







RESEARCH PAPER

Neurotrophic interactions between neurons and astrocytes following AAV1-Rheb(S16H) transduction in the hippocampus in vivo

Min-Tae Jeon^{1,2} | Gyeong Joon Moon^{1,2}  | Sehwan Kim^{1,2} | Minji Choi⁴ | Yong-Seok Oh⁵  | Dong Woon Kim⁶  | Hyung-Jun Kim⁷ | Kea Joo Lee⁷  | Youngshik Choe⁷ | Chang Man Ha⁷ | Il-Sung Jang^{8,9} | Michiko Nakamura⁹ | Catriona McLean^{10,11} | Won-Suk Chung¹² | Won-Ho Shin¹³ | Seok-Geun Lee^{4,14}  | Sang Ryong Kim^{1,2,3,9} 

¹School of Life Sciences, Kyungpook National University, Daegu, Korea

²BK21 plus KNU Creative BioResearch Group, Kyungpook National University, Daegu, Korea

³Institute of Life Science and Biotechnology, Kyungpook National University, Daegu, Korea

⁴Department of Science in Korean Medicine, Graduate School, Kyung Hee University, Seoul, Korea

⁵Department of Brain-Cognitive Science, Daegu-Gyeongbuk Institute of Science and Technology, Daegu, Korea

⁶Department of Anatomy, Brain Research Institute, Chungnam National University School of Medicine, Daejeon, Korea

⁷Department of Neural Development and Disease, Department of Structure and Function of Neural Network, Korea Brain Research Institute, Daegu, Korea

⁸Department of Pharmacology, School of Dentistry, Kyungpook National University, Daegu, Korea

⁹Brain Science and Engineering Institute, Kyungpook National University, Daegu, Korea

¹⁰Victorian Brain Bank Network, Florey Institute of Neuroscience and Mental Health, Parkville, Victoria, Australia

¹¹Department of Anatomical Pathology, Alfred Hospital, Melbourne, Victoria, Australia

¹²Department of Biological Sciences, Korea Advanced Institute of Science and Technology (KAIST), Daejeon, Korea

¹³Predictive Model Research Center, Korea Institute of Toxicology, Daejeon, Korea

¹⁴KHU-KIST Department of Converging Science and Technology, Kyung Hee University, Seoul, Korea

Correspondence

Sang Ryong Kim, School of Life Sciences, Kyungpook National University, Daegu 41566, Korea.

Email: srk75@knu.ac.kr

Seok-Geun Lee, Department of Science in Korean Medicine, Graduate School, Kyung Hee University, Seoul 02447, Korea.

Email: seokgeun@khu.ac.kr

Background and Purpose: We recently reported that AAV1-Rheb(S16H) transduction could protect hippocampal neurons through the induction of brain-derived neurotrophic factor (BDNF) in the rat hippocampus in vivo. It is still unclear how neuronal BDNF produced by AAV1-Rheb(S16H) transduction induces neuroprotective effects in the hippocampus and whether its up-regulation contributes to the enhance of a neuroprotective system in the adult brain.

Abbreviations: AAV, adeno-associated virus; AD, Alzheimer's disease; AP, anterior–posterior; BDNF, brain-derived neurotrophic factor; CA1, cornu ammonis 1; CM, conditioned medium; CNTF, ciliary neurotrophic factor; CNTFR, ciliary neurotrophic factor receptor; CNTFR α , ciliary neurotrophic factor receptor subunit α ; DV, dorsal–ventral; GDNF, glial cell line-derived neurotrophic factor; GFAP, glial fibrillary acidic protein; Iba1, ionized calcium-binding adapter molecule 1; MAP 2, microtubule-associated protein 2; ML, medial–lateral; mTORC1, mammalian target of rapamycin complex 1; MTT, 3-(4,5-dimethylthiazol-2-yl)-2,5-diphenyltetrazolium bromide; NeuN, neuronal nuclei; p-4E-BP1, phosphorylation of the Thr37/46 residues of 4E-BP1; PBST, Tween-20 in PBS; PD, Parkinson's disease; Rheb(S16H), Ras homolog enriched in brain containing a serine-to-histidine mutation at position 16; RT, room temperature; SD, Sprague–Dawley; SN, substantia nigra; TrkB, tropomyosin receptor kinase B

Min-Tae Jeon, Gyeong Joon Moon, Sehwan Kim, and Minji Choi contributed equally to this work.

This is an open access article under the terms of the Creative Commons Attribution-NonCommercial License, which permits use, distribution and reproduction in any medium, provided the original work is properly cited and is not used for commercial purposes.

© 2019 The Authors. British Journal of Pharmacology published by John Wiley & Sons Ltd on behalf of British Pharmacological Society

Funding information

KBRI Research Program of the Ministry of Science, ICT & Future Planning, Grant/Award Numbers: 18-BR-03-01, 18-RB-03-03 and 18-BR-02-06; Ministry of Health & Welfare, Grant/Award Numbers: HI15C1928 and HI14C1135; National Research Foundation of Korea, Grant/Award Numbers: NRF-2017R1D1A1B03032218 and NRF-2017R1A2B4002675

Experimental Approach: To determine the presence of a neuroprotective system in the hippocampus of patients with Alzheimer's disease (AD), we examined the levels of glial fibrillary acidic protein, BDNF and ciliary neurotrophic factor (CNTF) and their receptors, tropomyocin receptor kinase B (TrkB) and CNTF receptor α (CNTFR α), in the hippocampus of AD patients. We also determined whether AAV1-Rheb(S16H) transduction stimulates astroglial activation and whether reactive astrocytes contribute to neuroprotection in models of hippocampal neurotoxicity in vivo and in vitro.

Key Results: AD patients may have a potential neuroprotective system, demonstrated by increased levels of full-length TrkB and CNTFR α in the hippocampus. Further AAV1-Rheb(S16H) transduction induced sustained increases in the levels of full-length TrkB and CNTFR α in reactive astrocytes and hippocampal neurons. Moreover, neuronal BDNF produced by Rheb(S16H) transduction of hippocampal neurons induced reactive astrocytes, resulting in CNTF production through the activation of astrocytic TrkB and the up-regulation of neuronal BDNF and astrocytic CNTF which had synergistic effects on the survival of hippocampal neurons in vivo.

Conclusions and Implications: The results demonstrated that Rheb(S16H) transduction of hippocampal neurons could strengthen the neuroprotective system and this intensified system may have a therapeutic value against neurodegeneration in the adult brain.

1 | INTRODUCTION

Neurotrophic factors play important roles in enhancing neuronal survival, function and plasticity in the adult brain (Huang & Reichardt, 2001; Jeon et al., 2015; Jeong, Nam, Jin, & Kim, 2015; Siegel & Chauhan, 2000). The effects of neurotrophic factors are mediated by specific receptors, such as ciliary neurotrophic factor receptor (CNTFR), which is composed of its ligand-specific α subunit (CNTFR α), leukaemia inhibitory factor receptor β and gp130 (Siegel & Chauhan, 2000; Leibinger & Andreadaki et al., 2012) for ciliary neurotrophic factor (CNTF), as well as tropomyosin receptor kinase B (TrkB; Jeon et al., 2015; Jeong & Nam et al., 2015) for brain-derived neurotrophic factor (BDNF). The loss of neurotrophic factors such BDNF and CNTF may be associated with the pathogenesis of brain diseases (Chauhan, Siegel, & Lee, 2001; Jeon et al., 2015; Jeong et al., 2015; Phillips et al., 1991; Sopova, Gatsiou, Stellos, & Laske, 2014). On the other hand, the up-regulation of these factors induces neuroprotective effects through the activation of mammalian target of rapamycin complex 1 (mTORC1), which stimulates the neuronal survival signalling pathway against neurodegeneration in animal models of neurodegenerative diseases (Jeon et al., 2015; Leibinger, Andreadaki, & Fischer, 2012; Smith et al., 2014). These observations suggest that the sustained expression of neurotrophic factors may be useful for protecting neurons in the adult brain, and the regulatory system of a key molecule that can produce neurotrophic factors is a potential therapeutic target against neurodegenerative diseases.

We recently reported an increase in reactive astrocytes, which protect dopaminergic neurons, in the substantia nigra (SN) of patients with Parkinson's disease (PD; Nam et al., 2015), suggesting

What is already known

- AAV1-Rheb(S16H) transduction could protect hippocampal neurons by inducing brain-derived neurotrophic factor in the rat hippocampus.

What does this study add

- AAV1-Rheb(S16H) transduction of hippocampal neurons increased astrocytic ciliary neurotrophic factor expression.
- AAV1-Rheb(S16H) transduction strengthened the neuroprotective system through neuron–glia neurotrophic interactions in the rat hippocampus.

What is the clinical significance

- The AAV1-Rheb(S16H) transduction approach may be useful for protecting hippocampal neurons against neurodegenerative diseases.

the presence of an endogenous neuroprotective system in lesioned human brains. It is unclear whether reactive astrocytes in the brains of patients with Alzheimer's disease (AD) have a similar potential to protect hippocampal neurons. However, if a target molecule activated by cell-type-specific transduction consistently constructs and intensifies the neuroprotective system through neuron–glia interactions in the lesioned hippocampus of the adult brain, it may

be important to develop a novel therapeutic strategy against neurodegeneration.

Ras homolog enriched in brain (Rheb), which is a member of the **Ras** family of small GTP-binding proteins, is a key regulator of mTORC1 activation associated with neuronal survival and functional maintenance (Jeon et al., 2015; Jeong et al., 2015; Kim, Kareva, Yarygina, Kholodilov, & Burke, 2012). Neuronal transduction with Rheb(S16H), the constitutively active form of Rheb containing a serine-to-histidine mutation at position 16, sustains the activation of mTORC1 (Jeon et al., 2015; Jeong et al., 2015; Kim et al., 2011; Kim et al., 2012). This activation can protect hippocampal neurons by inducing neurotrophic effects, such as the production of neuronal BDNF against thrombin-induced neurotoxicity in vivo (Jeon et al., 2015). However, it is still unknown whether the production of BDNF following Rheb(S16H) transduction of hippocampal neurons contributes to glial activation, which can further produce neurotrophic factors and promote neuroprotective effects (Nam et al., 2015) via paracrine stimulation in the adult brain. In the present study in order to determine the presence of a neuroprotective system, which has been observed in the SN of patients with PD (Nam et al., 2015), we examined the protein levels of glial fibrillary acidic protein (GFAP), a marker of astrocytes, and neurotrophic factors such as BDNF and CNTF with their respective receptors. TrkB and CNTFR α . in the post-mortem hippocampus of patients with AD and age-matched controls. In addition, to verify the possibility of the development of a neuroprotective system following cell-type-specific gene transduction in the hippocampus of the adult brain, we examined whether adeno-associated virus 1 (AAV1)-Rheb(S16H) transduction of hippocampal neurons stimulates the activation of astrocytes in the hippocampus of the adult brain and whether the induction of reactive astrocytes following Rheb(S16H) transduction contributes to neuroprotection in models of hippocampal neurotoxicity in vivo and in vitro.

2 | METHODS

2.1 | Animals

Female Sprague-Dawley (SD) rats (200–220 g, 8-week-old; RRID: RGD_1566440) and pregnant SD rats were obtained from Daehan Biolink (Chungbuk, Korea). For hippocampal neuronal cultures, embryonic (embryonic days 18–19) SD rats were used. For astrocyte cultures, newborn (postnatal day 1) SD rats were used.

2.2 | Human brain tissues

Frozen brain tissues were obtained from the Victorian Brain Bank Network, supported by the Florey Institute of Neuroscience and Mental Health, The Alfred and the Victorian Forensic Institute of Medicine, and funded by Australia's National Health & Medical Research Council and Parkinson's Victoria. The information of the human post-mortem brain samples is shown in Table 1.

TABLE 1 Information of human post-mortem brain samples

	AD patients	Healthy controls	P value
Number of patients	5	5	
Region of brain	Hippocampus	Hippocampus	
Age (years)	76.4 \pm 2.6	76.6 \pm 2.5	>.05
Male gender, n (%)	4 (80)	4 (80)	>.05
PMI (hr)	18.5 \pm 3.5	40.4 \pm 2.4	<.05
Braak stage	6 \pm 0	0.4 \pm 0.4	<.05
Medication (n)	AChE inhibitor (3), antihypertensive (2), antidiabetic (1), antiplatelet (1), antipsychotic (2), antidepressant (2) and hypnotic (3) agents	PPI (1), NSAID (1), antigout agent (1), antiplatelet agent (1) and antihypertensive agent (2)	

Abbreviations: NSAID, nonsteroidal anti-inflammatory drug; PMI, post-mortem interval; PPI, proton-pump inhibitor.

2.3 | Ethics approval and consent to participate

All animal experimental procedures were carried out in accordance with the Guidelines for Animal Care and Use of Kyungpook National University, approved by the Animal Care and Use Committee of Kyungpook National University (No. KNU 2012-37 and 2016-42). All animals were maintained with a 12-h light-dark cycle in a temperature-controlled room and had free access to food and water. The animals were randomly divided into different groups. Human tissue experiments were approved by the Bioethics Committee, Institutional Review Board of Kyungpook National University Industry Foundation (IRB No. 2013-0016 and 2016-0011). Animal studies are reported in compliance with the ARRIVE guidelines (Kilkenny & Browne et al., 2010) and with the recommendations by the *British Journal of Pharmacology*.

2.4 | Production of AAV viral vectors

All vectors used were the AAV1 serotype as previously described (Kim et al., 2011; Kim et al., 2012). A plasmid carrying Rheb was purchased from OriGene Technologies (Rockville, MD, USA). Rheb DNA was amplified and modified to incorporate a FLAG-encoding sequence at the 3'-end by expand long-template PCR (Roche, Basel, Switzerland). Constitutively active Rheb [Rheb(S16H)] was generated with the Phusion Site-Directed Mutagenesis Kit (Cat No. E05545, New England Biolabs, Ipswich, MA, USA) in the pGEM-T vector (Cat No. A1360, Promega, Madison, WI, USA) and cloned into an AAV packaging construct that utilizes the chicken β -actin promoter and contains a

3' WPRE (pBL). AAVs were produced by the University of North Carolina Vector Core, and the genomic titre of Rheb(S16H) was 2×10^{12} viral genomes·ml⁻¹. eGFP, used as a control, was subcloned into the same viral backbone, and viral stocks were produced at titres of 1×10^{12} viral genomes·ml⁻¹.

2.5 | Intra-hippocampal injection

Animals were anaesthetized with 360 mg·kg⁻¹ chloral hydrate (Cat No. C8383; Sigma, St. Louis, MO, USA) by intraperitoneal injection and placed in a stereotaxic frame (David Kopf Instrument, Tujunga, CA, USA). AAV1-GFP or AAV1-Rheb(S16H) was infused unilaterally into the hippocampal cornu ammonis 1 (CA1) area of SD rats (anterior-posterior [AP]: -3.8 mm; medial-lateral [ML]: -2.4 mm; dorsal-ventral [DV]: -3.0 mm, relative to the bregma) according to the brain atlas (Paxinos & Watson, 2007) using 30-gauge injection needles connected to a 10- μ l Hamilton syringe. With an automated syringe pump, 2.0 μ l of viral vector suspension was infused at a rate of 0.1 μ l·min⁻¹ over 20 min and the injection needle was left in place for an additional 5 min to allow diffusion into the tissue and minimize dragging back along the injection track.

To determine the contribution of various signalling pathways, we used neutralizing antibodies and a recombinant protein. Three weeks after viral injection, 200 ng of neutralizing antibodies alone or in combination with 20 U of thrombin (Cat No. T4648; Sigma) dissolved in 4- μ l PBS were injected into the right hippocampal CA1 region as described above. Recombinant BDNF protein (200 ng in 2- μ l PBS; Cat No. 450-02; PeproTech, Rocky Hill, NJ, USA) was injected into the right hippocampal CA1 region of normal rats as described above. The following neutralizing antibodies were used: anti-BDNF (Cat No. SC-20981; RRID:AB_2064213; Santa Cruz Biotechnology, Dallas, TX, USA), anti-TrkB (Cat No. AF1494; RRID:AB_2155264; R&D Systems, Minneapolis, MN, USA), anti-CNTF (Cat No. AB-557-NA; RRID:AB_354368; R&D Systems) and anti-CNTFR α (Cat No. AF-303-NA; RRID:AB_2083208; R&D Systems).

2.6 | Astrocyte cultures and collection of conditioned medium

Primary astrocytes from the cerebral hemispheres of newborn SD rats were cultured as described previously (Kaeck & Banker, 2006). In brief, cortices were triturated into single cells in DMEM (Cat No. SH30243.01; Hyclone, Logan, UT, USA) containing 10% horse serum (Cat No. 16050130; Invitrogen, Carlsbad, CA, USA) and 1% antibiotics-antimycotics (Cat No. 30-004-CI; Corning, Inc., Corning, NY, USA) and plated on 75-cm² poly-D-lysine-coated culture flasks (0.5 hemisphere/flask; BD Biosciences, Franklin Lakes, NJ, USA). After 2 days, unattached cells and debris were removed by changing the medium. Cultures were fed every 2–3 days with fresh medium until they reached 70–80% confluence.

To determine whether BDNF-treated astrocyte-conditioned medium (CM) contributes to hippocampal neuronal survival after

exposure to thrombin, secondary astrocyte cultures were washed and treated with 80 ng·ml⁻¹ BDNF for 24 hr. The cells were washed three times with PBS and incubated in serum-free medium for 24 hr. The CM was then collected and centrifuged at 2,000 \times g for 10 min at 4°C and filtered through a 0.2- μ m filter to remove cell debris. The untreated astrocyte-CM served as the control. All media were stored at -70°C until use.

2.7 | Treatment of astrocyte cultures

To determine whether exogenous BDNF leads to the activation of mTORC1 and release of CNTF in astrocytes, the primary astrocyte cultures were treated with recombinant BDNF (10, 40, and 80 ng·ml⁻¹) for 24 hr in serum-free medium. We further examined whether the BDNF/TrkB signalling pathway is critical for increasing CNTF release by activating mTORC1 in astrocytes. The primary astrocyte cultures were exposed to 80 ng·ml⁻¹ recombinant BDNF alone or in combination with anti-TrkB neutralizing antibody, 10- μ M GNF5837 (Cat No. 4559; Tocris Bioscience, Bristol, UK), or 80-nM rapamycin (Cat No. 9904; Cell Signaling Technology, Danvers, MA, USA) in serum-free medium for 24 hr.

2.8 | Hippocampal neuronal cultures and treatment

Primary hippocampal neurons were obtained from fetal SD rats as described previously (Kaeck & Banker, 2006). In brief, hippocampal tissues were dissociated by mild mechanical trituration, plated at 4×10^5 cells·ml⁻¹ in 24-well culture plates previously coated with 1 mg·ml⁻¹ poly-D-lysine (Cat No. P6407; Sigma), and maintained in DMEM supplemented with 5% FBS (Cat No. SH30048.03; Hyclone). Three days after plating, the medium was replaced with Neurobasal/B27 medium (Cat No. 21103049; Invitrogen) containing cytosine arabinoside (final concentration 5 μ M; Cat No. C6645; Sigma) to halt the proliferation of non-neuronal cells. One third of the medium was replaced with fresh medium every 3 days. The cultures were maintained at 37°C in a humidified 5% CO₂ atmosphere. To determine the effects of BDNF-treated astrocyte-CM on hippocampal neuronal survival after thrombin treatment, hippocampal neuronal cultures were exposed to 50 U·ml⁻¹ thrombin alone or in combination with BDNF-treated astrocyte-CM or anti-CNTFR α neutralizing antibody for 48 hr.

2.9 | MTT assay

Hippocampal neuronal cell viability was evaluated by 3-(4,5-dimethylthiazol-2-yl)-2,5-diphenyltetrazolium bromide (MTT; Cat No. M5655; Sigma) assay. In brief, hippocampal neuronal cells were incubated with 0.5 mg·ml⁻¹ MTT in serum-free medium at 37°C under 5% CO₂ for 2 hr. After incubation, the medium was removed, and an equal volume of dimethylformamide (Cat No. 227056; Sigma) was added. The plate was gently mixed on an orbital shaker for 10 min. The absorbance was

measured at 570 nm using the VersaMax Microplate Reader (Molecular Devices, Sunnyvale, CA, USA).

2.10 | Immunohistochemical staining

The immuno-related procedures used comply with the recommendations made by the *British Journal of Pharmacology* (Alexander et al., 2018). Animals were transcardially perfused and their brains were fixed and brain sections (30 μm thick) were processed for immunohistochemical staining as previously described with some modifications (Kim et al., 2018). In brief, brain sections were rinsed with PBS and incubated at 4°C with primary antibodies for 48 hr. Then the sections were rinsed with PBS-0.5% BSA, incubated at room temperature (RT) with the appropriate biotinylated secondary antibodies, which included goat anti-mouse IgG (1:400; Cat No. 5450-0011; RRID: AB_2687537; SeraCare Life Sciences, Milford, MA, USA), anti-rabbit IgG (1:400; Cat No. BA-1000; RRID:AB_2313606; Vector Laboratories, Burlingame, CA, USA), and rabbit anti-goat IgG (1:400; Cat No. BA-5000; RRID:AB_2336126; Vector Laboratories), and processed with an avidin–biotin complex kit (Cat No. PK6100; RRID: AB_2336819; Vector Laboratories). The signal was detected by incubating the sections with 0.5 $\text{mg}\cdot\text{ml}^{-1}$ 3,3'-diaminobenzidine (Cat No. D5637; Sigma) in 0.1-M PBS containing 0.003% H_2O_2 . The stained samples were analysed under a bright-field microscope (Axio Imager; RRID:SCR_016980; Carl Zeiss, Jena, Germany). The primary antibodies were goat anti-CNTF (1:100; Cat No. AB-557-NA; RRID: AB_354368; R&D Systems), goat anti-TrkB (1:100; Cat No. AF1494; RRID:AB_2155264; R&D Systems), goat anti-CNTFR α (1:100; Cat No. AF-303-NA; RRID:AB_2083208; R&D Systems), rabbit anti-GFAP (1:2,000; Cat No. AB5804; RRID:AB_2109645; Millipore, Billerica, MA, USA), rabbit anti-ionized calcium-binding adapter molecule 1 (Iba1; 1:2,000; Cat No. 019-19741; RRID:AB_839504; Wako Pure Chemical Industries, Osaka, Japan), rabbit anti-phospho-4E-BP1 (1:1,000; Cat No. 2855; RRID:AB_560835; Cell Signaling Technology) and mouse anti-neuronal nuclei (NeuN; 1:500; Cat No. MAB377; RRID:AB_2298772; Millipore). For Nissl staining, brain sections were mounted on gelatin-coated slides, stained with 0.5% cresyl violet (Cat No. C5042; Sigma) and analysed using a bright-field microscope (RRID:SCR_016980; Axio Imager; Carl Zeiss).

For immunofluorescence labelling, brain sections were rinsed and incubated for 48 hr with one of the following pairs: rabbit anti-FLAG (1:3,000; Cat No. F7425; RRID:AB_439687; Sigma) and mouse anti-NeuN (1:500; Millipore), rabbit anti-FLAG (1:3,000; Sigma) and mouse anti-GFAP (1:2,000; Millipore), mouse anti-FLAG (1:2,000; Cat No. F3165; RRID:AB_259529; Sigma) and rabbit anti-Iba1 (1:2,000; Wako Pure Chemical Industries), mouse anti-NeuN (1:500; Millipore) and rabbit anti-BDNF (1:200; Santa Cruz Biotechnology), rabbit anti-GFAP (1:2,000; Millipore) and goat anti-TrkB (1:100; R&D Systems), mouse anti-OX-42 (1:500; Cat No. MCA275GA; RRID:AB_566455; Serotec, Oxford, UK) and goat anti-TrkB (1:100; R&D Systems), rabbit anti-p-4E-BP1 (1:1,000; Cell Signaling Technology) and mouse anti-GFAP (1:2,000; Millipore), mouse anti-GFAP (1:2,000; Millipore) and goat

anti-CNTF (1:100; R&D Systems), mouse anti-NeuN (1:500; Millipore) and goat anti-CNTFR α (1:100; R&D Systems), mouse anti-GFAP (1:2,000; Millipore) and goat anti-CNTFR α (1:100; R&D Systems), mouse anti-OX-42 (1:500; Serotec) and goat anti-CNTFR α (1:100; R&D Systems), mouse anti-OX-42 (1:500; Serotec) and rabbit anti-p-4E-BP1 (1:1,000; Cell Signaling Technology), rabbit anti-Iba1 (1:2,000; Wako Pure Chemical Industries) and goat anti-CNTF (1:100; R&D Systems), or mouse anti-microtubule-associated protein 2 (MAP 2; 1:500; Cat No. MAB3418; RRID:AB_94856; Millipore) and goat anti-TrkB (1:100; R&D Systems). The sections were then rinsed, incubated with Texas Red-conjugated second antibodies: goat anti-rabbit IgG (1:400; Cat No. TI-1000; RRID:AB_2336199; Vector Laboratories), rabbit anti-goat IgG (1:400; Cat No. TI-5000; RRID: AB_2336129; Vector Laboratories), horse anti-mouse IgG (1:400; Cat No. TI-2000; RRID:AB_2336178; Vector Laboratories), fluorescein isothiocyanate-conjugated donkey anti-rabbit IgG (1:200; Cat No. 711-095-152; RRID:AB_2315776; Jackson Lab, West Grove, PA, USA), or horse anti-mouse IgG (1:200; Cat No. FI-2000; RRID: AB_2336176; Vector Laboratories) for 1 hr, washed and mounted with Vectashield mounting medium (Cat No. H-1000; RRID: AB_2336789; Vector Laboratories). The stained sections were imaged using a fluorescence microscope (Axio Imager; Carl Zeiss). Negative controls for BDNF and TrkB were prepared by omitting the primary antibodies. The sections were incubated with 1.5 $\mu\text{g}\cdot\text{ml}^{-1}$ diaminodiphenylindole solution (Cat No. D1306; RRID:AB_2629482; Invitrogen) for 5 min, washed, and mounted with Vectashield mounting medium (Vector Laboratories).

2.11 | Western blot analysis

Western blot analysis was performed as described previously (Kim et al., 2018). In brief, the lysates obtained from astrocyte cultures and hippocampal tissues were homogenized and centrifuged at 4°C for 15 min at 14,000 \times g. The supernatant was transferred to a fresh tube, and the concentration was determined using a bicinchoninic acid assay kit (Cat No. 5000116; Bio-Rad Laboratories, Hercules, CA, USA). The samples were boiled at 100°C for 5 min before gel loading and equal amounts of protein (20 μg) were loaded into each lane with loading buffer. Proteins analysed by gel electrophoresis were transferred to polyvinylidene difluoride membranes (Millipore) using an electrophoretic transfer system (Bio-Rad Laboratories) and the membranes were incubated overnight at 4°C with specific primary antibodies. The following primary antibodies were used: mouse anti- β -actin (1:1,000; Cat No. SC-47778; RRID: AB_626632; Santa Cruz Biotechnology), rabbit anti-BDNF (1:500; Santa Cruz Biotechnology), goat anti-CNTF (1:1,000; R&D Systems), goat anti-TrkB (1:1,000; R&D Systems), goat anti-CNTFR α (1:1,000; R&D Systems), rabbit anti-IL-1 β (1:1,000; Cat No. SC-7884; RRID: AB_2124476; Santa Cruz Biotechnology), mouse anti-TNF- α (1:1,000; Cat No. SC-52746; RRID:AB_630341; Santa Cruz Biotechnology), rabbit anti-p-4E-BP1 (1:1,000; Cell Signaling Technology), rabbit anti-4E-BP1 (1:1,000; Cell Signaling Technology),

mouse anti-NeuN (1:500; Millipore), mouse anti-GFAP (1:500; Millipore) and rabbit anti-Iba1 (1:1,000; Wako Pure Chemical Industries). After washing, the membranes were incubated with HRP-conjugated secondary antibodies (1:5,000; Santa Cruz Biotechnology), which included goat anti-mouse IgG (Cat No. SC-2005; RRID:AB_631736), anti-rabbit IgG (Cat No. SC-2004; RRID:AB_631746) and mouse anti-goat IgG (Cat No. SC-2354; RRID:AB_628490) for 1 hr at RT and the blots were developed using enhanced chemiluminescence western blot detection reagents (GE Healthcare Life Sciences, Little Chalfont, UK) with an X-ray film (Agfa, Mortsel, Belgium) or the LAS-500 image analyser (GE Healthcare Life Sciences). The signal was analysed with Multi Gauge version 3.0 (RRID:SCR_014299; Fuji Photo Film, Tokyo, Japan). All histograms were quantitatively analysed based on the density of target proteins normalized to the β -actin band for each sample.

2.12 | Materials

The chemicals used in the present study were chloralhydrate, thrombin, cytosine arabinoside, MTT, and dimethylformamide (Sigma, St. Louis, MO, USA); GNF5837 (Tocris Bioscience, Bristol, UK); rapamycin (CellSignaling Technology, Danvers, MA, USA).

2.13 | Quantification of CNTF

To quantify the CNTF released in the BDNF-treated astrocyte culture medium, we used commercially available ELISA kits according to the manufacturer's protocol (Cat No. DY557; R&D Systems). In brief, a mouse-anti-rat CNTF capture antibody was coated at $2 \mu\text{g}\cdot\text{ml}^{-1}$ in 96-well immunoassay plates (Corning) overnight at RT. The plates were blocked with 1% BSA for at least 1 hr at RT and washed with 0.01% Tween-20 in PBS (PBST). Next, 100 μl of the CM of astrocyte cultures was added to each well, incubated for 2 hr at RT and washed with PBST. Biotinylated goat anti-rat CNTF detection antibody, diluted to $200 \text{ ng}\cdot\text{ml}^{-1}$ with blocking buffer, was added to the wells for 1 hr at RT. The assay was developed with substrate solution for 20 min at RT followed by the addition of stop solution. The absorbance was measured at 450 nm using the VersaMax Microplate Reader (Molecular Devices).

2.14 | Quantification of NeuN-positive neurons and p-4E-BP-1-positive cells in the hippocampus

As previously described (Jeon et al., 2015), alternate sections were obtained at 3.3-, 3.6-, 4.16- and 4.3-mm posterior to the bregma. NeuN-positive neurons were counted in a rectangular box ($1 \times 0.05 \text{ mm}$) for every selected section. The rectangular box was located over the CA1 cell layer beginning 1.5 mm lateral to the midline and only neurons with normal visible nuclei were counted for two sections in each level (eight regions for each animal). The number of p-4E-BP-1-positive cells in the hippocampus was quantified in six rectangular areas of $0.4 \times 0.3 \text{ mm}$ for each animal. The number of cells

was expressed as a percentage of the contralateral control using a light microscope (Carl Zeiss) at a magnification of 200 \times .

2.15 | Statistical analysis

All experiments were independently conducted five times, and all measurements were performed by an investigator who was blinded to the experimental groups. The precise number of animals used are given in the figure legend. The data and statistical analysis comply with the recommendations of the British Journal of Pharmacology on experimental design and analysis in pharmacology (Curtis et al., 2015). All values are expressed as the mean \pm SEM. Data normality was assessed using the Shapiro-Wilk test, and subsequent testing was performed using a Student's unpaired *t*-test, or one-way ANOVA with Tukey's post-hoc test. If the data is not satisfying the condition of normality then the non-parametric test was performed using the Kruskal-Wallis test. *P* values < 0.05 were considered statistically significant. All statistical analyses and graph generation were performed using SigmaPlot software (RRID:SCR_003210; Systat Software, San Jose, CA, USA).

2.16 | Nomenclature of targets and ligands

Key protein targets and ligands in this article are hyperlinked to corresponding entries in <http://www.guidetopharmacology.org>, the common portal for data from the IUPHAR/BPS Guide to PHARMACOLOGY (Harding et al., 2018) and are permanently archived in the Concise Guide to PHARMACOLOGY 2019/2020 (Alexander et al., 2019).

3 | RESULTS

3.1 | Elevation of the protein levels of GFAP, full-length TrkB and CNTFR α in the post-mortem hippocampus of patients with AD

To investigate the presence of an endogenous neuroprotective system in the hippocampus of patients with AD, which may be similar to the increase in reactive astrocytes exerting neuroprotection in the SN of patients with PD (Nam et al., 2015), we measured the protein levels of GFAP and neurotrophic factors such as BDNF and CNTF with their respective receptors TrkB and CNTFR α in the post-mortem hippocampus of AD patients and age-matched controls by western blotting (Figure 1). Western blot analysis of NeuN (a marker of neurons) showed significant neuronal loss in the hippocampus of AD patients compared with age-matched controls (Figure 1a). However, the protein levels of GFAP were significantly higher in the hippocampus of AD patients compared with age-matched controls (Figure 1a; **P* < .05 vs. CON). In addition, we observed a significant decrease in BDNF and a tendency for CNTF expression to decrease in the hippocampus of AD patients compared with age-matched controls (Figure 1b). In contrast, the levels of both full-length TrkB (TrkB-FL), which has a high

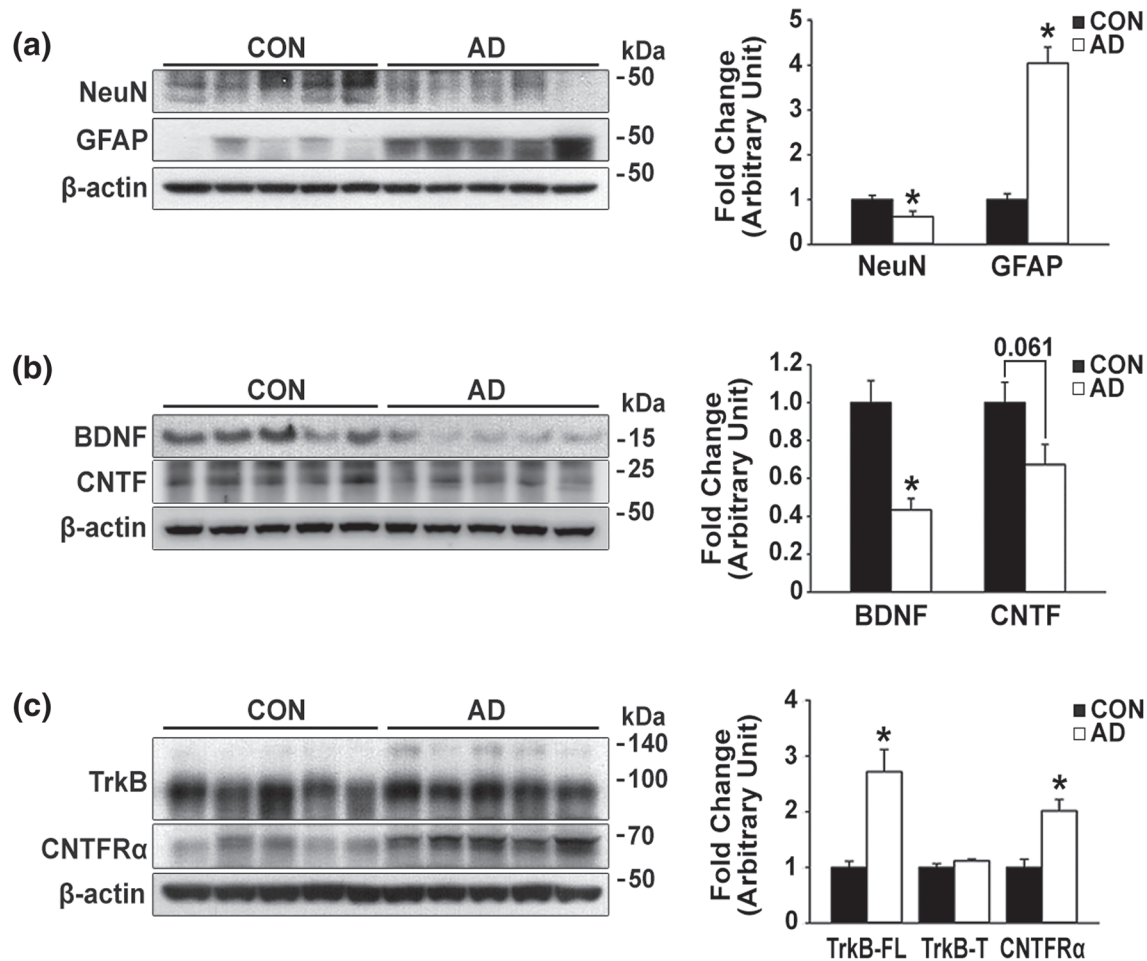


FIGURE 1 Comparison of protein expression profiles in the hippocampus of AD patients and age-matched controls. (a) Patient and tissue information. A total of 10 post-mortem brains were used: five from individuals with a clinical diagnosis of probable AD and five from individuals without neurological disorders (CON). (b–d) Western blot analysis of the levels of NeuN (neuronal marker), GFAP (astrocyte marker), BDNF, CNTF, TrkB, CNTFR α (trophic factors and their receptors) and β -actin in the hippocampus of CON ($n = 5$) and AD patients ($n = 5$). Differences between groups were evaluated by Student's unpaired t -test at a significance level of $*P < .05$

affinity for BDNF (Ferrer et al., 1999; Vidaurre et al., 2012) and CNTFR α , were significantly increased in the hippocampus of AD patients (Figure 1c). Therefore, similar to PD patients (Nam et al., 2015), a neuroprotective system involving an increase in reactive astrocytes may be present in the hippocampus of AD patients. However this neuroprotective system may not be sufficient to control neurodegeneration in the brain of AD patients.

3.2 | Astroglial activation following Rheb(S16H) transduction of hippocampal neurons in the adult brain

We previously reported that neuronal transduction with AAV1-Rheb(S16H) could induce neuroprotective effects against thrombin-induced neurotoxicity by increasing BDNF expression in hippocampal neurons in vivo (Jeon et al., 2015). Although the expression of a target protein following Rheb(S16H) transduction was limited to within neurons (Figures 2a,d and S1A), we recently found that its administration induced reactive astrocytes in the hippocampus of the rat brain.

Double immunofluorescence staining for GFP and NeuN or FLAG epitope and NeuN demonstrated that the target proteins GFP and FLAG were expressed in hippocampal neurons (Figure 2a) but not glial cells such as astrocytes (Figure 2d) and microglia (Figure S1A). These were stained with anti-GFAP and anti-Iba1, respectively, at 4 weeks after intra-hippocampal injection of AAV1-GFP or AAV1-Rheb(S16H). Consistent with our previous study (Jeon et al., 2015), the increased expression of neuronal BDNF was observed in the Rheb(S16H)-treated hippocampus, as demonstrated by immunohistochemical staining and western blotting (Figure 2b,c). However, there were apparent morphological changes in astrocytes (Figure 2d,e) and microglia (Figures S1A–C and S2A) following Rheb(S16H) transduction of hippocampal neurons. The protein levels of GFAP (Figure 2f; *) and Iba1 (Figure S2B) were significantly increased in the Rheb(S16H)-treated hippocampus compared with intact and GFP-transduced controls. There was no staining signal for BDNF in negative control tissues (Figure S3A). Moreover, western blot analysis revealed that the protein levels of IL-1 β and TNF- α , which are pro-inflammatory cytokines associated with neurodegeneration in the adult brain (Jang

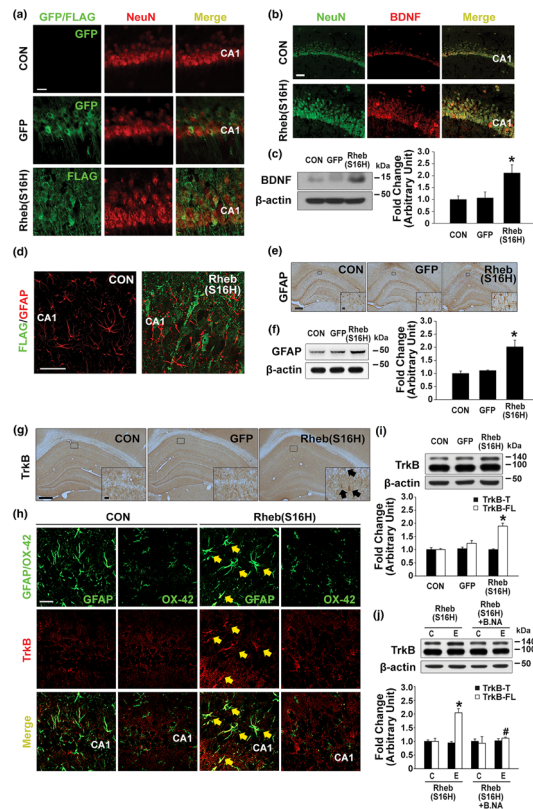


FIGURE 2 Induction of astrocytic activation and TrkB expression by transduction with Rheb(S16H) in hippocampal neurons. Hippocampal tissue sections and protein lysates were obtained from the AAV1-GFP-treated and AAV1-Rheb(S16H)-treated rat brain at 4 weeks after injection. (a) Double immunofluorescence staining against GFP (green) and NeuN (red) or FLAG epitope (green) and NeuN (red) showing the co-localization of these two markers in the CA1 region of the hippocampus. Scale bar, 20 μ m. (b) Double immunofluorescence staining against NeuN (green) and BDNF (red) showing the co-localization of these two markers in the CA1 region of the hippocampus of CON and AAV1-Rheb(S16H)-injected rats. Scale bar, 50 μ m. (c) Western blot analysis showing the levels of BDNF and β -actin in the hippocampus of CON, AAV1-GFP-injected, and AAV1-Rheb(S16H)-injected rats. Differences among groups were evaluated by one-way ANOVA and Tukey's post hoc analysis. * P < .05 versus CON and GFP (n = 5 for each group). (d) FLAG was not expressed in astrocytes as determined by anti-GFAP antibody staining in the CA1 region of the hippocampus of CON and AAV1-Rheb(S16H)-injected rats. Scale bar, 50 μ m. (e) Immunohistochemical staining showing the morphological changes in astrocytes in the CA1 region of the hippocampus of CON, AAV1-GFP-injected and AAV1-Rheb(S16H)-injected rats. Scale bars, 500 μ m (inset 20 μ m). (f) Western blot analysis showing the levels of GFAP and β -actin in the hippocampus of CON, AAV1-GFP-injected and AAV1-Rheb(S16H)-injected rats. Differences among groups were evaluated by one-way ANOVA and Tukey's post hoc analysis. * P < .01 versus CON and GFP (n = 5 for each group). (g) Immunohistochemical staining showing increased TrkB expression in glia-like cells of the hippocampal CA1 region of AAV1-Rheb(S16H)-injected rats (black arrows). Scale bars, 500 μ m (inset 20 μ m). (h) Double immunofluorescence staining against GFAP (green) and TrkB (red) or OX-42 (green) and TrkB (red) showing the co-localization of these two markers in the CA1 region of the hippocampus of CON and AAV1-Rheb(S16H)-injected rats (marked with a yellow arrow). Scale bar, 20 μ m. (i) Western blot analysis showing the levels of TrkB-FL (140 kDa), TrkB-T (100 kDa) and β -actin in the hippocampus of CON, AAV1-GFP-injected and AAV1-Rheb(S16H)-injected rats (n = 5 for each group). (j) Western blot analysis showing the levels of TrkB-FL, TrkB-T, and β -actin in the hippocampus after treatment with AAV1-Rheb(S16H) alone, with or without BDNF neutralizing antibody (B.NA). Differences among groups were evaluated by one-way ANOVA and Tukey's post hoc analysis. * P < .05 versus CON and # P < .05 versus Rheb(S16H) (n = 5 for each group)

et al., 2013; Nam et al., 2014; Shin et al., 2015), were not increased in the Rheb(S16H)-transduced hippocampus (Figure S2B). Therefore, these results suggest that Rheb(S16H) transduction of hippocampal neurons could induce glial activation in the hippocampus of the adult brain, which may not be related to neurotoxic inflammatory responses.

TrkB is a specific receptor involved in BDNF-mediated neurotrophic effects (Jeon et al., 2015; Jeong et al., 2015). The protein level of TrkB-FL, which contains a catalytic kinase domain that can be activated (Ferrer et al., 1999; Vidaurre et al., 2012), was significantly lower in the frontal cortex of AD patients (Ferrer et al., 1999) but not the

hippocampus of AD patients (Connor et al., 1996; Ferrer et al., 1999), suggesting a selective decrease in the BDNF/TrkB neurotrophic signalling pathway in AD. Our immunohistochemical staining results revealed increased TrkB expression in glia-like cells (marked with black arrows) in the Rheb(S16H)-treated hippocampus of the rat brain at 4 weeks after intra-hippocampal viral injection (Figure 2g). To identify the glial cell type overexpressing TrkB in the Rheb(S16H)-treated hippocampus, we further performed double immunofluorescence staining for GFAP and TrkB or OX-42, another marker of microglia (Shin et al., 2015) and TrkB. The results showed

the up-regulation of TrkB in GFAP-positive astrocytes but not microglia (Figure 2h). There was no TrkB-positive signal in negative control sections (Figure S3B). In addition, western blotting demonstrated a significant increase in TrkB-FL expression in the Rheb(S16H)-treated hippocampus compared with the controls (Figure 2i; $^*P < .05$ vs. CON); however, there was no significant change in truncated TrkB (TrkB-T) expression (Figure 2i). Moreover, the inhibition of BDNF activity induced by neutralizing antibodies against BDNF (Jeon et al., 2015; Jeong et al., 2015) significantly attenuated the increase in TrkB-FL following Rheb(S16H) administration as demonstrated by western blotting at 1 day after treatment with neutralizing antibodies (Figure 2j) and there was no change in TrkB-FL expression in the hippocampus treated with neutralizing antibodies alone (Figure S4A). These results were further confirmed by the expression of GFAP and TrkB-FL in astrocyte cultures after recombinant BDNF treatment *in vitro*. The results of preliminary western blot analysis showed that the expression of GFAP and TrkB-FL were increased in astrocytes exposed to recombinant BDNF ($80 \text{ ng}\cdot\text{ml}^{-1}$) for 24 hr (Figure S5). Therefore, these results suggest that the production of neuronal BDNF following Rheb(S16H) transduction could induce an increase in TrkB-FL expression in reactive astrocytes in the hippocampus.

3.3 | Activation of astrocytic mTORC1 by Rheb (S16H)-induced neuronal BDNF through the TrkB-dependent signalling pathway

In addition to the activation of mTORC1 in hippocampal neurons following Rheb(S16H) transduction (Jeon et al., 2015) the phosphorylation of the Thr37/46 residues of 4E-BP1 (p-4E-BP1), which is a representative substrate of mTORC1 (Jeon et al., 2015; Jeong et al., 2015), was increased in reactive astrocytes (yellow arrows) in the Rheb(S16H)-treated hippocampus (Figure 3a). However, it was not increased in the microglia (Figure S1B). In addition, p-4E-BP1-positive cells were significantly increased in the Rheb(S16H)-treated hippocampus compared with the control (Figure 3a). The inhibition of BDNF and TrkB activities by treatment with specific neutralizing antibodies significantly attenuated the up-regulation of p-4E-BP1 in the Rheb(S16H)-treated hippocampus of the rat brain (Figure 3b) and there was no significant change in p-4E-BP1 expression in the hippocampus treated with neutralizing antibodies alone (Figure S4B). To verify that neuronal BDNF produced by Rheb(S16H) transduction could induce mTORC1 activation in astrocytes, we further evaluated p-4E-BP1 expression following treatment with recombinant BDNF in rat astrocyte-enriched cultures. Treatment with recombinant BDNF induced a significant increase in p-4E-BP1 expression in astrocyte-enriched cultures compared with untreated controls (Figure 3c). Further the activation of astrocytic mTORC1 was significantly inhibited by co-treatment with recombinant BDNF and neutralizing antibodies against TrkB (Figure 3c), suggesting that Rheb(S16H)-induced neuronal BDNF could stimulate astrocytic mTORC1 activation through the TrkB-dependent signalling pathway in the hippocampus of the adult brain.

3.4 | Production of astrocytic CNTF through a BDNF/TrkB-dependent signalling pathway in the AAV1-Rheb(S16H)-treated hippocampus

Reactive astrocytes are involved in producing CNTF in the area surrounding the lesioned brain *in vivo* (Albrecht et al., 2003; Nam et al., 2015), suggesting that the induction of astroglial activation, which results in the production of neurotrophic factors such as CNTF. This may be a useful strategy for protecting neurons in the adult brain. In the present study, immunohistochemical staining for CNTF showed that Rheb(S16H) transduction of hippocampal neurons induced an increase in CNTF expression in glial cells in the hippocampus (marked with black arrows) at 4 weeks post intra-hippocampal viral injection (Figure 4a). Moreover, increased CNTF expression was observed in reactive astrocytes, as demonstrated by double immunofluorescence staining for GFAP and CNTF (Figure 4b). However, it was not observed in the microglia (Figure S1C). Consistent with these results of immunohistochemical staining for CNTF, western blotting also showed the up-regulation of CNTF following Rheb(S16H) transduction in the hippocampus (Figure 4c). To further clarify the mechanism of astrocytic CNTF production *in vivo* we examined whether the inhibition of BDNF or TrkB activity by treatment with specific neutralizing antibodies, attenuates the production of CNTF following Rheb(S16H) transduction in the hippocampus. Western blot analysis showed no significant change in CNTF expression in the hippocampus treated with neutralizing antibodies alone (Figure S4C). However, the neutralization of BDNF and TrkB significantly attenuated the increase in CNTF expression in the Rheb(S16H)-treated hippocampus (Figure 4d). In addition, treatment with recombinant BDNF by intra-hippocampal injection significantly increased the level of CNTF expression in reactive astrocytes in the hippocampus, as demonstrated by immunohistochemical staining (Figure 4e) and western blotting (Figure 4f). The increase in astrocytic CNTF following Rheb(S16H) transduction of hippocampal neurons *in vivo*, which may be controlled by inhibition of the BDNF/TrkB signalling pathway, was further confirmed by treatment with recombinant BDNF and co-treatment with GNF-5837 (a Trk inhibitor; Albaugh et al., 2012) and BDNF for 24 hr followed by ELISA (Figure 4g–i). Moreover, treatment with rapamycin as a specific inhibitor of mTORC1 (Jeon et al., 2015; Jeong et al., 2015) induced a significant decrease in CNTF expression in BDNF-treated astrocyte-enriched cultures (Figure 4i). Therefore, these *in vivo* and *in vitro* results suggest that the activation of astrocytic TrkB following an increase in neuronal BDNF in the Rheb(S16H)-treated hippocampus could promote the production of CNTF through the mTORC1-dependent signalling pathway.

3.5 | Contribution of Rheb(S16H)-induced astrocytic CNTF production to neuroprotection against thrombin-induced neurotoxicity in the hippocampus

The results of immunohistochemical staining for CNTFR α at 4 weeks after intra-hippocampal viral injection showed the expression of

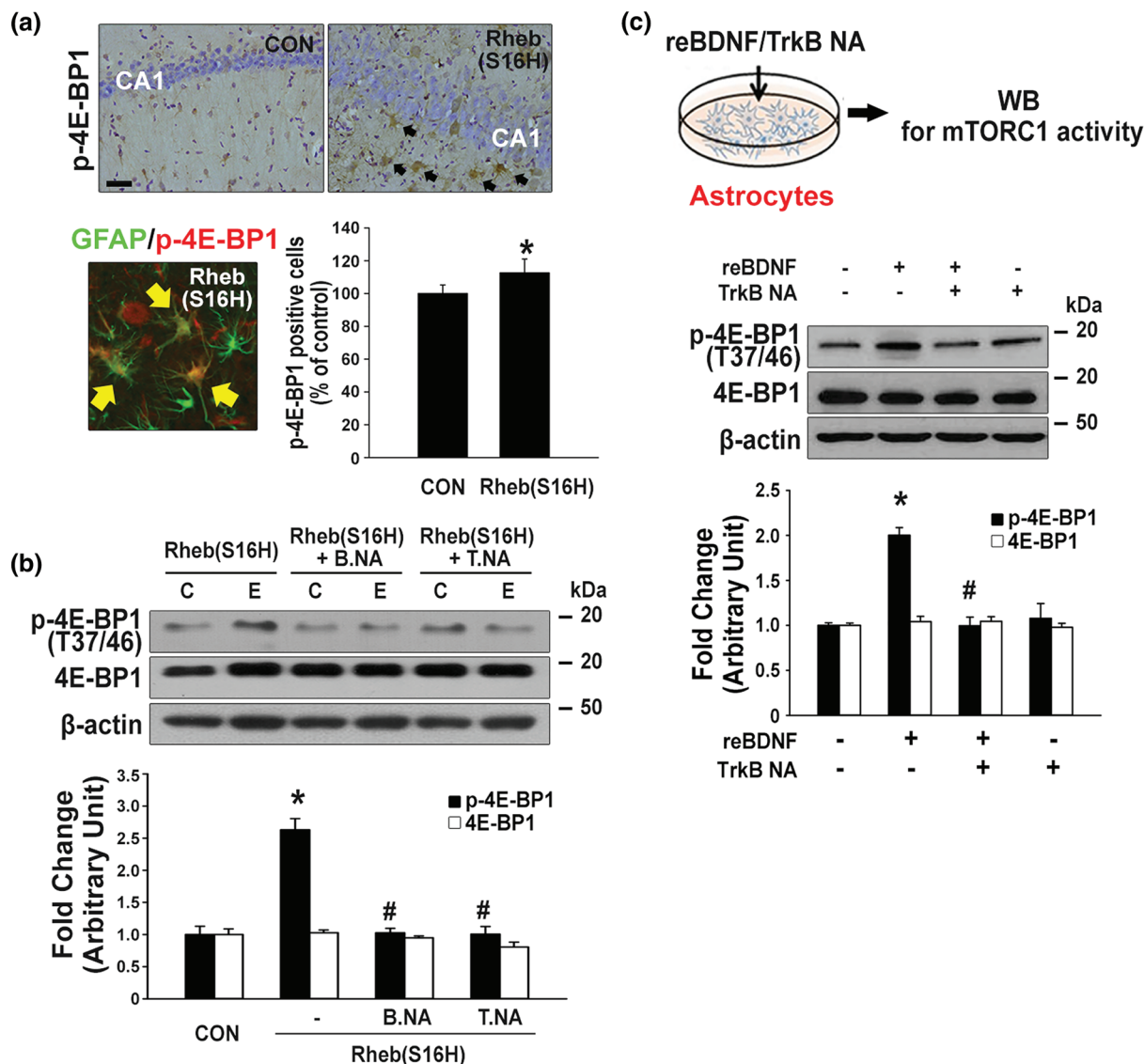


FIGURE 3 Induction of mTORC1 activation in hippocampal astrocytes via increased BDNF following Rheb(S16H) transduction of hippocampal neurons. Hippocampal tissue sections and protein lysates were obtained from the control (CON) or AAV1-Rheb(S16H)-treated rat brain at 4 weeks after injection. (a) Immunohistochemical staining showing increased p-4E-BP1 expression in glia-like cells of the hippocampal CA1 region of AAV1-Rheb(S16H)-injected rats (black arrows). Double immunofluorescence staining against GFAP (green) and p-4E-BP1 (red) showing the colocalization of these two markers in the CA1 region of the hippocampus of AAV1-Rheb(S16H)-injected rats (marked with a yellow arrow). Scale bar, 50 μ m. The bar graph represents the number of p-4E-BP1-positive cells in the CA1 region of the hippocampus. Differences between groups were evaluated by independent samples *t* test. **P* < .05 versus CON. (b) Western blot analysis showing the levels of p-4E-BP1, 4E-BP1, and β -actin in the hippocampus of AAV1-Rheb(S16H)-injected rats, with or without BDNF neutralizing antibody (B.NA) or TrkB neutralizing antibody (T.NA). Differences among groups were evaluated by one-way ANOVA and Tukey's post hoc analysis. **P* < .05 versus CON and #*P* < .05 versus Rheb(S16H) (*n* = 5 for each group). (c) Schematic of the experimental design for western blot analysis of mTORC1 activity after the treatment of hippocampal astrocyte cultures with recombinant BDNF (80 ng·ml⁻¹), with or without T.NA. Differences among groups were evaluated by Kruskal–Wallis test and Tukey's post hoc analysis. **P* < .05 versus CON and #*P* < .05 versus recombinant BDNF alone (*n* = 5 for each group)

CNTFR α in hippocampal neurons *in vivo* and AAV1-Rheb(S16H) transduction increased its expression in hippocampal neurons (Figure 5a). Consistent with the results of immunohistochemical staining, western blotting for CNTFR α showed a significant increase in CNTFR α expression in the Rheb(S16H)-treated hippocampus compared with control and GFP-transduced controls (Figure 5b). Double immunofluorescence staining for NeuN and CNTFR α also showed that increased CNTFR α

following Rheb(S16H) transduction was localized in hippocampal neurons (Figure 5c) but not GFAP-positive astrocytes and OX-42-positive microglia (Figure 5d). These results suggest that AAV1-Rheb(S16H) transduction could intensify the CNTF/CNTFR α neurotrophic signalling pathway in hippocampal neurons *in vivo*.

To verify that the increase in astrocytic CNTF following Rheb(S16H) transduction of hippocampal neurons contributes to

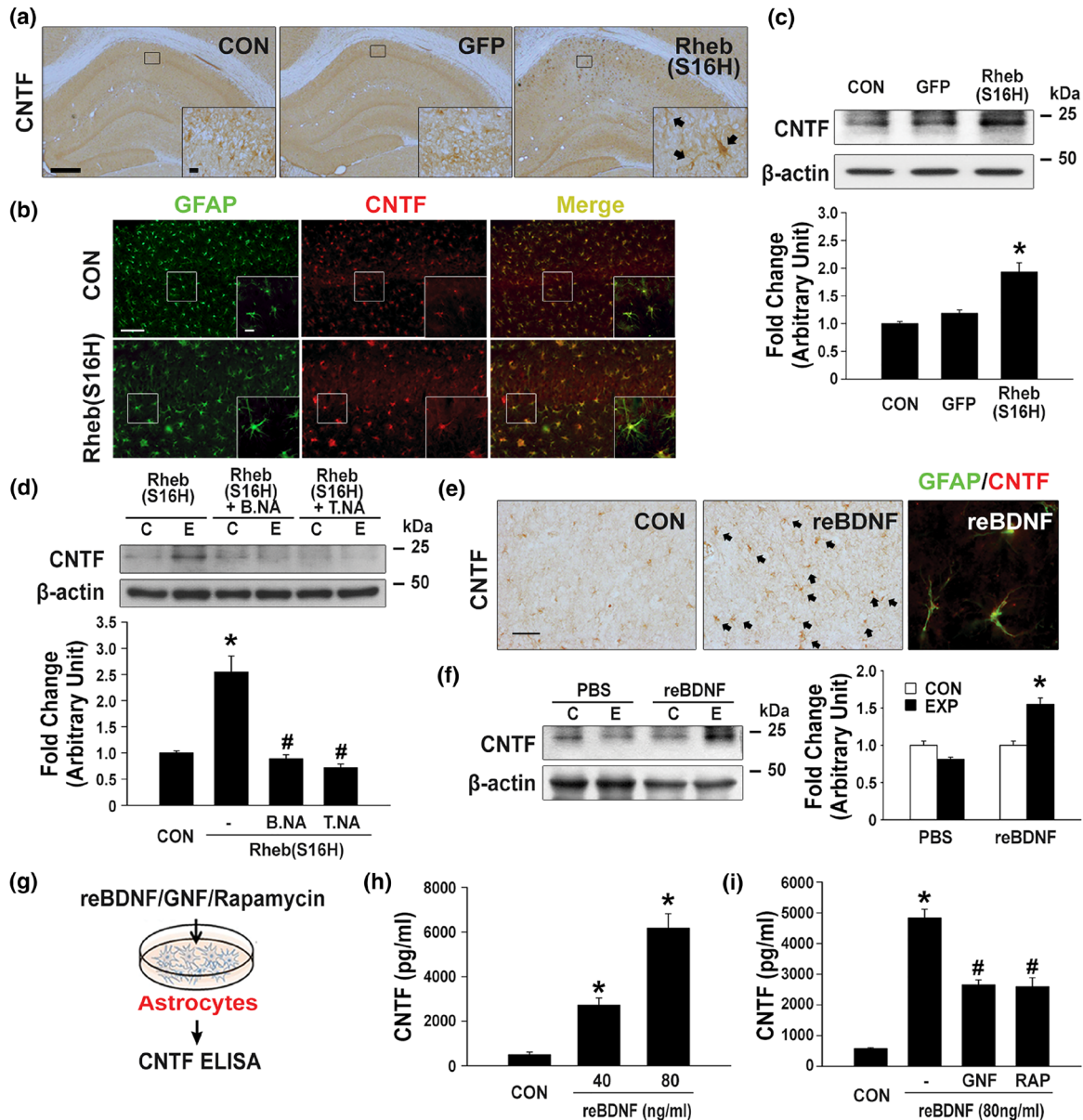


FIGURE 4 Elevation of CNTF production in hippocampal astrocytes via activation of the BDNF/TrkB/mTORC1 signalling pathway. (a) Immunohistochemical staining showing increased CNTF expression in glia-like cells of the hippocampal CA1 region of CON, AAV1-GFP-injected, and AAV1-Rheb(S16H)-injected rats (black arrows). Scale bars, 500 μ m (inset 20 μ m). (b) Double immunofluorescence staining against GFAP (green) and CNTF (red) showing the co-localization of these two markers in the CA1 region of the hippocampus of CON and AAV1-Rheb(S16H)-injected rats. Scale bars, 100 μ m (inset 10 μ m). (c, d) Western blot analysis of the levels of CNTF expression in the hippocampus of AAV1-GFP-injected or AAV1-Rheb(S16H)-injected rats, with or without BDNF neutralizing antibody (B.NA) or TrkB neutralizing antibody (T.NA). Differences among groups were evaluated by Kruskal-Wallis test (c) or one-way ANOVA (d) and Tukey's post hoc analysis. * $P < .05$ versus CON and # $P < .05$ versus Rheb(S16H) alone ($n = 5$ for each group). (e) Immunohistochemical staining showing increased CNTF expression in glia-like cells of the hippocampal CA1 region of recombinant BDNF-injected rats (black arrows). Double immunofluorescence staining against GFAP (green) and CNTF (red) showing the co-localization of these two markers in the CA1 region of the hippocampus of recombinant BDNF-injected rats. Scale bar, 50 μ m. (f) Western blot analysis of CNTF. Differences between groups were evaluated by Student's unpaired t test. * $P < .05$ versus CON ($n = 5$ for each group). (g-i) Measurement of CNTF concentration in the conditioned medium (CM) of recombinant BDNF-treated astrocyte cultures using ELISA kits. (g) Schematic of the experimental design for measuring the levels of CNTF in hippocampal astrocyte cultures. (h) Histogram showing the dose-dependent effects of recombinant BDNF on CNTF release in astrocyte cultures. Differences among groups were evaluated by Kruskal-Wallis test and Tukey's post hoc analysis. * $P < .05$ versus CON ($n = 5$ for each group). (i) Quantitative results showing the CNTF concentration after the treatment of astrocyte cultures with recombinant BDNF (80 $\text{ng}\cdot\text{ml}^{-1}$), with or without GNF-5837 (10 μM) or rapamycin (80 nM). Differences among groups were evaluated by one-way ANOVA and Tukey's post hoc analysis. * $P < .05$ versus CON and # $P < .05$ versus recombinant BDNF alone ($n = 5$ for each group).

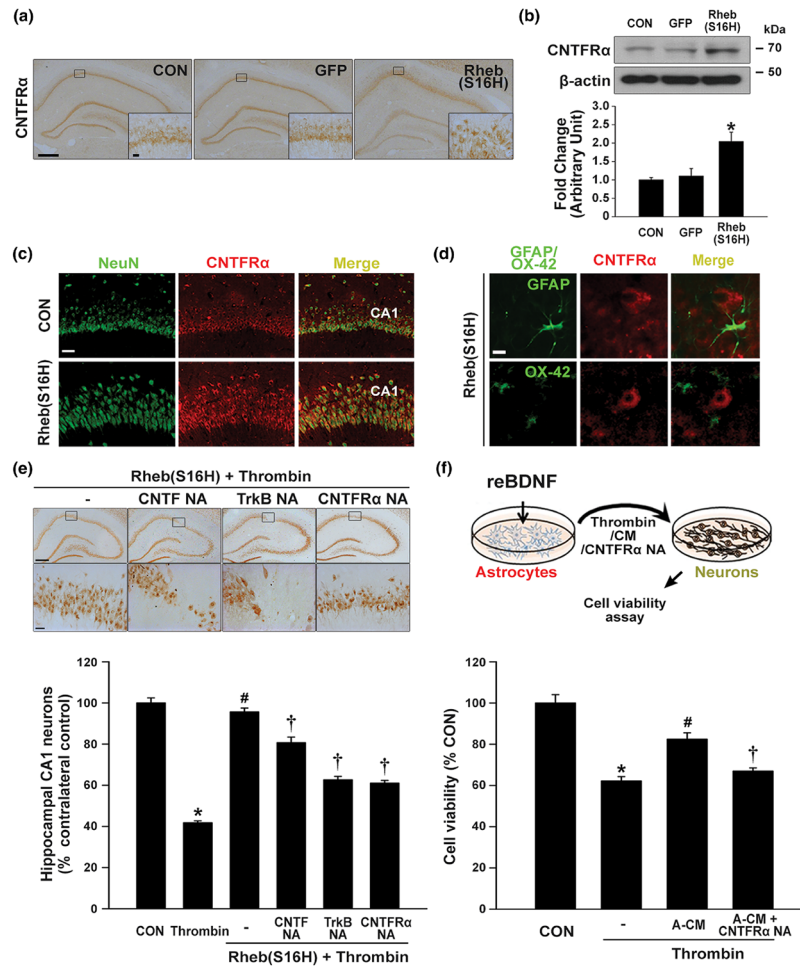


FIGURE 5 Contribution of increased neuronal CNTFRα and astrocytic CNTF to the neuroprotective effects of AAV1-Rheb(S16H) transduction in the hippocampus. (a, b) The expression and localization of CNTFRα in the hippocampus of the rat brain were determined by immunohistochemical staining and western blot analysis. (a) Immunohistochemical staining showing the differences in CNTFRα expression in neuronal cells of the hippocampal CA1 region of CON, AAV1-GFP-injected and AAV1-Rheb(S16H)-injected rats. Scale bars, 500 μm (inset 20 μm). (b) Western blot analysis showing the levels of CNTFRα and β-actin in the hippocampus of CON, AAV1-GFP-injected and AAV1-Rheb(S16H)-injected rats. Differences among groups were evaluated by one-way ANOVA and Tukey's post hoc analysis. **P* < .01 versus CON and GFP (*n* = 5 for each group). (c, d) The expression and localization of NeuN (green) and CNTFRα (red), GFAP (green) and CNTFRα (red) or OX-42 (green) and CNTFRα (red) in the hippocampus of the rat brain were determined by double immunofluorescence staining. (c) Double immunofluorescence staining against NeuN and CNTFRα showing the co-localization of these two markers in the CA1 region of the hippocampus of AAV1-Rheb(S16H)-injected rats. Scale bar, 40 μm. (d) CNTFRα was not expressed in GFAP-positive astrocytes and OX-42-positive microglia in the CA1 region of the hippocampus of AAV1-Rheb(S16H)-injected rats. Scale bar, 10 μm. (e) To assess the neuroprotective effects of AAV1-Rheb(S16H) transduction, NeuN staining was performed to quantify the number of hippocampal neurons. Surviving neurons in the CA1 region of the hippocampus were counted after treatment with AAV1-Rheb(S16H) and thrombin (20 U), with or without neutralizing antibodies (NA; 200 ng) against anti-CNTF, anti-TrkB, or anti-CNTFRα. Scale bars, 500 μm. The number of NeuN-positive hippocampal neurons in the CA1 region was expressed quantitatively as a percentage of the contralateral control. Differences among groups were evaluated by one-way ANOVA and Tukey's post hoc analysis. **P* < .05 versus CON, #*P* < .05 versus thrombin alone and †*P* < .05 versus Rheb(S16H) + thrombin (*n* = 5 for each group). (f) Neuronal cell viability was assessed by MTT assay after 48-hr treatment with thrombin (50 U·ml⁻¹), with or without recombinant BDNF-treated astrocyte-conditioned media (A-CM) or anti-CNTFRα-neutralizing antibody (CNTFRα NA) in primary hippocampal neuron cultures. The histogram shows the MTT assay results expressed as a percentage of the control. Differences among groups were evaluated by one-way ANOVA and Tukey's post hoc analysis. **P* < .05 versus CON, #*P* < .05 versus thrombin alone, and †*P* < .05 versus A-CM + thrombin (*n* = 5 for each group).

neuroprotection in the hippocampus of the adult brain, we determined whether the inhibition of CNTF, CNTFRα and TrkB activities by treatment with specific neutralizing antibodies diminished the Rheb(S16H)-induced neuroprotective effects against thrombin-induced neurotoxicity in the hippocampus of the rat brain (Jeong

et al., 2015). The results of immunohistochemical staining for NeuN showed that the administration of AAV1-Rheb(S16H) in the hippocampus of the rat brain protected NeuN-positive neurons from thrombin-induced neurotoxicity in the counting area of the hippocampal CA1 region by 95% compared with that of the contralateral

controls at 7 days post-lesion (Figure 5e). On the other hand, the inhibition of CNTF, TrkB and CNTFR α activities by intra-hippocampal injection of a mixture of specific neutralizing antibodies and thrombin significantly attenuated the preservation of NeuN-positive neurons by 15%, 33% and 34% respectively, compared with the preserved neurons with Rheb(S16H) treatment alone (Figure 5e;). Treatment with neutralizing antibodies alone did not result in significant changes in the number of NeuN-positive neurons compared with contralateral controls (Figure S6). In addition to the *in vivo* results, we observed that treatment with CM obtained from astrocyte-enriched cultures at 24 hr after recombinant BDNF treatment protected hippocampal neurons against thrombin-induced neurotoxicity *in vitro*, as demonstrated by cell viability analysis at 48 hr post-lesion (Figure 5f). CM-induced neuroprotection was significantly attenuated by co-treatment with neutralizing antibodies against CNTFR α (Figure 5f). Therefore, these results demonstrated that increased astrocytic CNTF following AAV1-Rheb(S16H) administration could contribute to the protection of hippocampal neurons against thrombin-induced neurotoxicity in the hippocampus.

In our previous study (Jeon et al., 2015), we found that Rheb(S16H)-induced neuronal BDNF could contribute to neuroprotection against thrombin-induced neurotoxicity in the hippocampus of the adult brain. However, it was unclear whether the activation of TrkB, expressed in hippocampal neurons is associated with neuroprotection

following Rheb(S16H) transduction. Similar to the results showing TrkB up-regulation in astrocytes (Figure 2h) double immunofluorescence staining for MAP 2, a marker of soma and dendrites (Chung, Jan, & Jan, 2006) and TrkB revealed an increase in TrkB expression in non-neuronal cells (white arrows; Figure 6a). However, TrkB expression was also observed in hippocampal dendrites *in vivo* (Figure 6a). Moreover, in comparison with treatment with thrombin alone, co-treatment with recombinant BDNF and thrombin significantly attenuated thrombin-induced neurotoxicity in hippocampal neuron-enriched cultures (Figure 6b). Taken together, in addition to the neuroprotective effects following activation of the paracrine signalling pathway between hippocampal neurons and astrocytes, Rheb(S16H)-induced neuronal BDNF could protect hippocampal neurons by activating neuronal TrkB, suggesting the activation of an autocrine signalling pathway in the hippocampus of the adult brain (Figure 7).

4 | DISCUSSION

Various studies have demonstrated the therapeutic potential of AAV vectors for treating neurodegeneration in the adult brain with the cell-type-specific transduction of AAVs in the CNS (Chtarto et al., 2013; Fitzsimons & During, 2006; Kim et al., 2011). The beneficial effects of neuronal transduction with AAV in the adult brain were first

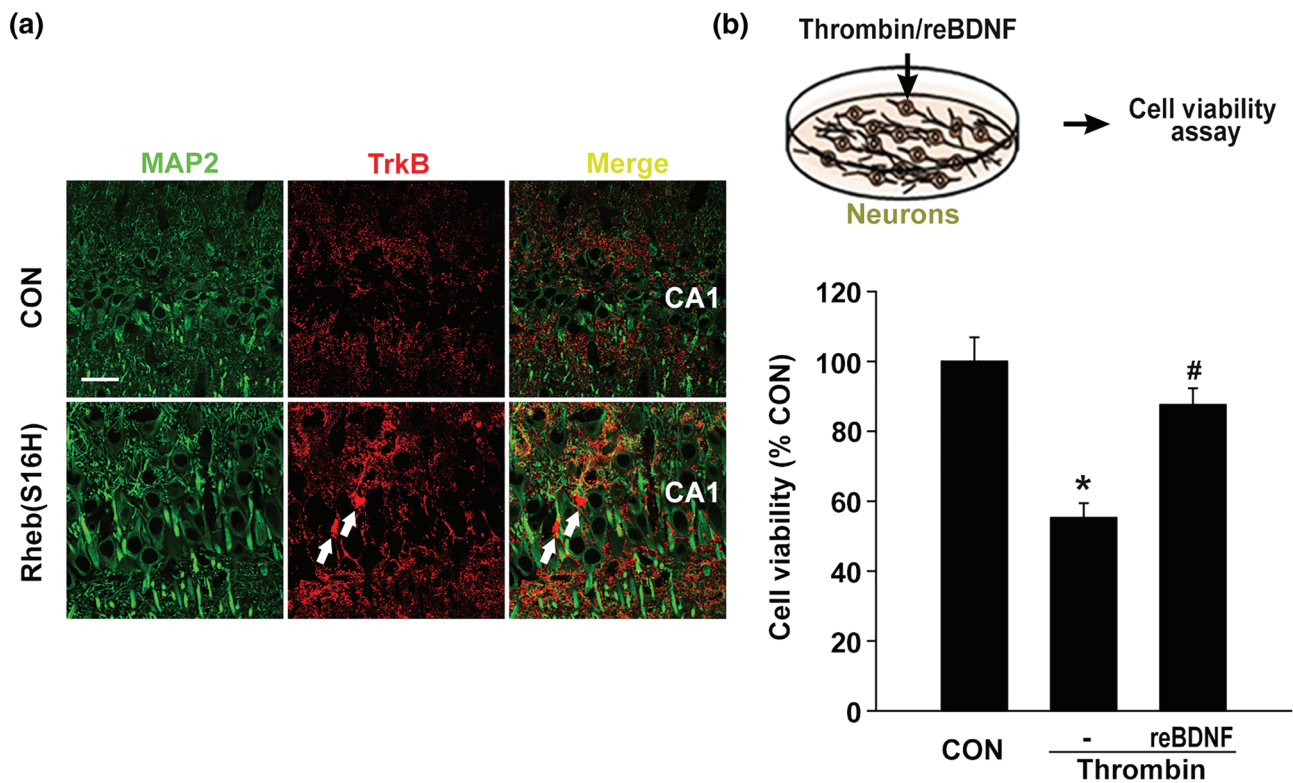


FIGURE 6 Neuroprotective effects of BDNF against thrombin-induced neurotoxicity *in vitro*. (a) Double immunofluorescence staining showing the expression pattern of MAP 2 and TrkB in the hippocampus of AAV1-Rheb(S16H)-injected rats. Scale bar, 40 μ m. White arrows indicate TrkB expression in MAP 2-negative cells. (b) Schematic of the experimental design. Neuronal cell viability was assessed by MTT assay 48 hr after treatment with thrombin (50 U·mL⁻¹) and BDNF. BDNF treatment significantly reduced thrombin-induced neuronal cell death. The histogram shows the results expressed as a percentage of the control ($n = 5$ for each group). Differences among groups were evaluated by one-way ANOVA and Tukey's *post hoc* analysis. * $P < .05$ versus CON and # $P < .05$ versus thrombin alone

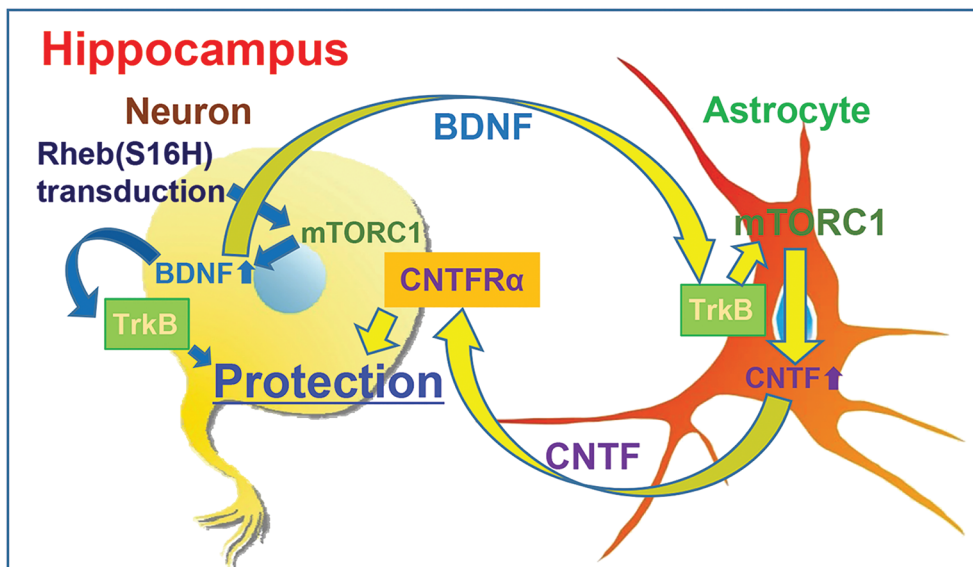


FIGURE 7 Schematic representation of the enhancing neuroprotective system following AAV1-Rheb(S16H) administration. The results suggest that AAV1-Rheb(S16H) transduction of hippocampal neurons could intensify a potential neuroprotective system through functional interactions between neurons and astrocytes triggered by an increase in neuronal BDNF. The intensified system could contribute to neuroprotection via both autocrine and paracrine signalling activation in the hippocampus *in vivo*

reported by Kaplitt et al., 1994, who showed that AAV transduction with a gene encoding TH in the nigral dopaminergic system induced functional recovery in a 6-hydroxydopamine-lesioned animal model of PD (Kaplitt et al., 1994). Transduction with AAV1 in the CNS, such as the hippocampus, SN and striatum, has shown that the expression of a transduced gene is limited to neurons without affecting glial cells such as astrocytes and microglia (Burger et al., 2004; Jeon et al., 2015; Kim et al., 2012), indicating that AAV1 is suitable for the neuronal transduction of a specific gene in the adult brain. In addition, specific gene delivery by AAV1 has been reported to have beneficial effects such as neuroprotection and functional recovery in animal models of neurodegenerative disorders such as PD and AD (Jeon et al., 2015; Jeong et al., 2015; Kim et al., 2011; Kim et al., 2012; Kiyota et al., 2015; Ries et al., 2006). However, although many studies have used AAV1 constructs for therapeutic interventions in the adult brain, it remains unclear whether the production of biomolecules following neuronal transduction with a specific gene using AAV1 stimulates glial activation, which simultaneously supports neuroprotection in the adult brain.

We previously reported that the induction of Rheb(S16H) expression in hippocampal neurons by AAV1 transduction could activate the mTORC1 signalling pathway, resulting in the production of BDNF as a neuroprotective biomolecule against thrombin-induced neurotoxicity in the hippocampus of the rat brain (Jeon et al., 2015). Similar to the beneficial effects in the hippocampus (Jeon et al., 2015), AAV1-Rheb(S16H) transduction in the SN of the murine brain could induce an increase in the protein levels of BDNF, CNTF and glial cell line-derived neurotrophic factor (GDNF) through the activation of mTORC1 in nigral dopaminergic neurons, resulting in neuroprotection in the nigrostriatal dopaminergic system *in vivo* (Jeong et al., 2015;

Kim et al., 2011; Kim et al., 2012). The expression patterns of neurotrophic factors are different between the Rheb(S16H)-treated SN and hippocampus and the mechanism involved remains unclear. However, the findings suggest that the sustained activation of the Rheb/mTORC1 signalling pathway in adult neurons can produce neurotrophic factors in the adult brain *in vivo*. In addition, we found that Rheb(S16H) transduction of hippocampal neurons increased astrocytic CNTF expression in the hippocampus but not hippocampal neurons of the rat brain (Figure 4a,b), suggesting the beneficial activation of astrocytes following the transduction of hippocampal neurons with a specific gene *in vivo*. To further investigate how Rheb(S16H) transduction of hippocampal neurons could induce glial activation in the hippocampus despite the limited expression of a target protein within neurons, we examined whether Rheb(S16H)-induced neuronal BDNF, which was observed previously (Jeon & Nam et al., 2015), stimulates glial activation in the hippocampus of the adult brain and secondly whether reactive astrocytes contribute to Rheb(S16H)-induced neuroprotection in the lesioned hippocampus through interactions between neurons and astrocytes.

Astrocytes are the most abundant cells in the brain and play an important role in the homeostatic maintenance of physiological environments, such as the ion concentration and pH in the CNS (Mulligan & MacVicar, 2004; Sofroniew, 2005). Although previous studies indicated that astrocyte activation is involved in neurodegeneration through inflammatory responses in the adult brain (Jang et al., 2013; Nam et al., 2014), reactive astrocytes can induce beneficial effects such as increasing neuronal survival, growth and activity through barrier function to restrict tissue damage and neuroinflammation (Cekanaviciute et al., 2014; Cekanaviciute et al., 2014; Sofroniew, 2005), also they produce neurotrophic factors such as GDNF and

CNTF in animal models of neurodegenerative diseases (Harada et al., 2002; Harada et al., 2011; Nam et al., 2015). Moreover, a study reported that the loss of astrocytes in the brain of GFAP-knockout mice resulted in a larger infarct area induced by ischaemic brain injury following middle cerebral artery occlusion (Nawashiro, Brenner, Fukui, Shima, & Hallenbeck, 2000). The loss of astrocytic density in the hippocampus may be associated with abnormal phenotypes in an animal model of post-traumatic stress disorder (Saur et al., 2016). Overall, these findings suggest that astrocytes are important for maintaining a normal neural system in the adult brain. Furthermore, the induction of reactive astrocytes, which can protect against neurodegeneration in patients with neurodegenerative diseases such as AD and PD, can support neuronal survival and functional maintenance *in vivo*. In the present study, morphological changes in astrocytes (Figures 2–4) and microglia (Figures 2, S1, and S2) were clearly observed in the Rheb(S16H)-treated hippocampus of the rat brain. However, there was no significant change in the levels of pro-inflammatory cytokines such as IL-1 β and TNF- α , which could induce neurodegeneration in the adult brain (Jang et al., 2013; Nam et al., 2014; Shin et al., 2015) and the Rheb(S16H)-treated hippocampus (Figure S2). Although the role of microglia, which are not involved in the production of BDNF and CNTF in the Rheb(S16H)-treated hippocampus, is unclear, the results suggest that astroglial activation following Rheb(S16H) transduction of hippocampal neurons is important in neuroprotection, but not for inducing neurotoxic inflammation in the adult brain.

The cellular effects of BDNF (a neurotrophin that mediates neuronal survival and differentiation) are initiated by its binding to the specific receptor TrkB expressed in hippocampal neurons resulting in the activation of mTORC1 (Bhave, Ghoda, & Hoffman, 1999; Jeon et al., 2015; Lambert et al., 2013). Increased BDNF expression induces neuroprotective effects, such as anti-neurodegeneration and improved cognitive ability in models of AD *in vivo* and *in vitro* (Jeon et al., 2015; Jeronimo-Santos et al., 2015; Massa et al., 2010; Nagahara et al., 2009). A recent study found that the primary recipient cells of neuronal BDNF are astrocytes in the visual system (Stahlberg, Kügler, & Dean, 2018). Several studies have shown that TrkB-FL and TrkB-T1 are highly expressed on astrocytes (Cheng et al., 2007; Climent, Sancho-Tello, Minana, Baretino, & Guerri, 2000; Cragnolini, Huang, Gokina, & Friedman, 2009; Gabryel & Bernacki, 2009). However, little is known about the role of BDNF in astrocytes. A previous study reported that BDNF delivery stimulated astrocytes to release neuroprotective factors, contributing to neuronal survival in the CNS (Saba et al., 2018), and the production of these neurotrophic factors was attenuated in GFAP-specific TrkB knockout mice (Harada et al., 2011), indicating that the induction of neurotrophic factors in astrocytes could be triggered by activation of the BDNF/TrkB signalling pathway.

CNTF, a neurotrophic cytokine belonging to the IL-6 family, is an important regulator of neuronal survival, neurogenesis and neuronal plasticity in the CNS (Blanchard et al., 2010; Chohan et al., 2011; Jeong et al., 2015; Lipinski et al., 2015; Nam et al., 2015). Its delivery has been found to alleviate cognitive impairments in models of AD *in vivo* and *in vitro* (Garcia et al., 2010; Wang et al., 2015). The effects of CNTF are mediated by the tripartite receptor complex including

leukemia inhibitory factor receptor β , gp130, and CNTFR α (Leibinger & Andreadaki et al., 2012; Jeong & Nam et al., 2015). Especially, CNTFR α is highly expressed in hippocampal neurons (MacLennan & Vinson et al., 1996; Lee & Hofmann et al., 1997). In animal models of PD, we recently found that the induction of astrocytic CNTF prevented the degeneration of dopaminergic neurons *in vivo* (Nam et al., 2015). In addition, the expression of CNTFR α and CNTF was significantly higher in dopaminergic neurons and reactive astrocytes respectively in the post-mortem SN of PD patients compared with age-matched controls (Nam et al., 2015), suggesting that there might be an endogenous neuroprotective system following injury in the SN.

BDNF expression levels have been reported to be decreased in the hippocampus and cortex of patients with AD (Peng, Wu, Mufson, & Fahnstock, 2005; Phillips et al., 1991). However, the increased expression of TrkB-FL, which contains a catalytic kinase domain that can be activated (Armanini, McMahon, Sutherland, Shelton, & Phillips, 1995), was detected in glial cells in the hippocampus of AD patients (Connor et al., 1996; Ferrer et al., 1999), suggesting the presence of a neuroprotective system in the lesioned hippocampus of AD patients, similar to the SN of PD patients. In the present study, the expression of CNTFR α , GFAP and TrkB-FL was significantly higher in the post-mortem hippocampus of AD patients compared with age-matched controls (Figure 1c). However, a decrease in CNTF and BDNF was observed in the hippocampus of AD patients (Figure 1b). These observations suggest that there may also be a compensatory system for neuroprotection in the lesioned hippocampus of AD patients. Although the compensatory system is not sufficient to control neurodegeneration, the induction of astrocytic CNTF and neuronal CNTFR α may be useful approach for strengthening the neuroprotective system, consequently resulting in the protection of hippocampal neurons in the adult brain. However, the current study has some limitations in defining the clinical relevance because measurements were restricted to a limited number of post-mortem human brain samples.

As demonstrated in Figures 2 and 3, Rheb(S16H) transduction of hippocampal neurons induced mTORC1 activation (Figure 3a) and increased the expression levels of both TrkB (Figure 2g–i) and CNTF (Figure 4a–c) in reactive astrocytes. In addition, BDNF was produced in hippocampal neurons *in vivo* (Figure 2b,c) even though transduction with Rheb(S16H) was limited in neurons (Figures 2a,d and S1A; Jeon et al., 2015). To clarify the mechanism of astrocytic CNTF production following AAV1-Rheb(S16H) transduction in the hippocampus, we investigated the effects of neutralizing antibodies against BDNF or TrkB on the production of astrocytic CNTF in the Rheb(S16H)-treated hippocampus *in vivo*. Our results demonstrated that increased astrocytic CNTF following Rheb(S16H) transduction was significantly attenuated by the neutralization of BDNF and TrkB (Figure 4d). Similar to the effects *in vivo*, the production of CNTF induced by treatment with recombinant BDNF was suppressed by inhibiting the TrkB/mTORC1 signalling pathway in rat astrocyte-enriched cultures *in vitro* (Figure 4g–i). In addition to increasing astrocytic CNTF, AAV1-Rheb(S16H) transduction induced a significant increase in CNTFR α expression in hippocampal neurons *in vivo* (Figure 5a–d). Moreover, Rheb(S16H)-induced neuroprotection against thrombin-

induced neurotoxicity in the hippocampus in vivo (Jeon et al., 2015) was inhibited by treatment with neutralizing antibodies against CNTF, TrkB or CNTFR α (Figure 5e) and CNTFR α inhibition in hippocampal neuron-enriched cultures decreased the neuroprotective effect of CM derived from astrocyte-enriched cultures treated with BDNF in vitro (Figure 5f). Therefore, these results indicated that neuronal BDNF produced by AAV1-Rheb(S16H) transduction activated astrocytes and these reactive astrocytes produced CNTF, contributes to neuroprotection in the hippocampus through the activation of TrkB-dependent mTORC1 signalling.

Our previous study demonstrated that neuronal BDNF produced by Rheb(S16H) transduction could protect hippocampal neurons through an autocrine signalling pathway, indicating neuronal TrkB activation in the hippocampus in vivo (Jeon et al., 2015). In the present investigation we have evaluated the neuroprotective effects of BDNF against thrombin-induced neurotoxicity in hippocampal neuron-enriched cultures. Our results showed that treatment with recombinant BDNF could attenuate thrombin-induced neurotoxicity in hippocampal neuron-enriched cultures (Figure 6b). In addition, TrkB expression was observed in hippocampal neurons in vivo (Figure 6a). Therefore, the production of neuronal BDNF following Rheb(S16H) transduction also contributed to neuroprotection through the direct activation of neuronal TrkB in the hippocampus.

This is the first study showing that the expression of a specific target protein induced by neuronal transduction with AAV1 could stimulate astroglial activation, exerting neuroprotective effects in the hippocampus of the adult brain. In conclusion, similar to glial activation in the SN of PD patients (Nam et al., 2015), analysis of the post-mortem hippocampus of AD patients suggests the presence of a neuroprotective system involving an increase in reactive astrocytes, which may protect neurons in the lesioned hippocampus. However, the endogenous of a neuroprotective system may not be sufficient to control neurodegeneration in AD. Our results suggest that viral transduction of hippocampal neurons using methods such as AAV1-Rheb(S16H) administration could intensify a potential neuroprotective system through functional interactions between neurons and astrocytes triggered by an increase in neuronal BDNF, contributing to neuroprotection via both autocrine and paracrine signalling activation in the hippocampus (Figure 7). A study reported that the induction of Rheb might reduce amyloid- β generation (Shahani et al., 2014). Although further studies are needed to determine the clinical relevance of the AAV1-Rheb(S16H) transduction approach for neurodegenerative diseases, this approach may be a useful strategy for protecting hippocampal neurons in the lesioned brain and its effects may be beneficial for patients with AD.

ACKNOWLEDGEMENTS

We thank Dr. Robert E. Burke, who had worked at Columbia University, for his thoughtful comments of the manuscript and results in life. This study was supported by grants from the National Research Foundation of Korea (NRF-2017R1A2B4002675 and NRF-2017R1D1A1B03032218), the Korea Healthcare Technology R&D

Project, Ministry of Health & Welfare (HI15C1928 and HI14C1135), and the KBRI Research Program of the Ministry of Science, ICT & Future Planning (18-BR-02-06, 18-RB-03-03, and 18-BR-03-01) by the Korean government.

AUTHOR CONTRIBUTIONS

M.T.J., G.J.M., S.K., M.C., S.G.L., and S.R.K. conceived and designed the experiments. M.T.J., S.K., G.J.M., Y.S.O., H.J.K., K.J.L., Y.C., W.H.S., S.G.L., and S.R.K. conducted the experiments in vivo, analysed the data, and generated the figures. M.C., W.S.C., I.S.J., M. N, S.G.L., and S.R.K. conducted the experiments in vitro, analysed the data and generated the figures. S.R.K. designed and provided the viral vectors. C.M. provided the human brain samples. G.J.M., D.W.K., C.M.H., and S.R.K. supervised the analysis of the data obtained from human brain samples. S.R.K. supervised the whole project and wrote the paper. All authors contributed to data analysis and preparation of the manuscript.

CONFLICT OF INTEREST

The authors declare no conflicts of interest.

DECLARATION OF TRANSPARENCY AND SCIENTIFIC RIGOUR

This Declaration acknowledges that this paper adheres to the principles for transparent reporting and scientific rigour of preclinical research as stated in the *BJP* guidelines for [Design & Analysis](#), [Immunoblotting and Immunochemistry](#) and [Animal Experimentation](#), and as recommended by funding agencies, publishers and other organizations engaged with supporting research.

AVAILABILITY OF DATA AND MATERIAL

All data generated in this study are included in the article and its supplementary information file.

ORCID

Gyeong Joon Moon  <https://orcid.org/0000-0002-2851-9563>

Yong-Seok Oh  <https://orcid.org/0000-0002-9039-2014>

Dong Woon Kim  <https://orcid.org/0000-0001-7691-1394>

Kea Joo Lee  <https://orcid.org/0000-0002-1366-4709>

Seok-Geun Lee  <https://orcid.org/0000-0002-0060-5777>

Sang Ryong Kim  <https://orcid.org/0000-0003-0299-1613>

REFERENCES

- Alexander, S. P. H., Roberts, R. E., Broughton, B. R. S., Sobey, S. G., George, C. H., Stanford, S. C., ... Ahluwalia, A. (2018). Goals and practicalities of immunoblotting and immunohistochemistry: A guide for submission to the *British Journal of Pharmacology*. *British Journal of Pharmacology*, 175, 407–411. <https://doi.org/10.1111/bph.14112>
- Alexander, S. P., Kelly, E., Mathie, A., Peters, J. A., Veale, E. L., Armstrong, J. F., ... Davies, J. A. (2019). The concise guide to pharmacology 2018/19:

- Introduction and Other Protein Targets. *British Journal of Pharmacology*, 176, S1–S120. <https://doi.org/10.1111/bph.14747>
- Albaugh, P., Fan, Y., Mi, Y., Sun, F., Adrian, F., Li, N., ... Molteni, V. (2012). Discovery of GNF-5837, a selective TRK inhibitor with efficacy in rodent cancer tumor models. *ACS Medicinal Chemistry Letters*, 3(2), 140–145. <https://doi.org/10.1021/ml200261d>
- Albrecht, P. J., Murtie, J. C., Ness, J. K., Redwine, J. M., Enterline, J. R., Armstrong, R. C., & Levison, S. W. (2003). Astrocytes produce CNTF during the remyelination phase of viral-induced spinal cord demyelination to stimulate FGF-2 production. *Neurobiology of Disease*, 13(2), 89–101. [https://doi.org/10.1016/S0969-9961\(03\)00019-6](https://doi.org/10.1016/S0969-9961(03)00019-6)
- Armanini, M. P., McMahon, S. B., Sutherland, J., Shelton, D. L., & Phillips, H. S. (1995). Truncated and catalytic isoforms of trkB are co-expressed in neurons of rat and mouse CNS. *European Journal of Neuroscience*, 7(6), 1403–1409. <https://doi.org/10.1111/j.1460-9568.1995.tb01132.x>
- Bhave, S. V., Ghoda, L., & Hoffman, P. L. (1999). Brain-derived neurotrophic factor mediates the anti-apoptotic effect of NMDA in cerebellar granule neurons: Signal transduction cascades and site of ethanol action. *Journal of Neuroscience*, 19(9), 3277–3286. <https://doi.org/10.1523/jneurosci.19-09-03277.1999>
- Blanchard, J., Chohan, M. O., Li, B., Liu, F., Iqbal, K., & Grundke-Iqbal, I. (2010). Beneficial effect of a CNTF tetrapeptide on adult hippocampal neurogenesis, neuronal plasticity, and spatial memory in mice. *Journal of Alzheimer's Disease*, 21(4), 1185–1195. <https://doi.org/10.3233/JAD-2010-1000069>
- Burger, C., Gorbatyuk, O. S., Velardo, M. J., Peden, C. S., Williams, P., Zolotukhin, S., ... Muzyczka, N. (2004). Recombinant AAV viral vectors pseudotyped with viral capsids from serotypes 1, 2, and 5 display differential efficiency and cell tropism after delivery to different regions of the central nervous system. *Molecular Therapy*, 10(2), 302–317. <https://doi.org/10.1016/j.ymthe.2004.05.024>
- Cecanaviciute, E., Dietrich, H. K., Axtell, R. C., Williams, A. M., Egusquiza, R., Wai, K. M., ... Buckwalter, M. S. (2014). Astrocytic TGF- β signaling limits inflammation and reduces neuronal damage during central nervous system *Toxoplasma* infection. *Journal of Immunology*, 193(1), 139–149. <https://doi.org/10.4049/jimmunol.1303284>
- Cecanaviciute, E., Fathali, N., Doyle, K. P., Williams, A. M., Han, J., & Buckwalter, M. S. (2014). Astrocytic transforming growth factor- β signaling reduces subacute neuroinflammation after stroke in mice. *Glia*, 62(8), 1227–1240. <https://doi.org/10.1002/glia.22675>
- Chauhan, N. B., Siegel, G. J., & Lee, J. M. (2001). Depletion of glial cell line-derived neurotrophic factor in substantia nigra neurons of Parkinson's disease brain. *Journal of Chemical Neuroanatomy*, 21(4), 277–288. [https://doi.org/10.1016/S0891-0618\(01\)00115-6](https://doi.org/10.1016/S0891-0618(01)00115-6)
- Cheng, A., Coksaygan, T., Tang, H., Khatri, R., Balice-Gordon, R. J., Rao, M. S., & Mattson, M. P. (2007). Truncated tyrosine kinase B brain-derived neurotrophic factor receptor directs cortical neural stem cells to a glial cell fate by a novel signaling mechanism. *Journal of Neurochemistry*, 100(6), 1515–1530. <https://doi.org/10.1111/j.1471-4159.2006.04337.x>
- Chohan, M. O., Li, B., Blanchard, J., Tung, Y. C., Heaney, A. T., Rabe, A., ... Grundke-Iqbal, I. (2011). Enhancement of dentate gyrus neurogenesis, dendritic and synaptic plasticity and memory by a neurotrophic peptide. *Neurobiology of Aging*, 32(8), 1420–1434. <https://doi.org/10.1016/j.neurobiolaging.2009.08.008>
- Chtarto, A., Bockstael, O., Tshibangu, T., Dewitte, O., Levivier, M., & Tenenbaum, L. (2013). A next step in adeno-associated virus-mediated gene therapy for neurological diseases: Regulation and targeting. *British Journal of Clinical Pharmacology*, 76(2), 217–232. <https://doi.org/10.1111/bcp.12065>
- Chung, H. J., Jan, Y. N., & Jan, L. Y. (2006). Polarized axonal surface expression of neuronal KCNQ channels is mediated by multiple signals in the KCNQ2 and KCNQ3 C-terminal domains. *Proceedings of the National Academy of Sciences of the United States of America*, 103(23), 8870–8875. <https://doi.org/10.1073/pnas.0603376103>
- Cilient, E., Sancho-Tello, M., Minana, R., Baretino, D., & Guerri, C. (2000). Astrocytes in culture express the full-length Trk-B receptor and respond to brain derived neurotrophic factor by changing intracellular calcium levels: Effect of ethanol exposure in rats. *Neuroscience Letters*, 288(1), 53–56. [https://doi.org/10.1016/S0304-3940\(00\)01207-6](https://doi.org/10.1016/S0304-3940(00)01207-6)
- Connor, B., Young, D., Lawlor, P., Gai, W., Waldvogel, H., Faull, R. L., & Dragunow, M. (1996). Trk receptor alterations in Alzheimer's disease. *Molecular Brain Research*, 42(1), 1–17. [https://doi.org/10.1016/S0169-328x\(96\)00040-x](https://doi.org/10.1016/S0169-328x(96)00040-x)
- Cragolini, A. B., Huang, Y., Gokina, P., & Friedman, W. J. (2009). Nerve growth factor attenuates proliferation of astrocytes via the p75 neurotrophin receptor. *Glia*, 57(13), 1386–1392. <https://doi.org/10.1002/glia.20857>
- Curtis, M. J., Bond, R. A., Spina, D., Ahluwalia, A., Alexander, S. P., Giembycz, M. A., ... McGrath, J. C. (2015). Experimental design and analysis and their reporting: New guidance for publication in *BJP*. *British Journal of Pharmacology*, 172(14), 3461–3471. <https://doi.org/10.1111/bph.12856>
- Ferrer, I., Marin, C., Rey, M. J., Ribalta, T., Goutan, E., Blanco, R., ... Marti, E. (1999). BDNF and full-length and truncated TrkB expression in Alzheimer disease. Implications in therapeutic strategies. *Journal of Neuropathology and Experimental Neurology*, 58(7), 729–739. <https://doi.org/10.1097/00005072-199907000-00007>
- Fitzsimons, H. L., & During, M. J. (2006). Design and optimization of expression cassettes including promoter choice and regulatory elements. In *Gene therapy of the central nervous system* (pp. 3–16). Amsterdam: Academic Press.
- Gabryel, B., & Bernacki, J. (2009). Effect of FK506 and cyclosporine A on the expression of BDNF, tyrosine kinase B and p75 neurotrophin receptors in astrocytes exposed to simulated ischemia in vitro. *Cell Biology International*, 33(7), 739–748. <https://doi.org/10.1016/j.cellbi.2009.04.006>
- Garcia, P., Youssef, I., Utvik, J. K., Florent-Bechard, S., Barthelemy, V., Malaplate-Armand, C., ... Pillot, T. (2010). Ciliary neurotrophic factor cell-based delivery prevents synaptic impairment and improves memory in mouse models of Alzheimer's disease. *Journal of Neuroscience*, 30(22), 7516–7527. <https://doi.org/10.1523/JNEUROSCI.4182-09.2010>
- Harada, C., Guo, X., Namekata, K., Kimura, A., Nakamura, K., Tanaka, K., ... Harada, T. (2011). Glia- and neuron-specific functions of TrkB signalling during retinal degeneration and regeneration. *Nature Communications*, 2, 189. <https://doi.org/10.1038/ncomms1190>
- Harada, T., Harada, C., Kohsaka, S., Wada, E., Yoshida, K., Ohno, S., ... Wada, K. (2002). Microglia-Muller glia cell interactions control neurotrophic factor production during light-induced retinal degeneration. *Journal of Neuroscience*, 22(21), 9228–9236. <https://doi.org/10.1523/JNEUROSCI.22-21-09228.2002>
- Harding, S. D., Sharman, J. L., Faccenda, E., Southan, C., Pawson, A. J., Ireland, S., et al. (2018). The IUPHAR/BPS guide to pharmacology in 2018: Updates and expansion to encompass the new guide to immunopharmacology. *Nucl Acids Res*, 46, D1091–D1106.
- Huang, E. J., & Reichardt, L. F. (2001). Neurotrophins: Roles in neuronal development and function. *Annual Review of Neuroscience*, 24, 677–736. <https://doi.org/10.1146/annurev.neuro.24.1.677>
- Jang, E., Kim, J. H., Lee, S., Kim, J. H., Seo, J. W., Jin, M., ... Suk, K. (2013). Phenotypic polarization of activated astrocytes: The critical role of

- lipocalin-2 in the classical inflammatory activation of astrocytes. *Journal of Immunology*, 191(10), 5204–5219. <https://doi.org/10.4049/jimmunol.1301637>
- Jeon, M. T., Nam, J. H., Shin, W. H., Leem, E., Jeong, K. H., Jung, U. J., ... Kim, S. R. (2015). In vivo AAV1 transduction with hRheb(S16H) protects hippocampal neurons by BDNF production. *Molecular Therapy*, 23(3), 445–455. <https://doi.org/10.1038/mt.2014.241>
- Jeong, K. H., Nam, J. H., Jin, B. K., & Kim, S. R. (2015). Activation of CNTF/CNTFR α signaling pathway by hRheb(S16H) transduction of dopaminergic neurons in vivo. *PLoS ONE*, 10(3), e0121803. <https://doi.org/10.1371/journal.pone.0121803>
- Jeronimo-Santos, A., Vaz, S. H., Parreira, S., Rapaz-Lerias, S., Caetano, A. P., Buee-Scherrer, V., ... Diogenes, M. J. (2015). Dysregulation of TrkB receptors and BDNF function by amyloid- β peptide is mediated by calpain. *Cerebral Cortex*, 25(9), 3107–3121. <https://doi.org/10.1093/cercor/bhu105>
- Kaech, S., & Banker, G. (2006). Culturing hippocampal neurons. *Nature Protocols*, 1(5), 2406–2415. <https://doi.org/10.1038/nprot.2006.356>
- Kaplitt, M. G., Leone, P., Samulski, R. J., Xiao, X., Pfaff, D. W., O'Malley, K. L., & During, M. J. (1994). Long-term gene expression and phenotypic correction using adeno-associated virus vectors in the mammalian brain. *Nature Genetics*, 8(2), 148–154. <https://doi.org/10.1038/ng1094-148>
- Kilkenny, C., Browne, W., Cuthill, I. C., Emerson, M., Altman, D. G., & Group NCRGW (2010). Animal research: Reporting in vivo experiments: The ARRIVE guidelines. *British Journal of Pharmacology*, 160(7), 1577–1579. <https://doi.org/10.1111/j.1476-5381.2010.00872.x>
- Kim, S., Moon, G. J., Oh, Y. S., Park, J., Shin, W. H., Jeong, J. Y., ... Kim, S. R. (2018). Protection of nigral dopaminergic neurons by AAV1 transduction with Rheb(S16H) against neurotoxic inflammation in vivo. *Experimental and Molecular Medicine*, 50(2), e440. <https://doi.org/10.1038/emm.2017.261>
- Kim, S. R., Chen, X., Oo, T. F., Kareva, T., Yarygina, O., Wang, C., ... Burke, R. E. (2011). Dopaminergic pathway reconstruction by Akt/Rheb-induced axon regeneration. *Annals of Neurology*, 70(1), 110–120. <https://doi.org/10.1002/ana.22383>
- Kim, S. R., Kareva, T., Yarygina, O., Kholodilov, N., & Burke, R. E. (2012). AAV transduction of dopamine neurons with constitutively active Rheb protects from neurodegeneration and mediates axon regrowth. *Molecular Therapy*, 20(2), 275–286. <https://doi.org/10.1038/mt.2011.213>
- Kiyota, T., Zhang, G., Morrison, C. M., Bosch, M. E., Weir, R. A., Lu, Y., ... Gendelman, H. E. (2015). AAV2/1 CD74 gene transfer reduces β -amyloidosis and improves learning and memory in a mouse model of Alzheimer's disease. *Molecular Therapy*, 23(11), 1712–1721. <https://doi.org/10.1038/mt.2015.142>
- Lambert, W. M., Xu, C. F., Neubert, T. A., Chao, M. V., Garabedian, M. J., & Jeanneteau, F. D. (2013). Brain-derived neurotrophic factor signaling rewrites the glucocorticoid transcriptome via glucocorticoid receptor phosphorylation. *Molecular and Cellular Biology*, 33(18), 3700–3714. <https://doi.org/10.1128/MCB.00150-13>
- Lee, M. Y., Hofmann, H. D., & Kirsch, M. (1997). Expression of ciliary neurotrophic factor receptor- α messenger RNA in neonatal and adult rat brain: An in situ hybridization study. *Neuroscience*, 77(1), 233–246. [https://doi.org/10.1016/s0306-4522\(96\)00476-9](https://doi.org/10.1016/s0306-4522(96)00476-9)
- Leibinger, M., Andreadaki, A., & Fischer, D. (2012). Role of mTOR in neuroprotection and axon regeneration after inflammatory stimulation. *Neurobiology of Disease*, 46(2), 314–324. <https://doi.org/10.1016/j.nbd.2012.01.004>
- Lipinski, D. M., Barnard, A. R., Singh, M. S., Martin, C., Lee, E. J., Davies, W. I. L., & MacLaren, R. E. (2015). CNTF gene therapy confers lifelong neuroprotection in a mouse model of human retinitis pigmentosa. *Molecular Therapy*, 23(8), 1308–1319. <https://doi.org/10.1038/mt.2015.68>
- MacLennan, A. J., Vinson, E. N., Marks, L., McLaurin, D. L., Pfeifer, M., & Lee, N. (1996). Immunohistochemical localization of ciliary neurotrophic factor receptor α expression in the rat nervous system. *Journal of Neuroscience*, 16(2), 621–630. <https://doi.org/10.1523/jneurosci.16-02-00621.1996>
- Massa, S. M., Yang, T., Xie, Y., Shi, J., Bilgen, M., Joyce, J. N., ... Longo, F. M. (2010). Small molecule BDNF mimetics activate TrkB signaling and prevent neuronal degeneration in rodents. *Journal of Clinical Investigation*, 120(5), 1774–1785. <https://doi.org/10.1172/JCI41356>
- Mulligan, S. J., & MacVicar, B. A. (2004). Calcium transients in astrocyte endfeet cause cerebrovascular constrictions. *Nature*, 431(7005), 195–199. <https://doi.org/10.1038/nature02827>
- Nagahara, A. H., Merrill, D. A., Coppola, G., Tsukada, S., Schroeder, B. E., Shaked, G. M., ... Tuszynski, M. H. (2009). Neuroprotective effects of brain-derived neurotrophic factor in rodent and primate models of Alzheimer's disease. *Nature Medicine*, 15(3), 331–337. <https://doi.org/10.1038/nm.1912>
- Nam, J. H., Park, E. S., Won, S. Y., Lee, Y. A., Kim, K. I., Jeong, J. Y., ... Jin, B. K. (2015). TRPV1 on astrocytes rescues nigral dopamine neurons in Parkinson's disease via CNTF. *Brain*, 138(Pt 12), 3610–3622. <https://doi.org/10.1093/brain/awv297>
- Nam, Y., Kim, J. H., Seo, M., Kim, J. H., Jin, M., Jeon, S., ... Suk, K. (2014). Lipocalin-2 protein deficiency ameliorates experimental autoimmune encephalomyelitis: The pathogenic role of lipocalin-2 in the central nervous system and peripheral lymphoid tissues. *Journal of Biological Chemistry*, 289(24), 16773–16789. <https://doi.org/10.1074/jbc.M113.542282>
- Nawashiro, H., Brenner, M., Fukui, S., Shima, K., & Hallenbeck, J. M. (2000). High susceptibility to cerebral ischemia in GFAP-null mice. *Journal of Cerebral Blood Flow and Metabolism*, 20(7), 1040–1044. <https://doi.org/10.1097/00004647-200007000-00003>
- Paxinos, G., & Watson, C. (2007). *The rat brain in stereotaxic coordinates* (6th ed.). Amsterdam, Boston: Academic Press/Elsevier.
- Peng, S., Wu, J., Mufson, E. J., & Fahnstock, M. (2005). Precursor form of brain-derived neurotrophic factor and mature brain-derived neurotrophic factor are decreased in the pre-clinical stages of Alzheimer's disease. *Journal of Neurochemistry*, 93(6), 1412–1421. <https://doi.org/10.1111/j.1471-4159.2005.03135.x>
- Phillips, H. S., Hains, J. M., Armanini, M., Laramée, G. R., Johnson, S. A., & Winslow, J. W. (1991). BDNF mRNA is decreased in the hippocampus of individuals with Alzheimer's disease. *Neuron*, 7(5), 695–702. [https://doi.org/10.1016/0896-6273\(91\)90273-3](https://doi.org/10.1016/0896-6273(91)90273-3)
- Ries, V., Henchcliffe, C., Kareva, T., Rzhetskaya, M., Bland, R., During, M. J., ... Burke, R. E. (2006). Oncoprotein Akt/PKB induces trophic effects in murine models of Parkinson's disease. *Proceedings of the National Academy of Sciences of the United States of America*, 103(49), 18757–18762. <https://doi.org/10.1073/pnas.0606401103>
- Saba, J., Turati, J., Ramirez, D., Carniglia, L., Durand, D., Lasaga, M., & Caruso, C. (2018). Astrocyte truncated tropomyosin receptor kinase B mediates brain-derived neurotrophic factor anti-apoptotic effect leading to neuroprotection. *Journal of Neurochemistry*, 146(6), 686–702. <https://doi.org/10.1111/jnc.14476>
- Saur, L., Baptista, P. P., Bagatini, P. B., Neves, L. T., de Oliveira, R. M., Vaz, S. P., ... Xavier, L. L. (2016). Experimental post-traumatic stress disorder decreases astrocyte density and changes astrocytic polarity in the CA1 hippocampus of male rats. *Neurochemical Research*, 41(4), 892–904. <https://doi.org/10.1007/s11064-015-1770-3>
- Shahani, N., Pryor, W., Swarnkar, S., Kholodilov, N., Thinakaran, G., Burke, R. E., & Subramaniam, S. (2014). Rheb GTPase regulates β -secretase levels and amyloid β generation. *Journal of Biological Chemistry*, 289(9), 5799–5808. <https://doi.org/10.1074/jbc.M113.532713>

- Shin, W. H., Jeon, M. T., Leem, E., Won, S. Y., Jeong, K. H., Park, S. J., ... Kim, S. R. (2015). Induction of microglial toll-like receptor 4 by prothrombin kringle-2: A potential pathogenic mechanism in Parkinson's disease. *Scientific Reports*, 5, 14764. <https://doi.org/10.1038/srep14764>
- Siegel, G. J., & Chauhan, N. B. (2000). Neurotrophic factors in Alzheimer's and Parkinson's disease brain. *Brain Research Reviews*, 33(2-3), 199–227. [https://doi.org/10.1016/S0165-0173\(00\)00030-8](https://doi.org/10.1016/S0165-0173(00)00030-8)
- Smith, E. D., Prieto, G. A., Tong, L., Sears-Kraxberger, I., Rice, J. D., Steward, O., & Cotman, C. W. (2014). Rapamycin and interleukin-1 β impair brain-derived neurotrophic factor-dependent neuron survival by modulating autophagy. *Journal of Biological Chemistry*, 289(30), 20615–20629. <https://doi.org/10.1074/jbc.M114.568659>
- Sofroniew, M. V. (2005). Reactive astrocytes in neural repair and protection. *The Neuroscientist*, 11(5), 400–407. <https://doi.org/10.1177/1073858405278321>
- Sopova, K., Gatsiou, K., Stellos, K., & Laske, C. (2014). Dysregulation of neurotrophic and haematopoietic growth factors in Alzheimer's disease: From pathophysiology to novel treatment strategies. *Current Alzheimer Research*, 11(1), 27–39. <https://doi.org/10.2174/1567205010666131120100743>
- Stahlberg, M. A., Kügler, S., & Dean, C. (2018). Visualizing BDNF Cell-to-Cell Transfer Reveals Astrocytes Are the Primary Recipient of Neuronal BDNF. *bioRxiv*, 255935. <https://doi.org/10.1101/255935>
- Vidaurre, O. G., Gascon, S., Deogracias, R., Sobrado, M., Cuadrado, E., Montaner, J., ... Diaz-Guerra, M. (2012). Imbalance of neurotrophin receptor isoforms TrkB-FL/TrkB-T1 induces neuronal death in excitotoxicity. *Cell Death and Disease*, 3, e256. <https://doi.org/10.1038/cddis.2011.143>
- Wang, K., Xie, M., Zhu, L., Zhu, X., Zhang, K., & Zhou, F. (2015). Ciliary neurotrophic factor protects SH-SY5Y neuroblastoma cells against Abeta1-42-induced neurotoxicity via activating the JAK2/STAT3 axis. *Folia Neuropathologica*, 53(3), 226–235. <https://doi.org/10.5114/fn.2015.54423>

SUPPORTING INFORMATION

Additional supporting information may be found online in the Supporting Information section at the end of the article.

How to cite this article: Jeon M-T, Moon GJ, Kim S, et al. Neurotrophic interactions between neurons and astrocytes following AAV1-Rheb(S16H) transduction in the hippocampus in vivo. *Br J Pharmacol*. 2020;177:668–686. <https://doi.org/10.1111/bph.14882>