Study: In heroin addiction, glial cells play key roles in regulating the motivation for the drug
29 January 2018, by Ellen Goldbaum

Published online last month in *Neuropsychopharmacology*, the paper describes how glial (non-neuronal) cells regulate both cellular and behavioral responses to heroin.

"Not much is known about glial cells in the context of addiction" said Dietz, a faculty member with UB's neuroscience program. "In the addiction field, most neuroscientists focus on neurons. Very rarely have they studied glial cells in psychiatric diseases. This work demonstrates an essential role of glia in addictive behaviors, and offers us the ability to provide a new set of targets for future therapies toward the treatment of addiction."

Dietz and his colleagues decided to study the potential role of glial cells in addiction when they found that RNA sequencing of tissue from heroin-addicted animals revealed changes in genes that are traditionally markers for a type of glial stem cell called oligodendrocyte precursor cells or OPCs.

**Opiates and the prefrontal cortex**

The research is likely the first to investigate how opiates affect adult OPCs in the brain's prefrontal cortex, which is involved in complex cognitive behaviors and is a main target of addictive drugs.

"We found that many of the genes regulated by heroin aligned with the profile of OPCs, so something was going on with them," he said.

OPCs, he explained, are cells that often become myelin, which is critical for efficient communication between neurons.

Dietz collaborated with his colleague Fraser Sim, PhD, associate professor in the Department of Pharmacology and Toxicology in the Jacobs School, co-author on the paper. In 2014, Sim...
identified one of the genes, SOX10, as a "master switch" for the differentiation of these stem cells towards myelination.

To determine what was happening when genes encoding OPCs were exposed to heroin, the scientists overexpressed them in addicted laboratory animals using viral gene therapy.

**Compensatory effect**

The result was surprising: when either of the two genes, SOX10 or BRG1, was overexpressed, the animals' motivation to take the drug was reduced.

"To our surprise, it reduced their drug-taking behavior," said Dietz. "It looks like the brain is trying to reconnect and possibly readapt myelin to normalize function, although that would need to be directly tested in future studies."

One way to think of what may be happening, he explained, is to imagine that the brain is responding to exposure to drugs of abuse by attempting to reconnect with the brain's other reward centers.

"As with any part of the body that sustains an insult, it seems that the addicted brain is trying to fix what went wrong," he said. "Our hypothesis is that after exposure to heroin, the brain starts to upregulate OPCs in an attempt to fix the altered connectivity that occurs in the addicted states. It is possible that when we facilitated OPCs, we may have reversed some of the disconnect between the prefrontal cortex and the brain's other reward regions."

**More information:** Jennifer A Martin et al. A Novel Role for Oligodendrocyte Precursor Cells (OPCs) and Sox10 in Mediating Cellular and Behavioral Responses to Heroin, *Neuropsychopharmacology* (2017). [DOI: 10.1038/npp.2017.303](https://doi.org/10.1038/npp.2017.303)

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