

Development of a Method to Treat Alzheimer's Disease by Protection of Synapses

Tech ID: 30459 / UC Case 2019-112-0

BACKGROUND

NMDA receptors (NMDARs) are principal regulators of synaptic signaling in the brain. Modulation of NMDARs' function and trafficking is important for the regulation of synaptic transmission and several forms of synaptic plasticity. Postsynaptic density protein 95 (PSD-95) acts as a scaffolding protein and stabilizes the surface and synaptic expression of NMDARs. NMDA receptors (NMDARs) are ionotropic glutamate receptors that are expressed throughout the nervous system and play crucial roles in neuronal development, synaptic plasticity, learning and memory.

PSD-95 (Post Synaptic Density protein) or SAP90, a membrane-associated guanylate kinase (MAGUK), is the major scaffolding protein in the excitatory postsynaptic density (PSD) and a potent regulator of synaptic strength. It is almost exclusively located in the post synaptic density of neurons and is involved in anchoring synaptic proteins. Its direct and indirect binding partners include neuroligin, NMDA receptors, AMPA receptors, and potassium channels. Postsynaptic loss does not precede obvious A β (beta-amyloid or amyloid beta) and Tau deposition, but instead appears to occur as A β and Tau pathologies advance. This indicates that PSD-95 is an excellent intrinsic biomarker for post synaptic mechanisms and its expression is reduced in brain tissue from patients with Alzheimer's Disease (AD) as well as in mouse models of AD.

TECHNOLOGY DESCRIPTION

Researchers at UC San Diego have shown that increased PSD-95, a major synaptic scaffolding molecule, blocks the effects of A β on synapses. These results indicate that increased PSD-95 protects synapses from A β toxicity by interfering with ion-flux independent NMDAR CTD signaling. Using a chemical inhibitor to block its depalmitoylation increases PSD-95 at synapses and rescues deficits caused by A β ; possibly opening a new therapeutic avenue against AD.

APPLICATIONS

The inventors have developed a method for treating a neurodegenerative disease by administering a therapeutically effective amount of a polypeptide or an effective amount of an inhibitor of a palmitoylation pathway to increase the amount of PSD-95 in the synapse.

STATE OF DEVELOPMENT

The state of the development is in the experimental stage.

INTELLECTUAL PROPERTY INFO

This technology is patent pending and available for licensing and/or research sponsorship.

PATENT STATUS

Patent Pending

CONTACT

University of California, San Diego
Office of Innovation and Commercialization
innovation@ucsd.edu
tel: 858.534.5815.



OTHER INFORMATION

KEYWORDS

Post Synaptic Density protein, PSD-95, neurodegenerative diseases, NMDA receptors, inhibitors of palmitoylation, Alzheimer's disease

CATEGORIZED AS

- **Medical**
- Other
- Therapeutics

RELATED CASES

2019-112-0

University of California, San Diego
Office of Innovation and Commercialization
9500 Gilman Drive, MC 0910, ,
La Jolla,CA 92093-0910

Tel: 858.534.5815
innovation@ucsd.edu
<https://innovation.ucsd.edu>
Fax: 858.534.7345

© 2019, The Regents of the
University of California
[Terms of use](#)
[Privacy Notice](#)