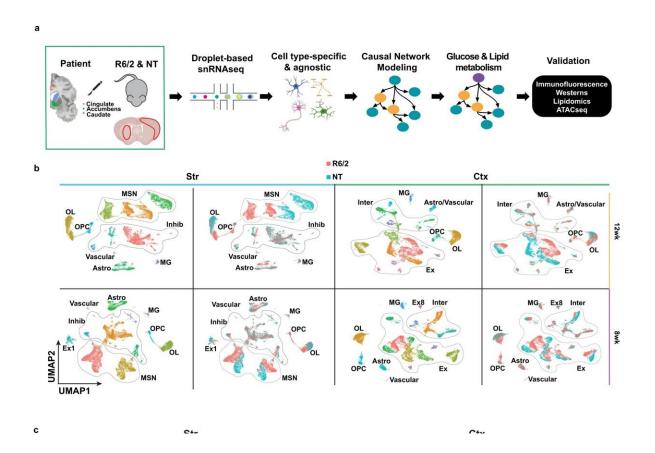


## Study links metabolism changes in certain brain cells to Huntington's disease

## December 21 2022, by Pat Harriman



Single nucleus RNAseq of mouse and human R6/2 and HD samples. **a** Illustration of workflow used for this study. After frozen tissue is microdissected from the Cingulate, Caudate, and nucleus Accumbens from 66 samples from 29 human donors (3 grade I, 4 grade II, 4 grade III, 3 grade IV, 5 juvenile-onset HD, and 10 matched controls), or the striatum and cortex of the mice (n = 3), nuclei are isolated,  $10 \times$  Libraries are prepared followed by next generation sequencing. Created with BioRender.com **b** Uniform manifold projection and approximation plots (UMAP) of the R6/2 and NT mouse data colored by cluster



or genotype. Initial QC and filtering led to the identification of 108,974 nuclei from mouse tissues. General cell type annotations: Astro Astrocytes, OL Oligodendrocyte, OPC Oligodendrocyte progenitors, MSN Medium spiny neurons, Inhib inhibitory neurons, MG Microglia, Ex Excitatory neurons, Inter Interneurons. c Barplot showing the number of up (orange) and down (blue) regulated DEGs per a cell type in the mouse data. b, c Striatal (Str, light blue bar) samples on the left and cortical (Ctx, light green bar) samples on the right, 12w samples marked by yellow bar and 8w marked by purple bar. d Proportion of R6/2 and NT cells within each cluster, red = R6/2 & blue = NT. e UMAP plots of the human snRNAseq results showing color-coded by cell type (Left), condition (Right). Credit: *Nature Communications* (2022). DOI: 10.1038/s41467-022-35388-x

A research team led by the University of California, Irvine has linked the mutation that causes Huntington's disease to developmental deficits in the brain's oligodendrocyte cells that are caused by changes in metabolism. They found that high doses of thiamine and biotin can restore normal processes.

OL cells generate the insulating coating around neurons, called myelin. The study, published online in the journal *Nature Communications*, provides detailed insight into the entire process of how these changes in the genes that regulate cell metabolism impair development of OLs, as well as the <u>therapeutic value</u> of treating HD with high doses of thiamine and biotin. Thiamine and biotin are both B vitamins and are involved in a wide range of metabolic processes that help keep the nervous system healthy.

"Our findings validate that the mutation that causes HD leads to maturation deficits in the myelin-producing cells and show that highdose thiamine and biotin treatment restores normal function of those cells," said Leslie Thompson, Ph.D., co-corresponding author and



Donald Bren and Chancellor's professor in the departments of psychiatry & <a href="https://human.behavior">human.behavior</a> and <a href="https://biological.chemistry">biological.chemistry</a> at the UCI School of Medicine, and neurobiology & behavior in the School of Biological Sciences.

Using advanced modeling methods, researchers confirmed that in mouse and human HD brain tissue, the maturation state of OL cells and their precursors are arrested in intermediate development, impairing production of the myelin that is critical for neuronal health and function. They found that high doses of thiamine and biotin were connected to significant rescue of gene expression changes in OL cells.

"The mechanisms of HD OL pathology and how these changes occur haven't been fully understood," said Ryan Lim, Ph.D., study co-first author and MIND Research Unit project scientist. "Our next steps will be to longitudinally track the effects of thiamine and biotin treatment on HD mice, so that we can further clarify those molecular and cellular processes, assess the efficacy of this therapeutic approach and identify other targets that may benefit HD patients."

**More information:** Ryan G. Lim et al, Huntington disease oligodendrocyte maturation deficits revealed by single-nucleus RNAseq are rescued by thiamine-biotin supplementation, *Nature Communications* (2022). DOI: 10.1038/s41467-022-35388-x

## Provided by University of California, Irvine

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