

Researchers pinpoint gene responsible for neurodevelopmental disorders, including autism

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Karun Singh, study co-author and researcher with McMaster's Stem Cell and Cancer Research Institute. Credit: McMaster University

A study led by researchers at McMaster University has pinpointed a gene



that is responsible for neurodevelopmental disorders, including autism.

Researchers found alterations of the gene thousand and one amino-acid kinase 2, known as TAOK2, plays a direct role in these disorders. This is the first comprehensive study that supports previous research suggesting the involvement of this gene.

The study was published in *Molecular Psychiatry*.

"Our studies reveal that in complex brain disorders that have a loss of many genes, a single deleted gene is sufficient to cause symptoms for the patients," said Karun Singh, study co-author and researcher with McMaster's Stem Cell and Cancer Research Institute.

"This is exciting because it focuses our research effort on the individual gene, saving us time and money as it will speed up the development of targeted therapeutics to this gene alone."

Many <u>neurodevelopmental disorders</u> are caused by large missing pieces of genetic material in a person's genome that contain several <u>genes</u>, termed a 'microdeletion'. Accurately diagnosing a gene microdeletion helps doctors to predict patient outcome and to determine if a new treatment is available.

The researchers used genetically engineered models and computer algorithms to study a human genome, which allowed them to pinpoint the single gene in question.

"Our next step is to screen candidate drugs that correct the cognitive brain deficits caused by genetic mutations in TAOK2, and identify candidates for pilot clinical trials," said Singh, who also holds the David Braley Chair in Human Stem Cell Research and is an assistant professor in biochemistry and biomedical sciences at McMaster.



The paper <u>complements a study</u> led by Singh on gene microdeletion published in *American Journal of Human Genetics* in early February.

"The investment into the Braley Chair for Dr. Singh and his development of key collaborations is building in multiple directions beyond what we initially imagined," said Mick Bhatia, director of McMaster's Stem Cell and Cancer Research Institute. "The combination of patient specific genetics and stem cell technologies is likely to be transformative in the near term for brain disorders."

More information: Melanie Richter et al. Altered TAOK2 activity causes autism-related neurodevelopmental and cognitive abnormalities through RhoA signaling, *Molecular Psychiatry* (2018). DOI: 10.1038/s41380-018-0025-5

Provided by McMaster University

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