Molecular mechanisms explain why people respond differently to stress, study finds

November 28 2023, by Anke Schlee

Dex SNP-DREs modulate the physiological stress response. (A) Experimental setup of a psychosocial stress task. Salivary cortisol was measured at baseline, prior to the stress task, after completion of the task, and following a 30-min rest period. (B) Association of salivary cortisol changes compared to baseline with the FGS for the pretest, posttest, and postbreak measurements. (C) Experimental setup for a fear acquisition task eliciting an eyeblink startle response. (D) Association between startle amplitude and habituation and the FGS. Note that an increased habituation z-score indicates a persisting startle response with multiple
Researchers from the Max Planck Institute of Psychiatry asked themselves which genetic variants might be involved in the response to stress. Using a new methodology, they were able to test more than 3,600 specific variants in parallel to determine which were functional.

This pioneering study of the molecular level of stress found that the genetic variants that modulate the sensitivity to the consequences of stress have an effect on the risk of psychiatric disorders. The findings are published in the journal *Proceedings of the National Academy of Sciences*.

The interaction of genes and the environment is critical in modulating the risk of developing psychiatric disorders. This has been known for many years. Stress is one of the most relevant environmental factors. But why are some people able to deal well with stress and stressful life events, while others go on to develop psychiatric disorders? Scientists from the MPI of Psychiatry (MPI) approached this question on a molecular level. They used new genetic and statistical techniques to look at many genetic variants simultaneously, in order to understand how they affect cells in their reaction to stress.

Led by postdoc and the study's first author, Signe Penner-Goeke, the team at the MPI used statistical methods which allowed them to better understand how these variants work in the brain, and even allowed them to predict causal effects. The scientists modeled stress using dexamethasone, a synthetic molecule that mimics the effect of the stress hormone cortisol.
Using cells that are known to be highly reactive to stress, they identified more than 500 regions that responded to stress as well as 79 variants that affected gene expression only when treated with dexamethasone. The findings showed that some of the variants had an effect on the risk of developing psychiatric disorders. "We found a number of variants associated with psychiatric disorders," says Penner-Goeke.

**Stress tests**

Researchers at MPI then sought to identify how the combination of variants modulated risk. In an experiment where they stressed individuals in a stress task, results showed that a higher number of variants were associated with an increase in cortisol. This difference was not seen before the stress task, meaning these variants were only relevant in stressful situations. Another test indicated that individuals with more risk variants had a more intense reaction when startled. These individuals also showed an increased amplitude of being startled with repeated exposures.

This is the first study explaining the molecular level of stress: The genetic variants that modulate the physiological response to stress are involved in the risk of psychiatric disorders. "Genetics have an effect on the sensitivity to the consequences of stress. This molecular mechanism could explain why stressful life events correlate more or less with psychiatric disorders," MPI’s director Elisabeth Binder says. These findings are essential for predicting which individuals may have a higher risk for developing psychiatric disorders in response to stress, which could be harnessed to better prevent and treat psychiatric disorders.

Provided by Max Planck Society

Citation: Molecular mechanisms explain why people respond differently to stress, study finds
(2023, November 28) retrieved 19 December 2023 from

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