

Recent developments suggest potential new therapeutic role for caffeine

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Credit: Journal of Caffeine and Adenosine Research



An important new article reviews recently discovered properties of adenosine A2A receptor-dopamine D2 receptor (A2AR-D2R) and adenosine A1 receptor-dopamine D1 receptor (A1R-D1R) heteromers—both of which are main targets for caffeine—and discusses the therapeutic implications of these findings. Preclinical evidence is reviewed which indicates that caffeine and selective A2AR antagonists could be used to treat the motivational symptoms of depression as well as cognitive and emotional impairments in attention-deficit/hyperactivity disorder.

In addition, new research suggests that the A1R-D1R heteromer, which modulates the excitability of the spinal motoneuron, could be targeted by A1R antagonists to therapeutic effect in spinal cord injury. Caffeine and selective adenosine receptor antagonists as potential new therapeutic tools are discussed in an article published in the *Journal of Caffeine and Adenosine Research*, a peer-reviewed journal fromMary Ann Liebert, Inc., publishers. Click here to read the article free on the *Journal of Caffeine and Adenosine Research* website.

The article, titled "New Developments on the Adenosine Mechanisms of the Central Effects of Caffeine and Their Implications for Neuropsychiatric Disorders," is part of a series of reviews from a symposium on caffeine and adenosine presented at the meeting "Purines 2018 International," in Foz do Iguaçu, Brazil. Groundbreaking findings regarding caffeine and adenosine-dopamine receptor heteromers are reviewed by Sergi Ferré, MD, Ph.D., National Institute on Drug Abuse, National Institutes of Health (Baltimore, MD) and colleagues from the University of Puerto Rico (San Juan), the University of Connecticut (Storrs), and the Federal University of Santa Catarina (Florianópolis, Brazil).

More information: Sergi Ferré et al. New Developments on the Adenosine Mechanisms of the Central Effects of Caffeine and Their



Implications for Neuropsychiatric Disorders, *Journal of Caffeine and Adenosine Research* (2018). DOI: 10.1089/caff.2018.0017

Provided by Mary Ann Liebert, Inc

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