

Researchers identify new drug target for schizophrenia

August 13 2012

(Medical Xpress) -- Researchers at Mount Sinai School of Medicine may have discovered why certain drugs to treat schizophrenia are ineffective in some patients. Published online in *Nature Neuroscience*, the research will pave the way for a new class of drugs to help treat this devastating mental illness, which impacts one percent of the world's population, 30 percent of whom do not respond to currently available treatments.

A team of researchers at Mount Sinai School of Medicine set out to discover what epigenetic factors, or external factors that influence [gene expression](#), are involved in this treatment-resistance to atypical antipsychotic drugs, the standard of care for schizophrenia. They discovered that, over time, an enzyme in the brains of schizophrenic patients analyzed at autopsy begins to compensate for the prolonged [chemical changes](#) caused by antipsychotics, resulting in reduced efficacy of the drugs.

"These results are groundbreaking because they show that [drug resistance](#) may be caused by the very medications prescribed to treat schizophrenia, when administered chronically," said Javier Gonzalez-Maeso, PhD, Assistant Professor of Psychiatry and Neurology at Mount Sinai School of Medicine and lead investigator on the study.

They found that an enzyme called HDAC2 was highly expressed in the brain of mice chronically treated with antipsychotic drugs, resulting in lower expression of the receptor called mGlu2, and a recurrence of [psychotic symptoms](#). A similar finding was observed in the postmortem

brains of [schizophrenic patients](#). The research team administered a chemical called suberoylanilide hydroxamic acid (SAHA), which inhibits the entire family of HDACs. They found that this treatment prevented the detrimental effect of the antipsychotic called clozapine on mGlu2 expression, and also improved the therapeutic effects of [atypical antipsychotics](#) in mouse models.

Previous research conducted by the team showed that chronic treatment with the antipsychotic clozapine causes repression of mGlu2 expression in the frontal cortex of mice, a brain area key to cognition and perception. The researchers hypothesized that this effect of clozapine on mGlu2 may play a crucial role in restraining the therapeutic effects of antipsychotic drugs.

"We had previously found that chronic antipsychotic drug administration causes biochemical changes in the brain that may limit the therapeutic effects of these drugs," said Dr. Gonzalez-Maeso. "We wanted to identify the molecular mechanism responsible for this biochemical change, and explore it as a new target for new drugs that enhance the therapeutic efficacy of antipsychotic drugs."

Mitsumasa Kurita, PhD, a postdoctoral fellow at Mount Sinai and the lead author of the study, said, "We found that [atypical antipsychotic drugs](#) trigger an increase of HDAC2 in frontal cortex of individuals with schizophrenia, which then reduces the presence of mGlu2, and thereby limits the efficacy of these drugs," said

Dr. Gonzalez-Maeso's team is now developing compounds that specifically inhibit HDAC2 as adjunctive treatments to antipsychotics. The study was funded by the National Institutes of Health.

Provided by The Mount Sinai Hospital

Citation: Researchers identify new drug target for schizophrenia (2012, August 13) retrieved 20 March 2024 from <https://medicalxpress.com/news/2012-08-drug-schizophrenia.html>

This document is subject to copyright. Apart from any fair dealing for the purpose of private study or research, no part may be reproduced without the written permission. The content is provided for information purposes only.