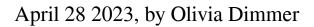
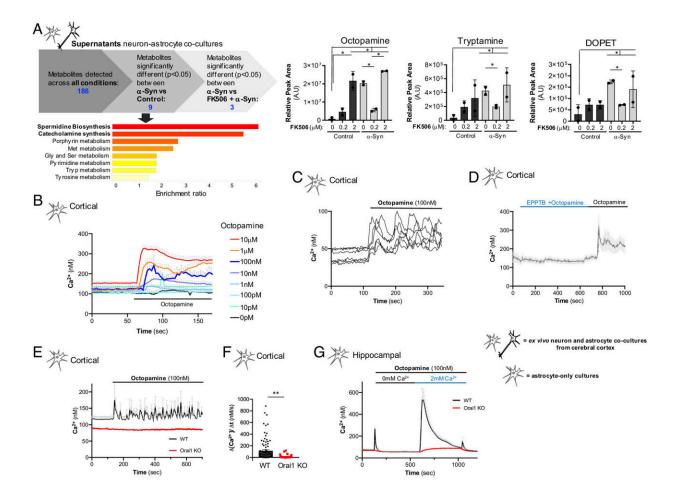


Understanding the role of octopamine in neurodegeneration





Octopamine is a calcineurin-dependent metabolite released under α -syn proteotoxicity that mediates Ca²⁺ influx in astrocytes. (*A*) Metabolite screen from supernatants of rat primary cortical neuron–astrocyte cocultures transduced with either control lentivirus or α -synA53T driven by the synapsin promoter and treated with either vehicle (DMSO) or subsaturating (0.2 μ M) doses of FK506 5 to 6 wk posttransduction. Metabolites that were significantly different (*P* P B)



Ca²⁺ imaging of primary rat cortical astrocyte-only cultures treated with a range of octopamine concentrations (0 to 10 μ M). N = 3; 7 to 10 cells/biological replicate. (C) Representative single-cell Ca^{2+} imaging traces of primary rat cortical astrocyte-only cultures treated with 100 nM octopamine (N = 6). (D) Ca²⁺ imaging of primary rat cortical astrocyte-only cultures cotreated with octopamine (100 nM) and the TAAR1 receptor inhibitor, EPPTB (500 nM), washed, and then rechallenged with octopamine (100 nM). N = 3; 7 to 10 cells/biological replicate. (E) Ca^{2+} imaging of wild type (WT) or Orai1-KO (Orai1_{fl/fl}, GFAP-Cre) primary mouse cortical astrocyte-only cultures treated with octopamine (100 nM). N = 3; 7 to 10 cells/biological replicate. (F) Ca^{2+} influx rates of WT and Orai1-KO (Orai1_{fl/fl}, GFAP-Cre) transients from E. N = 3; 7 to 10 cells/biological replicate. (G) Ca^{2+} imaging of WT or Orai1-KO (Orai1_{fl/fl}, GFAP-Cre) primary mouse hippocampal astrocyte-only cultures treated with octopamine (100 nM) in the absence of extracellular Ca^{2+} and in $2mM \text{ Ca}^{2+}$. N = 3; 7 to 10 cells/biological replicate. Credit: *Proceedings of the* National Academy of Sciences (2023). DOI: 10.1073/pnas.2217396120

Northwestern Medicine scientists have discovered how octopamine, the major "fight-or-flight" neurotransmitter in invertebrates, communicates with other cells in mammalian brains to prevent cell death, according to a study published in the *Proceedings of the National Academy of Sciences*.

Although octopamine is still found in the mammalian brain in trace amounts, its function has been replaced by epinephrine. Long thought to have been an evolutionary leftover in mammals, the role of octopamine in the <u>human brain</u> has not previously been well understood.

In the current study, investigators first set out to understand how <u>astrocytes</u>, which make up the majority of cells in the human central nervous system, contribute to brain dysfunction in <u>neurodegenerative</u> <u>diseases</u>. In astrocyte cultures from the cerebral cortex of mice, scientists



found that introducing octopamine at certain levels prompted the production of <u>lactate</u> in the astrocytes, promoting cell survival.

"Our findings are significant because we found a way in which this trace amine, octopamine, operates in the mammalian brain," said Gabriela Caraveo Piso, Ph.D., assistant professor in the Ken and Ruth Davee Department of Neurology Division of Movement Disorders.

"Think of it like an SOS signal; Neurons that are stressed send out this signal to astrocytes to send them energy, to send lactate. At the right level, octopamine allows astrocytes to read this distress signal and start making energy which will protect the cells from death by lack of ATP. If there's too much octopamine, it's sort of like smoke getting in the way of the SOS. It can't be read by the astrocytes."

The findings could help inform future therapies for Alzheimer's disease, Parkinson's and <u>bipolar disorder</u>, all of which have been associated with dysregulated levels of octopamine in the brain, Caraveo Piso said.

"Lactate was thought of as a waste product for a long time. But it turns out that it is not, it is a very important fuel that the neurons need to convert to higher forms of energy," Caraveo Piso said. "We think this is important because this can impact other diseases where octopamine levels are altered, including Alzheimer's disease and <u>psychiatric</u> <u>disorders</u>."

Moving forward, Piso and her collaborators hope to better understand how octopamine operates in healthy brains.

"What we want to know now is: Does this only happen under disease-like conditions? Or does octopamine play a role under physiological conditions such as learning and memory, where neurons also experience high energic demands?" Caraveo Piso said. "Given that octopamine can



harness lactate metabolism in astrocytes, we are also interested in understanding the role of lactate metabolism in the brain in this context of memory and learning and aging."

More information: Andrew Shum et al, Octopamine metabolically reprograms astrocytes to confer neuroprotection against α-synuclein, *Proceedings of the National Academy of Sciences* (2023). DOI: 10.1073/pnas.2217396120

Provided by Northwestern University

Citation: Understanding the role of octopamine in neurodegeneration (2023, April 28) retrieved 3 May 2024 from https://medicalxpress.com/news/2023-04-role-octopamine-neurodegeneration.html

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