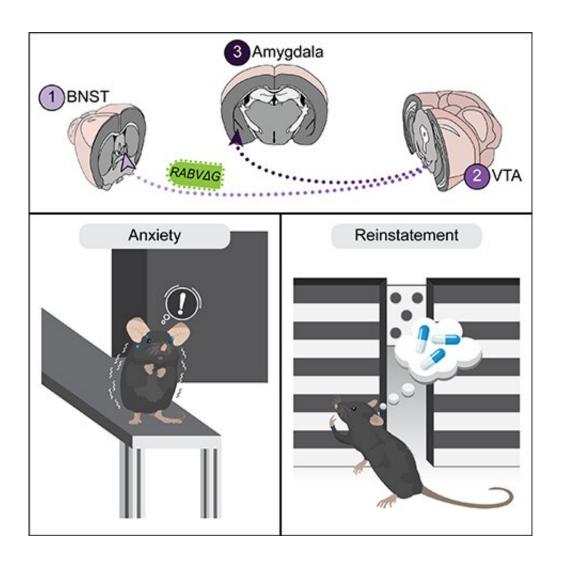


New study reveals brain circuit responsible for cocaine withdrawal-induced anxiety and relapse-related behavior

May 3 2022



(Top panel) A schematic of the brain circuit responsible for mediating drug withdrawal-induced anxiety (bottom left panel) and drug-seeking behavior (bottom right panel). Rabies (RABV\bar{2}G) tracing helped identify the circuit, comprised of the bed nucleus of the stria terminalis (BNST), ventral tegmental



area (VTA), and amygdala. Credit: UCI School of Medicine

New research from the University of California, Irvine, finds that drug withdrawal-induced anxiety and reinstatement of drug seeking behaviors are controlled by a single pathway in the brain and centered around dopamine cells.

The study, "An extended amygdala-midbrain circuit controlling <u>cocaine</u> <u>withdrawal</u>-induced anxiety and reinstatement," was published today in *Cell Reports*.

Addiction occurs in phases: Initial <u>drug</u> exposures are rewarding, repeated administration leads to tolerance or sensitization to the drug's effects, and withdrawal leads to anxiety and a negative affective state, which in turn contributes to reinstatement of drug taking/seeking.

"In order to prevent relapse among drug users, specifically cocaine users, we need to understand the factors in the brain that contribute to drug seeking behaviors and the vulnerability to relapse," said Kevin Beier, Ph.D., assistant professor of physiology and biophysics at UCI School of Medicine. "In this study, we identified a brain circuit that is responsible for drug withdrawal-induced anxiety as well as relapse-related behavior, along with the identification of a potential target for therapeutic interventions."

The negative affective state induced by withdrawal from use of drugs of abuse is a critical factor causing <u>drug users</u> to relapse.

"Both the drug withdrawal-induced anxiety and reinstatement of drug seeking are controlled by a single pathway centered around <u>dopamine</u> cells in the ventral midbrain," explained Beier. "That a single pathway



controls both sets of behavioral changes may help to explain many addiction-related behavioral phenomena. Importantly, it links them both directly to dopamine, which is more typically linked to reward-related behaviors."

Although midbrain dopamine circuits are central to motivated behaviors, the knowledge of how experience modifies these circuits to facilitate subsequent behavioral adaptations is limited. This study demonstrates the selective role of a ventral tegmental area dopamine projection to the amygdala for cocaine induced anxiety, but not cocaine reward or sensitization. Silencing this projection prevents development of anxiety during protracted withdrawal after cocaine use.

According to the National Center for Drug Abuse Statistics, there are roughly 70,000 drug overdoses each year in the United States. In 2017, nearly one in five drug overdose deaths was cocaine-related, with the highest rate of cocaine-related overdoses and deaths occurring among non-Hispanic black populations. Between 2012 and 2018, the rate of cocaine-related overdose deaths increased from 1.4 to 4.5 percent. The American Addiction Centers state recent drug relapse statistics show that more than 85 percent of individuals relapse and return to drug use within a year following treatment.

More information: Kevin Beier, An extended amygdala-midbrain circuit controlling cocaine withdrawal-induced anxiety and reinstatement, *Cell Reports* (2022). <u>DOI: 10.1016/j.celrep.2022.110775</u>. <u>www.cell.com/cell-reports/full ... 2211-1247(22)00539-3</u>

Provided by University of California, Irvine

Citation: New study reveals brain circuit responsible for cocaine withdrawal-induced anxiety and



relapse-related behavior (2022, May 3) retrieved 24 March 2023 from https://medicalxpress.com/news/2022-05-reveals-brain-circuit-responsible-cocaine.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.