

# Search Results

## Table of Contents

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Search History .....	page 2
1. Internet addiction and other behavioral addictions. ....	page 3
2. Opioid use disorders. ....	page 3
3. Expression profile of nicotinic acetylcholine receptor subunits in the brain of hiv-1 transgenic rats given chronic nicotine treatment. ....	page 3
4. Mechanisms of Habitual Approach: Failure to Suppress Irrelevant Responses Evoked by Previously Reward-Associated Stimuli. ....	page 4
5. Effects of dorsal hippocampal orexin-2 receptor antagonism on the acquisition, expression, and extinction of morphine-induced place preference in rats. ....	page 4
6. Sex differences in reinstatement of alcohol seeking in response to cues and yohimbine in rats with and without a history of adolescent corticosterone exposure. ....	page 5

## Search History

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1. PsycInfo; exp ADDICTION/ OR DRUG ABUSE [+NT]/ OR DRUG USAGE; 39753 results.
2. PsycInfo; addict\*.ti,ab; 37548 results.
3. PsycInfo; 1 OR 2; 67864 results.

**1. Internet addiction and other behavioral addictions.**

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**Citation:** Child and Adolescent Psychiatric Clinics of North America, Apr 2016, (Apr 11, 2016), 1056-4993 (Apr 11, 2016)

**Author(s):** Jorgenson, Alicia Grattan; Hsiao, Ray Chih-Jui; Yen, Cheng-Fang

**Abstract:** The Internet is increasingly influential in the lives of adolescents. Although there are many positives, there are also risks related to excessive use and addiction. It is important to recognize clinical signs and symptoms of Internet addiction (compulsive use, withdrawal, tolerance, and adverse consequences), treat comorbid conditions (other substance use disorders, attention deficit hyperactivity disorder, anxiety, depression, and hostility), and initiate psychosocial interventions. More research on this topic will help to provide consensus on diagnostic criteria and further clarify optimal management. (PsycINFO Database Record (c) 2016 APA, all rights reserved)(journal abstract)

**Subject Headings:** [No terms assigned](#)

**Source:** PsycInfo

**Full Text:** Available from *Elsevier* in [Child and Adolescent Psychiatric Clinics of North America](#)

**2. Opioid use disorders.**

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**Citation:** Child and Adolescent Psychiatric Clinics of North America, Apr 2016, (Apr 9, 2016), 1056-4993 (Apr 9, 2016)

**Author(s):** Sharma, Bikash; Bruner, Ann; Barnett, Gabrielle; Fishman, Marc

**Abstract:** Opioid use and addiction in adolescents and young adults is a health problem of epidemic proportions, with devastating consequences for youth and their families. Opioid overdose is a life-threatening emergency that should be treated with naloxone, and respiratory support if necessary. Overdose should always be an opportunity to initiate addiction treatment. Detoxification is often a necessary, but never sufficient, component of treatment for OUDs. Treatment for OUDs is effective but treatment capacity is alarmingly limited and under-developed. Emerging consensus supports the incorporation of relapse prevention medications such as buprenorphine and extended release naltrexone into comprehensive psychosocial treatment including counseling and family involvement. (PsycINFO Database Record (c) 2016 APA, all rights reserved)(journal abstract)

**Subject Headings:** [No terms assigned](#)

**Source:** PsycInfo

**Full Text:** Available from *Elsevier* in [Child and Adolescent Psychiatric Clinics of North America](#)

**3. Expression profile of nicotinic acetylcholine receptor subunits in the brain of hiv-1 transgenic rats given chronic nicotine treatment.**

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**Citation:** Journal of Neurovirology, Apr 2016, (Apr 7, 2016), 1355-0284 (Apr 7, 2016)

**Author(s):** Cao, Junran; Nesil, Tanseli; Wang, Shaolin; Chang, Sulie L.; Li, Ming D.

**Abstract:** Abuse of addictive substances, including cigarettes, is much greater in HIV-1-infected individuals than in the general population and challenges the efficiency of highly active anti-retroviral therapy (HAART). The HIV-1 transgenic (HIV-1Tg) rat, an animal model used to study drug addiction in HIV-1-infected patients on HAART, displays abnormal neurobehavioral responses to addictive substances. Given that the cholinergic system plays an essential part in the central reward circuitry, we evaluated the expression profile of nine nicotinic acetylcholine receptor (nAChR) subunit genes in the central nervous system (CNS) of HIV-1Tg rats. We found that nAChR subunits were differentially expressed in various brain regions in HIV-1Tg rats compared to F344 control rats, with more subunits altered in the ventral tegmental area (VTA) and nucleus accumbens (NAc) of the HIV-1Tg rats than in other brain regions. We also found that chronic nicotine treatment (0.4 mg/kg/day) decreased the mRNA expression of nAChR subunits  $\alpha 6$ ,  $\beta 3$ , and  $\beta 4$  in the VTA of HIV-1Tg rats, whereas expression of  $\alpha 4$  and  $\alpha 6$  subunits in the NAc increased. No such changes were observed in F344 rats. Together, our data suggest that

HIV-1 proteins alter the expression of nAChRs, which may contribute to the vulnerability to cigarette smoking addiction in HIV-1 patients. (PsycINFO Database Record (c) 2016 APA, all rights reserved)(journal abstract)

**Subject Headings:** No terms assigned

**Source:** PsycInfo

#### 4. Mechanisms of Habitual Approach: Failure to Suppress Irrelevant Responses Evoked by Previously Reward-Associated Stimuli.

**Citation:** Journal of Experimental Psychology: General, Apr 2016, (Apr 7, 2016), 0096-3445 (Apr 7, 2016)

**Author(s):** Anderson, Brian A.; Folk, Charles L.; Garrison, Rebecca; Rogers, Leeland

**Abstract:** Reward learning has a powerful influence on the attention system, causing previously reward-associated stimuli to automatically capture attention. Difficulty ignoring stimuli associated with drug reward has been linked to addiction relapse, and the attention system of drug-dependent patients seems especially influenced by reward history. This and other evidence suggests that value-driven attention has consequences for behavior and decision-making, facilitating a bias to approach and consume the previously reward-associated stimulus even when doing so runs counter to current goals and priorities. Yet, a mechanism linking value-driven attention to behavioral responding and a general approach bias is lacking. Here we show that previously reward-associated stimuli escape inhibitory processing in a go/no-go task. Control experiments confirmed that this value-dependent failure of goal-directed inhibition could not be explained by search history or residual motivation, but depended specifically on the learned association between particular stimuli and reward outcome. When a previously high-value stimulus is encountered, the response codes generated by that stimulus are automatically afforded high priority, bypassing goal-directed cognitive processes involved in suppressing task-irrelevant behavior. (PsycINFO Database Record (c) 2016 APA, all rights reserved)(journal abstract)

**Subject Headings:** No terms assigned

**Source:** PsycInfo

**Full Text:** Available from *ProQuest* in *Journal of Experimental Psychology: General*

#### 5. Effects of dorsal hippocampal orexin-2 receptor antagonism on the acquisition, expression, and extinction of morphine-induced place preference in rats.

**Citation:** Psychopharmacology, Apr 2016, (Apr 6, 2016), 0033-3158 (Apr 6, 2016)

**Author(s):** Sadeghi, Bahman; Ezzatpanah, Somayeh; Haghparast, Abbas

**Abstract:** Rationale: Orexinergic system is involved in reward processing and drug addiction. Objectives: Here, we investigated the effect of intrahippocampal CA1 administration of orexin-2 receptor (OX2r) antagonist on the acquisition, expression, and extinction of morphine-induced place preference in rats. Methods: Conditioned place preference (CPP) was induced by subcutaneous injection of morphine (5 mg/kg) during a 3-day conditioning phase. Three experimental plots were designed; TCS OX2 29 as a selective antagonist of orexin-2 receptors (OX2rs) was dissolved in DMSO, prepared in solutions with different concentrations (1, 3, 10, and 30 nM), and was bilaterally microinjected into the CA1 and some neighboring regions (0.5 µl/side). Conditioning scores and locomotor activities were recorded during the test. Results: Results demonstrate that intra-CA1 administration of the OX2r antagonist attenuates the induction of morphine CPP during the acquisition and expression phases. Effect of TCS OX2 29 on reduction of morphine CPP was dose-dependent and was more pronounced during the acquisition than the expression. Furthermore, higher concentrations of TCS OX2 29 facilitated the extinction of morphine-induced CPP and reduced extinction latency period. Nevertheless, administration of TCS OX2 29 solutions did not have any influence on locomotor activity of all phases. Conclusions: Our findings suggest that OX2rs in the CA1 region of hippocampus are involved in the development of the acquisition and expression of morphine CPP. Moreover, blockade of OX2rs could facilitate extinction and may

abrogate or extinguish the ability of drug-related cues, implying that the antagonist might be considered as a propitious therapeutic agent in suppressing drug-seeking behavior. (PsycINFO Database Record (c) 2016 APA, all rights reserved)(journal abstract)

**Subject Headings:** [No terms assigned](#)

**Source:** PsycInfo

**6. Sex differences in reinstatement of alcohol seeking in response to cues and yohimbine in rats with and without a history of adolescent corticosterone exposure.**

**Citation:** Psychopharmacology, Apr 2016, (Apr 6, 2016), 0033-3158 (Apr 6, 2016)

**Author(s):** Bertholomey, M. L.; Nagarajan, V.; Torregrossa, Mary M.

**Abstract:** Rationale: Women represent a vulnerable and growing population with respect to alcohol abuse. Elevated glucocorticoid exposure in adolescence increases addiction risk and stress sensitivity in adulthood. However, little is known about sex differences in ethanol craving-like behavior. Objective: This study characterized sex differences in ethanol-motivated behavior following ethanol-paired cues and/or acute stimulation of the HPA axis in male and female rats with or without exposure to chronically elevated glucocorticoids in adolescence. Methods: Adolescent corticosterone-treated (Experiment 1) or naïve (Experiment 2) male and female rats were trained as adults to self-administer ethanol paired with a cue, and tested for the effects of this cue, alone or in combination with yohimbine, on the reinstatement of ethanol seeking. Results: Females showed elevated ethanol self-administration and seeking compared to males. In Experiment 1, corticosterone exposure in adolescence augmented cue-induced reinstatement of ethanol seeking in females only, and females were more sensitive to yohimbine in promoting reinstatement. Experiment 2 replicated these findings and showed that exposure to both yohimbine and alcohol-related cues enhanced the reinstatement of alcohol seeking, producing additive effects in females. Corticosterone levels were higher in females and in yohimbine-treated rats, and corticosterone and estradiol correlated with responding during reinstatement. Conclusions: Chronic manipulations in adolescence and acute manipulations in adulthood of the HPA axis increase cue-induced reinstatement of ethanol seeking to a greater degree in females than in males. Elucidating the mechanisms that underlie these effects may lead to the development of sex-specific interventions aimed at mitigating alcohol relapse risk in females. (PsycINFO Database Record (c) 2016 APA, all rights reserved)(journal abstract)

**Subject Headings:** [No terms assigned](#)

**Source:** PsycInfo