

Won-Jong Oh, PhD

Principal Investigator

NEUROVASCULAR UNIT GROUP
Korea Brain Research Institute (KBRI)

Office : 5-3

Lab : wet lab 5-2

Tel : +82-53-980-8360

Fax : +82-53-980-8339

E-mail : ohwj@kbri.re.kr

<http://sites.google.com/site/neurovascularbiologylab/home>

Molecular mechanism of neurovascular unit and blood-brain barrier

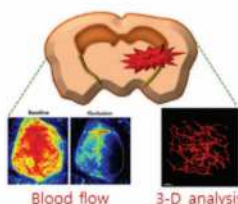
Evolution has crafted vertebrates with two exquisitely elaborate systems each specialized to perform their unique and vital functions: the nervous and vascular systems. These two systems are wired intimately to perform many physiological processes such as energy metabolism and hemodynamic control. However, in contrast to our advanced understanding of neural circuit assembly, how the architecture of the neurovascular system is wired and subsequently establishes the proper functional unit in the mammalian brain is still poorly understood. In addition, it has been widely highlighted that the impairments of the neurovascular structure and functions are crucial etiological factors leading the neuropathological conditions such as Alzheimer's disease, Parkinson's disease, and vascular dementia. Using a combination of advanced mouse genetic, histological, molecular and cellular approaches, we are currently investigating basic principles of brain neurovascular development and disease pathogenesis and will facilitate the development of a therapeutic strategy for curing neurological disorders.

Aim Understanding neurovascular unit formation and functional maintenance

Tool *In vivo* mouse model + Molecular histology + Brain vasculature physiology

T
A
R
G
E
T

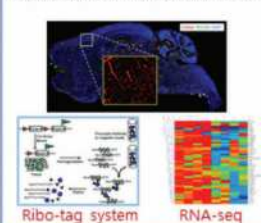
Vascular damage and remodeling



Vascular dementia



Neurovascular interaction factor



Curriculum Vitae

2014~Present : Principal Investigator, KBRI
 2007~2014 : Research Fellow, Harvard Medical School, USA
 1996~1999 : Researcher, Samsung Biomedical Research
 Institute, Korea

Academic Credential

2006 : Ph.D., Molecular Medicine, Medical College of
 Georgia, USA
 1995 : M.S., Molecular Biology, Chung-Ang Univ., Korea
 1993 : B.S., Biology, Chung-Ang Univ.

Awards/Honors/Memberships

2002~Present : Member, Society for Neuroscience
 2015~Present : Member (Academic committee), Korean Society for
 Vascular Biology and Medicine
 2014~Present : Member, The Korean Society for Brain and Neural Sciences
 2010~2011 : Alice and Joseph Brooks Fund Postdoctoral Fellowship, USA
 2008~2010 : Lefler Fellowship, USA
 2006 : Ph.D. with Distinction, Medical College of Georgia, USA

Research keywords

Brain Vasculature, Neurovascular Unit, Blood-Brain Barrier, Vascular Dementia.

Key techniques

Genetics- and surgery-based model mouse development, Molecular Histology, Blood flow and permeability analysis.

Research Interests/Topics

- Mechanism about establishment and maintenance of the neurovascular unit, and screening of novel neurovascular interaction factors.
- Basic mechanism of neuronal disorders by vascular impairment and therapeutic strategy.

Research Publications (selected)

- Choi MG, Kim MJ, Kim DG, Yu R, Jang YN, **Oh WJ**. Sequestration of synaptic proteins by alpha-synuclein aggregates leading to neurotoxicity is inhibited by small peptide. *Plos One*, 13(4):e0195339, 2018.
- **Oh WJ**, Gu C. Establishment of neurovascular congruency in the mouse whisker system by an independent patterning mechanism. *Neuron*, 80: 458-469, 2013.
- Ding JB*, **Oh WJ***, Sabatini BL, Gu C. Semaphorin3E-Plexin-D1 signaling controls pathway-specific synapse formation in the striatum. *Nature Neuroscience*, 15:215-223, 2012. (*equal contribution)
- Kim JH*, **Oh WJ***, Gaiano N, Yoshida Y, Gu C. Semaphorin3E-Plexin-D1 signaling regulates VEGF function in developmental angiogenesis via a feedback mechanism. *Genes and Development*, 25:1399-1411, 2011. (*equal contribution)

Patents (selected)

- Choi MG and **Oh WJ**. A method for suppressing alpha-Synuclein aggregates-mediated cellular toxicity. (10-1896182, patent registered)