Epigenetics contribute to male and female differences in fear memory

A study in Biological Psychiatry investigates sex-specific epigenetic regulation of fear memory

Philadelphia, January 17, 2019 — In a mouse model of traumatic memory, male mice recall fear-related memories better than female mice, according to a study in Biological Psychiatry. The difference between sexes was attributed to a gene important for creating fear memories and stress behavior, called cyclin dependent kinase 5 (Cdk5), which was naturally activated in male but not in female mice. The findings could help explain why fear and stress-related disorders affect men and women differently.

Fear and memory produce changes to genes that modulate gene expression, called epigenetic modifications. Epigenetic activation of Cdk5 increased naturally in males, but not in females, after the mice recalled a fear-related memory. Artificial activation of Cdk5 had no effect in male mice, in which Cdk5 was already naturally increased, but reduced the strength of fear memories in female mice, indicating sex differences in how fear is remembered.

“There is growing evidence for sex differences in the neurobiology of fear. These differences may provide important new insights into novel sex-specific treatments for anxiety disorders,” said John Krystal, MD, Editor of Biological Psychiatry.

Although previous research had already shown that Cdk5 is activated by stress and regulates the strength of fear-related memories, it had only been studied in male mice. “We examined both sexes, and found male-specific epigenetic activation of Cdk5 expression after fear conditioning, a model of traumatic memory,” said senior author Elizabeth A. Heller, PhD, University of Pennsylvania.

Dr. Heller and colleagues then used epigenetic editing to artificially increase Cdk5 activation in the hippocampus, the brain’s memory hub. “Remarkably, this manipulation reduced fear memory retrieval and increased Tau phosphorylation in female, but not male mice,” said Dr. Heller. Phosphorylation of the protein tau by Cdk5 regulates learning and memory.

“Taken together, epigenetic editing uncovered a female-specific role of Cdk5 activation in repressing fear-induced memory,” said Dr. Heller. Cdk5 activation and tau phosphorylation have been shown to cause negative effects on learning and memory in female mice, but not male mice. The authors suggest that Cdk5 expression is naturally blocked in females to protect them from these negative effects.

The epigenetic differences in male and female mice indicate sex differences in the biology of how fearful events are remembered, which highlights that sex should be an important consideration in the research and treatment of neuropsychiatric diseases that involve fear and stress, such as posttraumatic stress disorder, depression, and anxiety.
Notes for editors

Copies of this paper are available to credentialed journalists upon request; please contact Rhiannon Bugno at Biol.Psych@sobp.org or +1 214 648 0880. Journalists wishing to interview the authors may contact Elizabeth A. Heller at eheller@pennmedicine.upenn.edu.

The authors’ affiliations and disclosures of financial and conflicts of interests are available in the article.

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