guo X,Huang Y,Zhang W,Xu Y,Zhang Y,Bai X,Wu J,Zu H PubMed Article URL:http://dx.doi.org/10.1007/s12035-022-02945-w
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Not Applicable / 1:500	AHB0042 was used in western blot to study the tauopathic changes in the striatum of the alpha-synuclein A53T mutant mouse PloS one (2011; 6:) **"Tauopathic changes in the striatum of A53T -synuclein mutant mouse model of Parkinson's disease."** Author(s):Wills J,Credle J,Haggerty T,Lee JH,Oaks AW,Sidhu A PubMed Article URL:http://dx.doi.org/10.1371/journal.pone.0017953
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Not Applicable / 1:1000	Neuroscience (2007; 144: 119) **"Caspase-3- and calpain-mediated tau cleavage are differentially prevented by estrogen and testosterone in beta-amyloid-treated hippocampal neurons."** Author(s):Park SY,Tournell C,Sinjoanu RC,Ferreira A PubMed Article URL:http://dx.doi.org/10.1016/j.neuroscience.2006.09.012
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Human / Not Cited	AHB0042 was used in Western Blotting to propose that dysregulation of neural gene networks may set in motion the pathologic cascade that leads to AD. Cell reports (2019; 26: 1112) **"REST and Neural Gene Network Dysregulation in iPSC Models of Alzheimer's Disease."** Author(s):Meyer K,Feldman HM,Lu T,Drake D,Lim ET,Ling KH,Bishop NA,Pan Y,Seo J,Lin YT,Su SC,Church GM,Tsai LH,Yankner BA PubMed Article URL:http://dx.doi.org/10.1016/j.celrep.2019.01.023
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Mouse / Not Cited	The Journal of biological chemistry (2003; 278: 10506) **"Tau phosphorylation by cyclin-dependent kinase 5/p39 during brain development reduces its affinity for microtubules."** Author(s):Takahashi S,Saito T,Hisanaga S,Pant HC,Kulkarni AB PubMed Article URL:http://dx.doi.org/10.1074/jbc.M211964200
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	AHB0042 was used in Western Blotting to investigate to what extent tau cleavage and the generation of the neurotoxic tau45-230 fragment is associated with amyotrophic lateral sclerosis.
Human / Not Cited Mouse / Not Cited	Molecular medicine (Cambridge, Mass.) (2016; 22: 477) **"The Neurotoxic TAU₄₅₋₂₃₀ Fragment Accumulates in Upper and Lower Motor Neurons in Amyotrophic Lateral Sclerosis Subjects."** Author(s):Vintilescu CR,Afreen S,Rubino AE,Ferreira A PubMed Article URL:http://dx.doi.org/10.2119/molmed.2016.00095
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	Anesthesiology (2012; 116: 779) **"Tau phosphorylation and sevoflurane anesthesia: an association to postoperative cognitive impairment."** Author(s):Le Freche H,Brouillette J,Fernandez-Gomez FJ,Patin P,Caillierez R,Zommer N,Sergeant N,Buée-Scherrer V,Lebuffe G,Blum D,Buée L PubMed Article URL:http://dx.doi.org/10.1097/ALN.0b013e31824be8c7
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	Molecular and cellular neurosciences (2020; 108:) **"Dystrophic Dmd^{mdx} rats show early neuronal changes (increased S100 and Tau5) at 8 months, supporting severe dystropathology in this rodent model of Duchenne muscular dystrophy."** Author(s):Krishnan VS,Thanigaiarasu LP,White R,Crew R,Larcher T,Le Guiner C,Grounds MD PubMed Article URL:http://dx.doi.org/10.1016/j.mcn.2020.103549

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	AHB0042 was used in western blot to elucidate how liraglutide affects disease and cognitive function in a mouse model of Alzheimer disease
Mouse / 1:1000	Neurochemical research (2017; 42: 2326) **"Liraglutide Improves Water Maze Learning and Memory Performance While Reduces Hyperphosphorylation of Tau and Neurofilaments in APP/PS1/Tau Triple Transgenic Mice."** Author(s):Chen S,Sun J,Zhao G,Guo A,Chen Y,Fu R,Deng Y PubMed Article URL:http://dx.doi.org/10.1007/s11064-017-2250-8
	AHB0042 was used in Western Blotting to study UPS14 aptamers for degrading proteotoxic proteins to protect cells from neurodegenerative stressors.
Human / Not Cited	Scientific reports (2015; 5:) **"Facilitated Tau Degradation by USP14 Aptamers via Enhanced Proteasome Activity."** Author(s):Lee JH,Shin SK,Jiang Y,Choi WH,Hong C,Kim DE,Lee MJ PubMed Article URL:http://dx.doi.org/10.1038/srep10757
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Human / 1:500	The European journal of neuroscience (2011; 33: 1598) **"Hyperphosphorylated Tau in an -synuclein-overexpressing transgenic model of Parkinson's disease."** Author(s):Haggerty T,Credle J,Rodriguez O,Wills J,Oaks AW,Masliah E,Sidhu A PubMed Article URL:http://dx.doi.org/10.1111/j.1460-9568.2011.07660.x
	AHB0042 was used in western blot to investigate the neurotoxicity of 2,3,7,8-tetrachlorodibenzo-p-dioxin
Not Applicable / 1:1000	Toxicology (2009; 255: 65) **"2,3,7,8-TCDD neurotoxicity in neuroblastoma cells is caused by increased oxidative stress, intracellular calcium levels, and tau phosphorylation."** Author(s):Sul D,Kim HS,Cho EK,Lee M,Kim HS,Jung WW,Hwang KW,Park SY PubMed Article URL:http://dx.doi.org/10.1016/j.tox.2008.10.006
	AHB0042 was used in Western Blot to investigate whether trazodone, a licensed antidepressant with hypnotic efficacy in dementia, can reduce disease-related cellular pathways and improve memory and sleep in male rTg4510 mice with a tauopathy-like phenotype.
Mouse / 1:4000	The Journal of neuroscience : the official journal of the Society for Neuroscience (2022; 42: 3494) **"Improved Sleep, Memory, and Cellular Pathological Features of Tauopathy, Including the NLRP3 Inflammasome, after Chronic Administration of Trazodone in rTg4510 Mice."** Author(s):de Oliveira P,Cella C,Locker N,Ravindran KKG,Mendis A,Wafford K,Gilmour G,Dijk DJ,Winsky-Sommerer R PubMed Article URL:http://dx.doi.org/10.1523/JNEUROSCI.2162-21.2022
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Mouse / Not Cited	Biochimica et biophysica acta (2014; 1842: 1556) **"Early alterations in energy metabolism in the hippocampus of APPswe/PS1dE9 mouse model of Alzheimer's disease."** Author(s):Pedrós I,Petrov D,Allgaier M,Sureda F,Barroso E,Beas-Zarate C,Auladell C,Pallàs M,Vázquez-Carrera M,Casadesús G,Folch J,Camins A PubMed Article URL:http://dx.doi.org/10.1016/j.bbadis.2014.05.025
	AHB0042 was used in western blot to investigate the axon initial segment in neurological disorders
Human / Not Cited	Molecular neurodegeneration (2016; 11:) **"Acetylated tau destabilizes the cytoskeleton in the axon initial segment and is mislocalized to the somatodendritic compartment."** Author(s):Sohn PD,Tracy TE,Son HI,Zhou Y,Leite RE,Miller BL,Seeley WW,Grinberg LT,Gan L PubMed Article URL:http://dx.doi.org/10.1186/s13024-016-0109-0
	AHB0042 was used in Western Blot to open new perspectives for the design of highly specific ligands with diagnostic and therapeutic potential.
Human / 1:20000	The Journal of biological chemistry (2021; 297:) **"Structural mapping techniques distinguish the surfaces of fibrillar 1N3R and 1N4R human tau."** Author(s):Caroux E,Redeker V,Madiona K,Melki R PubMed Article URL:http://dx.doi.org/10.1016/j.jbc.2021.101252
	AHB0042 was used in western blot to investigate survival and neuronal differentiation of human BM-MSCs by pulsed electromagnetic fields
Not Applicable / Not Cited	Life sciences (2016; 151: 130) **"Pulsed electromagnetic fields promote survival and neuronal differentiation of human BM-MSCs."** Author(s):Urnukhsaikhan E,Cho H,Mishig-Ochir T,Seo YK,Park JK PubMed Article URL:http://dx.doi.org/10.1016/j.lfs.2016.02.066

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	AHB0042 was used in Western Blot to demonstrate a simple and precise method to simultaneously generate iPSC lines with different gene dosages using paired Cas9 nickases.
Human / 1:20000	Communications biology (2021; 4:) **"Efficient manipulation of gene dosage in human iPSCs using CRISPR/Cas9 nickases."** Author(s):Ye T,Duan Y,Tsang HWS,Xu H,Chen Y,Cao H,Chen Y,Fu AKY,Ip NY PubMed Article URL:http://dx.doi.org/10.1038/s42003-021-01722-0
Human / Not Cited	AHB0042 was used in Western Blotting to discover whether tau biosensors capable of monitoring tau oligomer conformation are able to identify tool compounds that modulate the structure and conformation of these tau assemblies, providing key insight into the unique structural fingerprints of toxic tau oligomers.
	Alzheimer's & dementia : the journal of the Alzheimer's Association (2019; 15: 1489) **"Targeting the ensemble of heterogeneous tau oligomers in cells: A novel small molecule screening platform for tauopathies."** Author(s):Lo CH,Lim CK,Ding Z,Wickramasinghe SP,Braun AR,Ashe KH,Rhoades E,Thomas DD,Sachs JN PubMed Article URL:http://dx.doi.org/10.1016/j.jalz.2019.06.4954
Not Applicable / 1:1000	AHB0042 was used in immunohistochemistry - paraffin section and western blot to evaluate motor function and tau pathology of P301S tau transgenic mice
	Molecular neurodegeneration (2014; 9:) **"Long-term treadmill exercise attenuates tau pathology in P301S tau transgenic mice."** Author(s):Ohia-Nwoko O,Montazari S,Lau YS,Eriksen JL PubMed Article URL:http://dx.doi.org/10.1186/1750-1326-9-54
Not Applicable / 1:1000	AHB0042 was used in western blot to elucidate the neuroprotective mechanisms by which curcumin protects against Abeta-induced toxicity
	Food and chemical toxicology : an international journal published for the British Industrial Biological Research Association (2008; 46: 2881) **"Curcumin protected PC12 cells against beta-amyloid-induced toxicity through the inhibition of oxidative damage and tau hyperphosphorylation."** Author(s):Park SY,Kim HS,Cho EK,Kwon BY,Phark S,Hwang KW,Sul D PubMed Article URL:http://dx.doi.org/10.1016/j.fct.2008.05.030
Not Applicable / 1:5000	AHB0042 was used in western blot to assess the deletion of Ccr2 and cavity size and tau pathology after mild traumatic brain injury
	Journal of neuroinflammation (2015; 12:) **"Ccr2 deletion dissociates cavity size and tau pathology after mild traumatic brain injury."** Author(s):Gyoneva S,Kim D,Katsumoto A,Kokiko-Cochran ON,Lamb BT,Ransohoff RM PubMed Article URL:http://dx.doi.org/10.1186/s12974-015-0443-0

1 Dot blot References

Species / Dilution	Summary
Mouse / Not Cited	AHB0042 was used in Dot blot to suggest that conformation-specific antibodies targeting -synuclein aggregates are promising therapeutic agents for PD and related synucleinopathies.
	Experimental neurobiology (2022; 31: 29) **"Conformation-specific Antibodies Targeting Aggregated Forms of -synuclein Block the Propagation of Synucleinopathy."** Author(s):Choi M,Kim TK,Ahn J,Lee JS,Jung BC,An S,Kim D,Lee MJ,Mook-Jung I,Lee SH,Lee SJ PubMed Article URL:http://dx.doi.org/10.5607/en21039

1 Immunoelectrophoresis References

Species / Dilution	Summary
Human / 1:1000	AHB0042 was used in Immunoelectrophoresis to suggest that PTM profiles occur through the post-translational control of the master PTM remodeling enzymes themselves.
	The Journal of biological chemistry (2022; 298:) **"Identification of a reciprocal negative feedback loop between tau-modifying proteins MARK2 kinase and CBP acetyltransferase."** Author(s):Tabassum Z,Tseng JH,Isemann C,Tian X,Chen Y,Herring LE,Cohen TJ PubMed Article URL:http://dx.doi.org/10.1016/j.jbc.2022.101977

5 Immunoprecipitation References

Species / Dilution	Summary

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	AHB0042 was used in IP/Western Blotting to identify molecules which may provide new tools to study a neuronal migration and cortical plate development.
Mouse / Not Cited	Cerebral cortex (New York, N.Y. : 1991) (2007; 17: 211) **"Identification of small molecules that interfere with radial neuronal migration and early cortical plate development."** Author(s):Zhou L,Jossin Y,Goffinet AM PubMed Article URL:http://dx.doi.org/10.1093/cercor/bhj139
	AHB0042 was used in Immunoprecipitation to show that Rac1 levels decreased in the frontal cortex of Alzheimer's disease patients compared to non-demented controls.
Mouse / 1:1,000	Acta neuropathologica communications (2018; 6:) **"Rac1 activation links tau hyperphosphorylation and A dysmetabolism in Alzheimer's disease."** Author(s):Borin M,Saraceno C,Catania M,Lorenzetto E,Pontelli V,Paterlini A,Fostinelli S,Avesani A,Di Fede G,Zanusso G,Benussi L,Binetti G,Zorzan S,Ghidoni R,Buffelli M,Bolognin S PubMed Article URL:http://dx.doi.org/10.1186/s40478-018-0567-4
	AHB0042 was used in Immunoprecipitation to identify salivary Tau species as potential biomarkers of Alzheimer's disease.
Human / Not Cited	Journal of Alzheimer's disease : JAD (2012; 27: 299) **"Salivary tau species are potential biomarkers of Alzheimer's disease."** Author(s):Shi M,Sui YT,Peskind ER,Li G,Hwang H,Devic I,Ginghina C,Edgar JS,Pan C,Goodlett DR,Furay AR,Gonzalez-Cuyar LF,Zhang J PubMed Article URL:http://dx.doi.org/10.3233/JAD-2011-110731
	AHB0042 was used in Immunoprecipitation to find that mitochondrial SOD2 deficiency exacerbates amyloid burden and significantly reduces metal levels in the brain, while increasing levels of Ser-396 phosphorylated tau.
Human / Not Cited	PloS one (2007; 2:) **"Mitochondrial oxidative stress causes hyperphosphorylation of tau."** Author(s):Melov S,Adlard PA,Morten K,Johnson F,Golden TR,Hinerfeld D,Schilling B,Mavros C,Masters CL,Volitakis I,Li QX,Laughton K,Hubbard A,Cherny RA,Gibson B,Bush AI PubMed Article URL:http://dx.doi.org/10.1371/journal.pone.0000536
	AHB0042 was used in Proximity Ligation Assay (PLA) to suggest an enabling role of tau in the pathogenesis of autism and identify tau reduction as a potential therapeutic strategy for some of the disorders that cause this condition.
Mouse / 1:1000	Neuron (2020; 106: 421) **"Tau Reduction Prevents Key Features of Autism in Mouse Models."** Author(s):Tai C,Chang CW,Yu GQ,Lopez I,Yu X,Wang X,Guo W,Mucke L PubMed Article URL:http://dx.doi.org/10.1016/j.neuron.2020.01.038

1 in situ PLA References

Species / Dilution	Summary
	AHB0042 was used in Proximity Ligation Assay (PLA) to suggest an enabling role of tau in the pathogenesis of autism and identify tau reduction as a potential therapeutic strategy for some of the disorders that cause this condition.
Mouse / 1:4000	Neuron (2020; 106: 421) **"Tau Reduction Prevents Key Features of Autism in Mouse Models."** Author(s):Tai C,Chang CW,Yu GQ,Lopez I,Yu X,Wang X,Guo W,Mucke L PubMed Article URL:http://dx.doi.org/10.1016/j.neuron.2020.01.038

8 Immunohistochemistry References

Species / Dilution	Summary
	AHB0042 was used in Immunohistochemistry to show that the hydrophobic patches of aggregates from acetylated peptides were different when compared to wild-type (WT) peptide.
Rat / 1:200	ACS chemical neuroscience (2020; 11: 1178) **"Acetylation of A42 at Lysine 16 Disrupts Amyloid Formation."** Author(s):Adhikari R,Yang M,Saikia N,Dutta C,Alharbi WFA,Shan Z,Pandey R,Tiwari A PubMed Article URL:http://dx.doi.org/10.1021/acschemneuro.0c00069
	AHB0042 was used in Immunohistochemistry to study the pathological association of tau dynamics in both cerebral ischemia and Alzheimer's disease.
Rat / 1:400	Journal of cerebral blood flow and metabolism : official journal of the International Society of Cerebral Blood Flow and Metabolism (2017; 37: 2441) **"Modifications of tau protein after cerebral ischemia and reperfusion in rats are similar to those occurring in Alzheimer's disease - Hyperphosphorylation and cleavage of 4- and 3-repeat tau."** Author(s):Fujii H,Takahashi T,Mukai T,Tanaka S,Hosomi N,Maruyama H,Sakai N,Matsumoto M PubMed Article URL:http://dx.doi.org/10.1177/0271678X16668889

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Not Applicable / Not Cited	AHB0042 was used in immunohistochemistry to determine the distribution of tauopathy in different regions of the brain using the alpha-Syn overexpressing mouse model BMC neuroscience (2011; 12:) **"Region-specific tauopathy and synucleinopathy in brain of the alpha-synuclein overexpressing mouse model of Parkinson's disease."** Author(s):Kaul T,Credle J,Haggerty T,Oaks AW,Masliah E,Sidhu A PubMed Article URL:http://dx.doi.org/10.1186/1471-2202-12-79
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Human / Not Cited	Acta neuropathologica (2002; 103: 599) **"Ubiquitin-positive neuronal and tau 2-positive glial inclusions in frontotemporal dementia of motor neuron type."** Author(s):Forno LS,Langston JW,Herrick MK,Wilson JD,Murayama S PubMed Article URL:http://dx.doi.org/10.1007/s00401-001-0509-1
Not Applicable / 1:2000	AHB0042 was used in immunohistochemistry to examine the brains of 6-month-old rats treated neonatally with the glutamatergic beta-N-methylamino-L-alanine Toxicological sciences : an official journal of the Society of Toxicology (2012; 130: 391) **"Neonatal exposure to the cyanobacterial toxin BMAA induces changes in protein expression and neurodegeneration in adult hippocampus."** Author(s):Karlsson O,Berg AL,Lindström AK,Hanrieder J,Arnerup G,Roman E,Bergquist J,Lindquist NG,Brittebo EB,Andersson M PubMed Article URL:http://dx.doi.org/10.1093/toxsci/kfs241
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5 Immunocytochemistry References

Species / Dilution	Summary
Mouse / Not Cited	AHB0042 was used in Immunoprecipitation to show that Rac1 levels decreased in the frontal cortex of Alzheimer's disease patients compared to non-demented controls. Acta neuropathologica communications (2018; 6:) **"Rac1 activation links tau hyperphosphorylation and A dysmetabolism in Alzheimer's disease."** Author(s):Borin M,Saraceno C,Catania M,Lorenzetto E,Pontelli V,Paterlini A,Fostinelli S,Avesani A,Di Fede G,Zanusso G,Benussi L,Binetti G,Zorzan S,Ghidoni R,Buffelli M,Bolognin S PubMed Article URL:http://dx.doi.org/10.1186/s40478-018-0567-4
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Non-human primate / Not Cited Human / Not Cited	Toxicologic pathology (2007; 35: 972) **"Expression of serine/threonine protein-kinases and related factors in normal monkey and human retinas: the mechanistic understanding of a CDK2 inhibitor induced retinal toxicity."** Author(s):Saturno G,Pesenti M,Cavazzoli C,Rossi A,Giusti AM,Gierke B,Pawlak M,Venturi M PubMed Article URL:http://dx.doi.org/10.1080/01926230701748271

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Human / Not Cited	Molecular biology of the cell (2004; 15: 2720) **"Modulation of microtubule dynamics by tau in living cells: implications for development and neurodegeneration."** Author(s):Bunker JM,Wilson L,Jordan MA,Feinstein SC PubMed Article URL:http://dx.doi.org/10.1091/mbc.e04-01-0062
Rat / Not Cited	The EMBO journal (2005; 24: 209) **"A Cdk5 inhibitory peptide reduces tau hyperphosphorylation and apoptosis in neurons."** Author(s):Zheng YL,Kesavapany S,Gravell M,Hamilton RS,Schubert M,Amin N,Albers W,Grant P,Pant HC PubMed Article URL:http://dx.doi.org/10.1038/sj.emboj.7600441

2 Immunohistochemistry (Frozen) References

Species / Dilution	Summary
Not Applicable / 1:500	AHB0042 was used in immunohistochemistry - frozen section and western blot to utilize a time course study of sciatic nerves from aging mice to gain a neurogenic perspective of sarcopenia Journal of neuropathology and experimental neurology (2016; 75: 464) **"A Neurogenic Perspective of Sarcopenia: Time Course Study of Sciatic Nerves From Aging Mice."** Author(s):Krishnan VS,White Z,McMahon CD,Hodgetts SI,Fitzgerald M,Shavlakadze T,Harvey AR,Grounds MD PubMed Article URL:http://dx.doi.org/10.1093/jnen/nlw019
Mouse / 1:100	AHB0042 was used in immunohistochemistry - frozen section to investigate the expression patterns of HA synthases in the murine central nervous system Journal of Alzheimer's disease : JAD (2018; 57: 395) **"Tau Pathology Promotes the Reorganization of the Extracellular Matrix and Inhibits the Formation of Perineuronal Nets by Regulating the Expression and the Distribution of Hyaluronic Acid Synthases."** Author(s):Li Y,Li ZX,Jin T,Wang ZY,Zhao P PubMed Article URL:http://dx.doi.org/10.3233/JAD-160804

5 Miscellaneous PubMed References

Species / Dilution	Summary
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Rat / 1:1000	AHB0042 was used in immunohistochemistry - paraffin section and western blot to elucidate the molecular mechanisms of ischemic tolerance Neurological sciences : official journal of the Italian Neurological Society and of the Italian Society of Clinical Neurophysiology (2011; 32: 229) **"Ischemic preconditioning attenuates of ischemia-induced degradation of spectrin and tau: implications for ischemic tolerance."** Author(s):Nakajima T,Ochi S,Oda C,Ishii M,Ogawa K PubMed Article URL:http://dx.doi.org/10.1007/s10072-010-0359-5
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Mouse / Not Cited	AHB0042 was used in western blot to investigate the contribution of Ang II to Alzheimer disease. FEBS letters (2012; 586: 3737) **"Central angiotensin II-induced Alzheimer-like tau phosphorylation in normal rat brains."** Author(s):Tian M,Zhu D,Xie W,Shi J PubMed Article URL:http://dx.doi.org/10.1016/j.febslet.2012.09.004
Mouse / Not Cited	AHB0042 was used in immunohistochemistry (frozen) to study the composition of pathological granules that appear in degenerative brain diseases. Age (Dordrecht, Netherlands) (2014; 36: 151) **"Presence of a neo-epitope and absence of amyloid beta and tau protein in degenerative hippocampal granules of aged mice."** Author(s):Manich G,del Valle J,Cabezón I,Camins A,Pallàs M,Pelegrí C,Vilaplana J PubMed Article URL:http://dx.doi.org/10.1007/s11357-013-9560-9

2 ELISA References

Species / Dilution	Summary

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	AHB0042 was used in ELISA to test if HFE polymorphisms are associated with alterations in tau phosphorylation in a human neuroblastoma cell line
Not Applicable / Not Cited	Neurobiology of aging (2011; 32: 1409) **"Expression of the HFE allelic variant H63D in SH-SY5Y cells affects tau phosphorylation at serine residues."** Author(s):Hall EC,Lee SY,Mairuae N,Simmons Z,Connor JR PubMed Article URL:http://dx.doi.org/10.1016/j.neurobiolaging.2009.08.012
Human / Not Cited	Journal of biomolecular screening (2004; 9: 122) **"Development of an assay to screen for inhibitors of tau phosphorylation by cdk5."** Author(s):Ahn JS,Musacchio A,Mapelli M,Ni J,Scinto L,Stein R,Kosik KS,Yeh LA PubMed Article URL:http://dx.doi.org/10.1177/1087057103260594

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