arXiv.org > q-bio > arXiv:1204.1395

Search or Article-id

(Help | Advan

All papers

**Download:** PDF only

Current browse cont a-bio.NC < prev | next >

new | recent | 1204

q-bio q-bio.GN

Change to browse b

References & Citation

NASA ADS

Bookmark(what is this?)









Quantitative Biology > Neurons and Cognition

## mGluR5 Knockout mice exhibit normal conditioned place-preference to cocaine

Melissa A. Fowler, Andrew L. Varnell, Donald C. Cooper

(Submitted on 6 Apr 2012)

Metabotropic glutamate receptor 5 (mGluR5) null mutant (-/-) mice have been reported to totally lack the rein- forcing or locomotor stimulating effects of cocaine. We tested mGluR5 -/- and +/+ mice for their locomotor and conditioned place- preference response to cocaine. Unlike the previous finding, here we show that compared to mGluR5 +/+ mice, -/- mice exhibit no difference in the locomotor response to low to moderate doses of cocaine (10 or 20 mg/kg). A high dose of cocaine (40 mg/kg) resulted in a blunted rather than absent locomo- tor response. We tested mGluR5 -/- and +/+ mice for conditioned place-preference to cocaine and found no group differences at a conditioning dose of 10 mg/kg, suggesting normal conditioned rewarding properties of cocaine. These results differ substantially from Chiamulera et al. (2001) and replicates Olsen et al., (2010), who found normal cocaine place-preference in mGluR5 -/- mice at 5 mg/kg. Our results indicate mGluR5 receptors exert a modulatory rather than necessary role in cocaine-induced locomotor stimulation and exert no effect on the conditioned rewarding effects of cocaine.

Comments: 2 pages, 2 figures Nature Precedings this http URL

Subjects: Neurons and Cognition (q-bio.NC); Genomics (q-bio.GN)

10.1038/npre.2011.6180.2 DOI: Cite as: arXiv:1204.1395 [q-bio.NC]

(or arXiv:1204.1395v1 [q-bio.NC] for this version)

## **Submission history**

From: Donald Cooper Ph.D. [view email] [v1] Fri, 6 Apr 2012 02:04:55 GMT (332kb)

Which authors of this paper are endorsers?

Link back to: arXiv, form interface, contact.