Learning disabilities commonly found in Wiedemann-Steiner Syndrome can now be investigated in the lab
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Genetic analysis of Wiedemann-Steiner Syndrome patients revealed that a variety of changes in one gene (MLL1) causes the disease. Researchers of the Max Planck Institute of Psychiatry in Munich and the Icahn School of Medicine at Mount Sinai in New York were able to copy the memory problems, one of the common symptoms of the syndrome, in a mouse model. When the scientists deleted part of the MLL1 gene in brain cells it led to an astonishing behavioral impairment.

MLL1 is able to change the regulation of other genes via so-called epigenetic mechanisms. The term “epigenetic” describes all processes that do not change the actual genetic code in a cell but alter its accessibility. Epigenetically regulated genes can be tagged with a chemical mark which acts like a switch. "Thus in theory, these changes could be reversible" states Mira Jakovcevski, leading scientist in the recent study. The researchers investigated such epigenetic mechanisms in their mouse model and focused on nerve cells in the brain, since these cells are required for learning. Screening of all genes present in nerve cells revealed altered epigenetic marks on many genes in the brain of mice with impaired MLL1.

For some of these genes, such as MEIS2, the epigenetic changes led to changes of related gene products as well. According to a recent genetic study, MEIS2 has been associated with learning difficulties in humans. "Now, it is important to test whether we can improve the memory performance of these mice by restoring the amount of the respective gene products" says Mira Jakovcevski. "This would be the very first step towards the development of specific therapies for the Wiedemann-Steiner Syndrome."

More information: "Neuronal Kmt2a/Mll1 Histone