



RESEARCH ARTICLE

Adolescent THC exposure does not sensitize conditioned place preferences to subthreshold d-amphetamine in male and female rats [version 1; referees: 2 approved]

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Abstract

The acute effects of marijuana consumption on brain physiology and behaviour are well documented, but the long-term effects of its chronic use are less well known. Chronic marijuana use during adolescence is of increased interest, given that the majority of individuals first use marijuana during this developmental stage, and adolescent marijuana use is thought to increase the susceptibility to abusing other drugs when exposed later in life. It is possible that marijuana use during critical periods in adolescence could lead to increased sensitivity to other drugs of abuse later on. To test this, we chronically administered Δ^9 -tetrahydrocannabinol (THC) to male and female Long-Evans (LER) and Wistar (WR) rats directly after puberty onset. Rats matured to postnatal day 90 before being exposed to a conditioned place preference task (CPP). A subthreshold dose of d-amphetamine, found not to induce place preference in drug naïve rats, was used as the unconditioned stimulus. The effect of d-amphetamine on neural activity was inferred by quantifying *cfos* expression in the nucleus accumbens and dorsal hippocampus following CPP training. Chronic exposure to THC post-puberty had no potentiating effect on a subthreshold dose of d-amphetamine to induce CPP. No differences in *cfos* expression were observed. These results show that chronic exposure to THC during puberty did not increase sensitivity to d-amphetamine in adult LER and WR rats. This supports the concept that THC may not sensitize the response to all drugs of abuse.

Keywords

THC, adolescence, d-amphetamine, strain, sex, conditioned place preference

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Introduction

Marijuana is one of the most commonly used drugs of abuse worldwide¹, and the psychoactive properties of marijuana are a result of the actions of Δ^9 -tetrahydrocannabinol (THC)^{2,3}. Chronic marijuana use is associated with an increased risk of psychosis and depression⁴, and these relationships are even more concerning when use occurs during adolescence (for example, 5–7). In addition to the reported increased sensitivity of the adolescent period to the effects of marijuana, sex may also play a role in the consequences of both short- and long-term marijuana use with females more sensitive to depression and anxiety following marijuana exposure in adolescence⁸.

In addition to sex differences in the outcome of adolescent marijuana use, genetic background, including rat strain, can change the long-term consequences of THC exposure^{9,10}. Rat strains vary on measures related to learning and memory^{11–18}, anxiety¹⁸ and development¹⁰ as well as in response to drugs of abuse^{19–25}. Given that rat strains are used interchangeably in drug abuse research despite their innate differences, the inclusion of multiple strains of rat in any one study can help determine the strength and reproducibility of the long-term consequences of marijuana.

Marijuana use during adolescence may increase the likelihood of engaging in other physiologically and sociologically harmful drugs of abuse in adulthood. THC administration can potentiate the response to opioids²⁶ and nicotine²⁷, through the facilitation of brain reward mechanisms^{28,29}. However, the interaction between the consumption of one drug of abuse and initiating use of another is complex, and individual differences may predict sensitivity to other drugs, including amphetamine^{19,23,30–32}. The use of multiple rat strains, including Long-Evans (LER) and Wistar (WR) rats that have previously been observed to have differential sensitivity to THC, can model individual differences in response to THC.

This study sought to determine the long-term consequences of THC administration during the post-pubertal period in two previously studied strains of rats⁹. Following systemic administration of THC for 14 days after puberty onset, rats were aged to 90 days, at which point all rats were trained in a conditioned place preference (CPP) task to a subthreshold dose of d-amphetamine. It was hypothesized that if a particular strain and sex group was more sensitive to the effects of THC and if THC exposure increased the sensitivity to other drugs of abuse, sensitive rats would develop CPP to the sub-threshold dose of d-amphetamine and show increased neural activation, as inferred by protein expression of the immediate early gene, *Cfos*, in reward (nucleus accumbens) and context-specific (dorsal hippocampus) brain regions. However, if THC administration does not increase the sensitivity of rats to amphetamine, then no strain or sex group should show CPP behaviour in response to a sub-threshold dose of d-amphetamine and no differences in *Cfos* expression should be observed.

Methods

Experiment 1: Determining a subthreshold dose of d-amphetamine in drug naïve rats

Subjects. Subjects were purchased and shipped from Charles River (Semmeville, Quebec) as adults (250–300g) (LER female: N = 16; LER male: N = 24; WR female: N = 16; WR male: N = 16). All rats were housed in standard laboratory conditions (21°C and 35% relative humidity; 12D:12L) in Plexiglas tubs (46cm × 25cm × 20cm) with *ad libitum* access to food and water. All rat handling and procedures were done in accordance to the University of Lethbridge's Animal Welfare Committee and the Canadian Council on Animal Care guidelines.

D-amphetamine doses. Drug naïve adult rats were tested using three doses of d-amphetamine, 0.5mg/kg and 0.7mg/kg (0.49mg/ml d-amphetamine in saline, Sigma Aldrich). These doses were chosen as 1mg/kg of d-amphetamine has been shown to induce CPP in multiple research groups (as reviewed in 33) and was confirmed here in naïve LER male rats (Figure 1C). N = 8 for each strain, sex and drug dosage group.

CPP: Apparatus and training. *Apparatus* – A similar apparatus and procedure to that used for discriminative appetitive^{34,35} and fear conditioning^{36,37} to context tasks were implemented here. Briefly, opaque Plexiglas contexts that differed in shape (triangle versus square), colour (black versus white) and odour (amyl acetate versus eucalyptus), were connected with a grey alleyway. Both contexts and the alleyway were placed upon a clear Plexiglas table, and underneath the table, a mirror was inclined at a 45° angle which allowed for viewing by both an observer and a video camera.

Training – Pre-exposure: Rats were placed in the grey alleyway and allowed to freely explore both contexts for 10min then returned to their home cage. Dwell time in each chamber was recorded by an observer.

Training: The context to be paired with d-amphetamine injection (paired) and the context to be paired with a saline injection (unpaired) were assigned to each rat in a counterbalanced, quasi-random fashion. For training, rats were given 6 consecutive daily exposures³³, where they were given an injection of either saline or d-amphetamine then placed in one of the contexts for 30min. Injection type and context exposure alternated each day.

Preference: Rats explored the contexts connected by a grey alleyway for 10min. Dwell time in both contexts was recorded.

Experiment 2: CPP in adolescent THC exposed rats

Subjects, puberty onset and drug administration. Subjects were acquired, bred and handled as previously described^{9,10,15}. Briefly, male and female LER and WR (N = 9/strain and sex group) were obtained from Charles River (Semmeville, Quebec). Rats acclimated for 2 weeks before breeding. Pups were

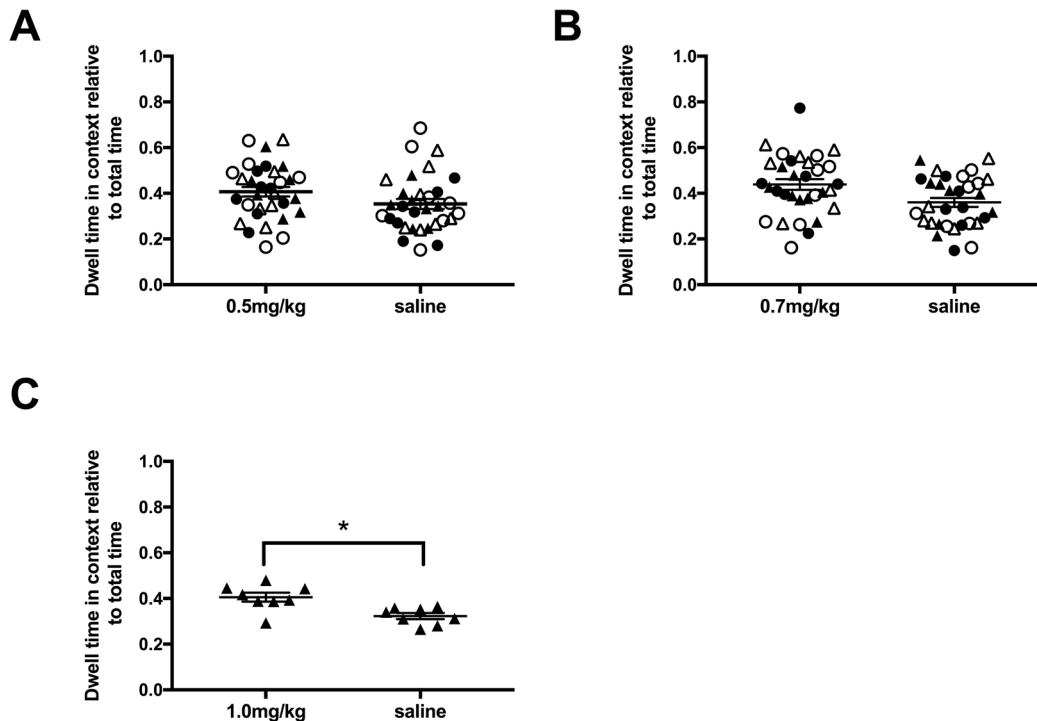


Figure 1. Dwell time in the previously paired (black) and unpaired (white) contexts during CPP. A. 0.5mg/kg d-amphetamine. **B.** 0.7mg/kg d-amphetamine. **C.** 1.0mg/kg d-amphetamine. Note 0.5 and 0.7mg/kg d-amphetamine was tested in all strain and sex groups and 1mg/kg was tested only in LER males to confirm previously published work. * $p < 0.05$. Individual data plus mean and SEE. LER females (closed circle), LER male (closed triangle), WR female (open circle), WR male (open triangle).

weaned at postnatal day 21 (p21) and placed into sex-matched pairs or triplets. $N = 8$ for all strain and sex groups for all experiments.

Puberty onset, group assignment and injection procedures were conducted as previously described¹⁰. Puberty onset was determined using the external features of the genitalia (vaginal opening and preputial separation), which correlate with gonadal hormone changes associated with puberty^{38,39}. On weaning day, rats were assigned to their experimental groups: handled control (CON), vehicle (VEH; 1:1:18 ethanol:cremaphor:saline) or 5mg/kg THC (THC). I.p. injection procedures and handling were conducted as previously described⁹. On the day of determination of puberty onset, rats were brought to a dark injection room. All rats were weighed before treatment. All rats received treatment for 14 consecutive days following determination of puberty onset. After the treatment period, rats were aged to adulthood (p90) before behavioural testing.

CPP to a subthreshold Dose of d-amphetamine: Apparatus & training. From the results of Experiment 1 (see Results section),

a subthreshold dose of d-amphetamine was determined to be 0.7mg/kg. This dose was used for all rats exposed to adolescent THC. Apparatus and training were conducted as described.

Perfusion & fixation. *Cfos* protein is present in neurons that were active 20–30min after an experience⁴⁰, and in rats, d-amphetamine will reach the brain within 5min of an i.p. injection and remain stable for 1hr⁴¹. Any *cfos* protein signal detected 1hr after d-amphetamine injection represents the neurons active 30min after d-amphetamine injection. One week after the final day of CPP, rats were injected with a single 1mg/kg dose of d-amphetamine and sat in their home cage for 1hr. Rats were euthanized with a single i.p. injection of sodium pentobarbital (120mg/kg) and transcardially perfused with approximately 150mL of 1x phosphate-buffered saline (PBS) followed by 4% paraformaldehyde (PFA) in 1xPBS. Brains were immersion fixed in 4% PFA in 1xPBS. PFA was replaced 24h after perfusion with 30% sucrose and 0.2% Na azide in 1xPBS. Brains were sectioned at 40 μ m using a cryostat (CM1900, Leica, Germany) and placed directly into Eppendorf tubes containing 0.2% Na azide in 1xPBS.

Cfos immunohistochemistry & quantification. The amount of cfos protein was stained as previously described⁴². Briefly, free-floating tissue was washed (1xPBS), followed by a 30min quenching step (0.3% H₂O₂ in 1xPBS). Tissue was blocked (1.5% goat serum in 0.3% triton-X 1xPBS) for 30min then incubated in 1° antibody (rabbit; 1:1000, 0.33% triton-X in 1xPBS with 1.5% normal goat serum; Santa Cruz, California) for 24hrs. Then, tissue was washed followed by a 24hr incubation in 2° antibody (anti-rabbit; 1:1000, Vector Labs, Canada) at room temperature. On the third day, tissue was washed then placed in AB Complex (Vector labs, Canada) for 45min. Tissue was washed then bathed for 5min in a 0.5% 3,3'-diaminobenzidine (DAB) solution (1xPBS with NiCl₂·6H₂O and 0.05% H₂O₂). Sections were washed then mounted on 1% gelatin coated slides left to dry for 24hrs, dehydrated and coverslipped with Permount.

Representative images from NAc and dorsal hippocampus were taken and quantified using particle analysis in Image J (NIH, USA). Regions of interest were defined using the Rat Brain Atlas⁴³, and particles were counted per unit area.

Vaginal cytology and determination of estrous cycle

Vaginal cytology and the determination of estrous cycle was conducted as previously described^{9,10,34}. Sterile Q-tips were dipped in sterile distilled water to collect samples onto standard glass slides (Vector labs, Canada). Vaginal smears were collected during all behavioural testing days and examined using brightfield microscopy on a Zeiss Axio Imager MT (Carl Zeiss, MicroImaging GmbH, Germany) using the 20X objective.

Statistical analysis

All raw data can be found in the raw dataset. Statistical tests were conducted using SPSS (IBM, ver 17), and estrous cycle phase was used as a covariate. For Experiment 1, a repeated measures ANOVA was conducted for percent dwell time in either context with strain and sex as the between subjects factors. Since we were interested in whether a preference for one context over another had occurred, *a priori* comparisons were conducted within each strain and sex group comparing dwell time in each context. We report partial η^2 for effect size and observed power for all results.

For Experiment 2, percent dwell time in the paired and unpaired contexts on the pre-exposure and preference days were compared within strain and sex groups using drug condition (group) as a between subjects factor. *A priori* hypotheses were established such that within each drug group and within

each strain and sex group, comparisons between the paired and unpaired contexts were always conducted. For *cfos* quantification, between subjects comparisons within strain and sex groups were conducted in order to determine the effects of drug exposure on a specific strain and sex group.

Results

Estrous cycle did not significantly alter any of the results and was not included as a covariate in subsequent analyses.

Experiment 1: Determination of a subthreshold dose of d-amphetamine

No initial preference nor any preference after training was observed for 0.5 (Figure 1A) or 0.7mg/kg (Figure 1B) of d-amphetamine for any strain or sex group (see Table 1 for statistical results). A dose of 1mg/kg d-amphetamine was used to confirm previous experiments and did induce significant place preference (see Figure 1C), thus 0.7mg/kg dose was considered subthreshold for all subsequent experiments.

Experiment 2: CPP to a sub-threshold dose of d-amphetamine

There were no pre-existing bias to spend more time in the paired or unpaired context, regardless of strain, sex or drug administration. No interaction between drug or context were observed in any strain and sex group. On the preference day, LER females overall spent significantly more time in the paired context ($F_{(1, 21)} = 17.483$, $p < 0.001$; Figure 2A). No overall effect of group was observed. Individual comparisons within groups revealed that CON ($p = 0.04$) and VEH ($p = 0.028$) LER females spent significantly more time in the context paired with d-amphetamine. No such difference was observed within LER females exposed to THC, although this value did approach statistical significance ($p = 0.065$). LER males (Figure 2B), WR females (Figure 2C) and WR males (Figure 2D) showed no significant effect of drug as well as did not show an overall preference for one context over the others (see Table 2 for statistical results).

Cfos immunohistochemistry

No significant effects were observed for any strain and sex group for *cfos* expression in dorsal hippocampus (Figure 3) and NAc (Figure 4) following a 1mg/kg injection of d-amphetamine.

Dataset 1. Raw data associated with Figure 1 – Figure 4

<http://dx.doi.org/10.5256/f1000research.14029.d196720>

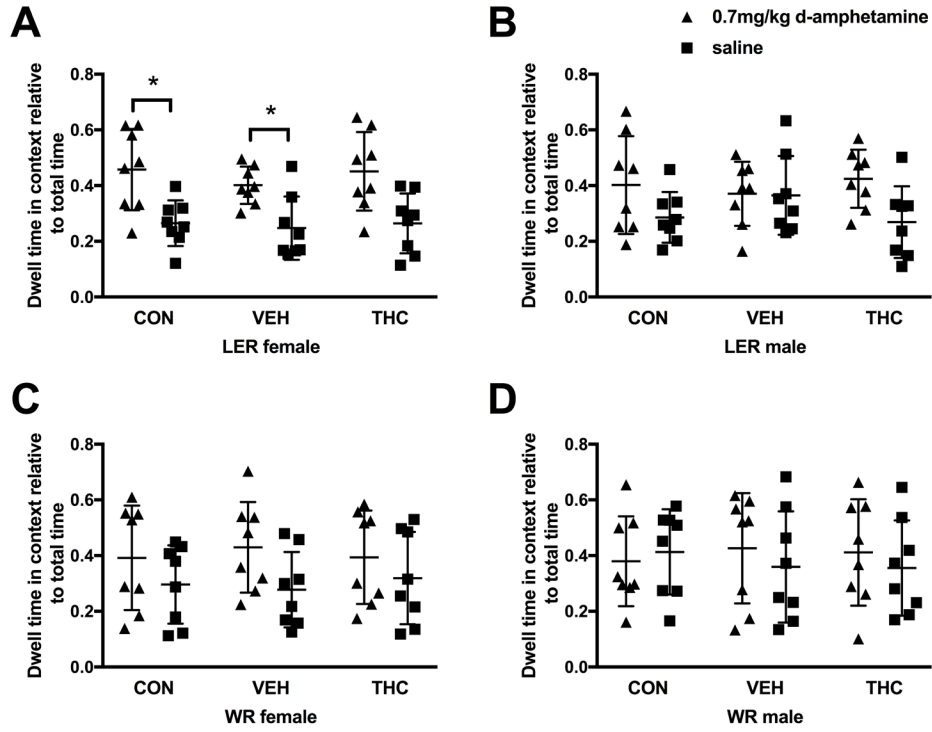


Figure 2. CPP to 0.7mg/kg d-amphetamine in THC exposed adolescent LER and WR male and female rats. Preference for **A)** LER females, **B)** LER males, **C)** WR females and **D)** WR males. * $p < 0.05$.

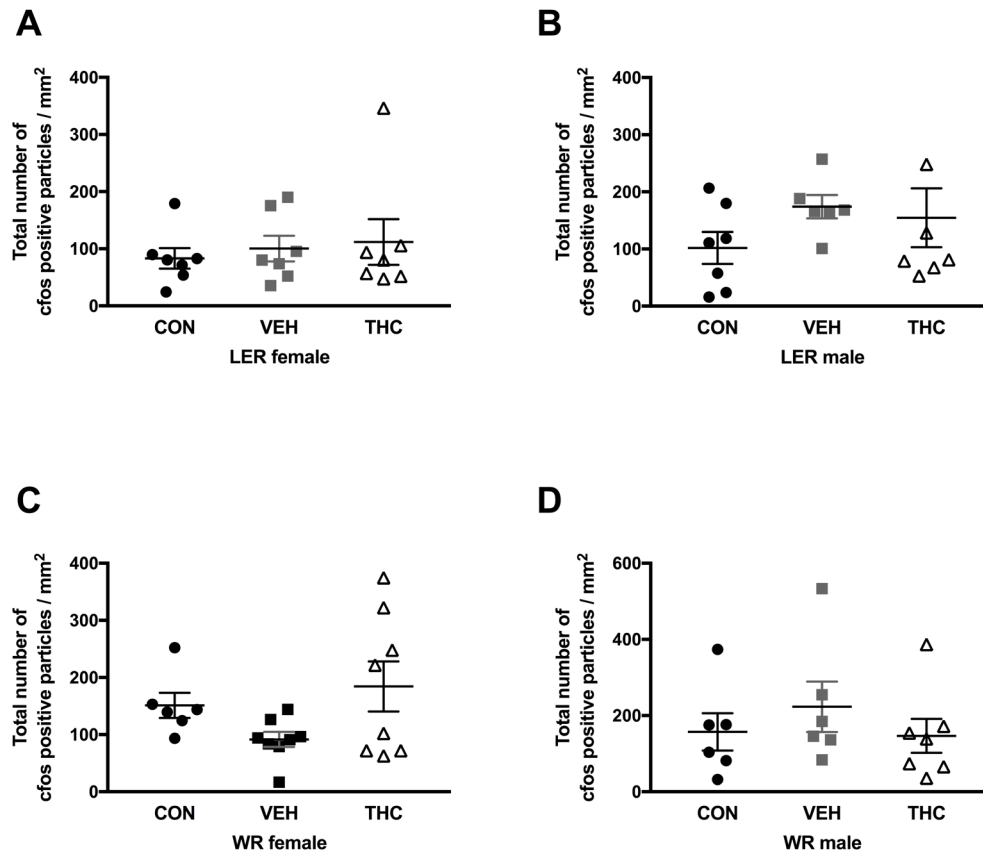


Figure 3. *Cfos* staining in dorsal hippocampus in LER and WR male and female rats exposed to THC as adolescents.

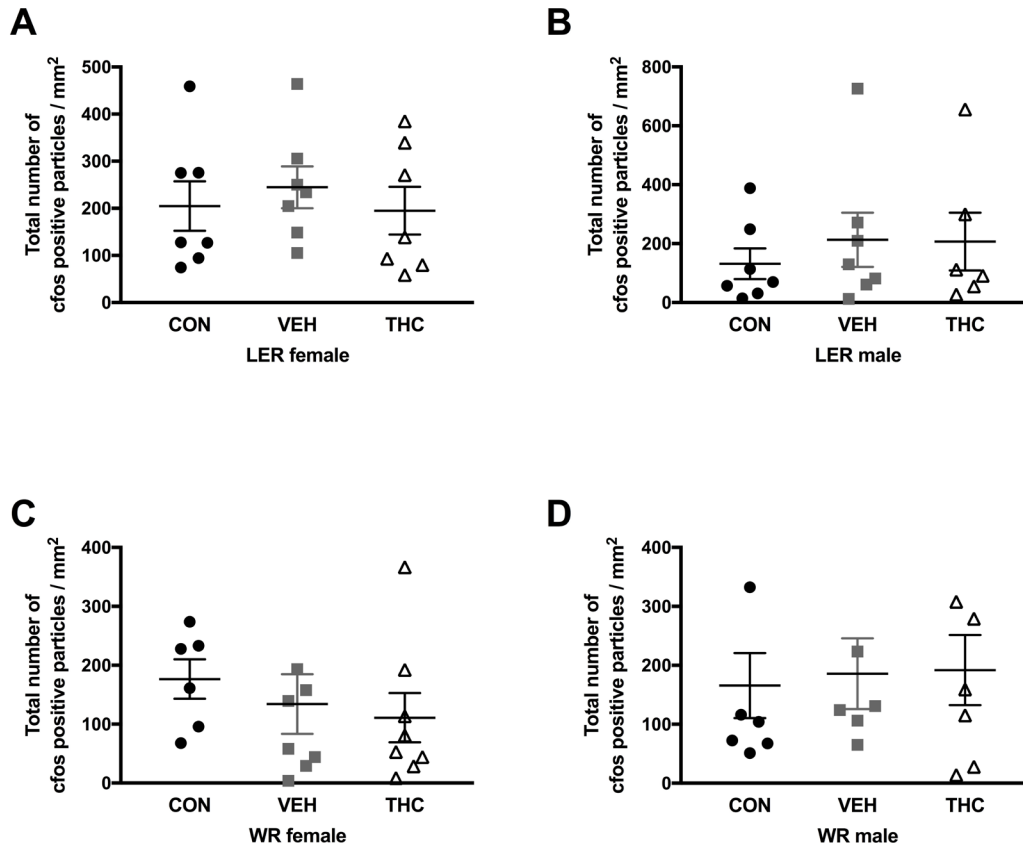


Figure 4. *Cfos* staining in NAC in LER and WR male and female rats exposed to THC as adolescents.

Discussion

Here, we report no long-term consequences of adolescent THC exposure on sensitivity to d-amphetamine in adulthood. We did observe that rearing environment affected sensitivity to d-amphetamine in LER females; CON LER females bred in house expressed CPP behaviour to a 0.7mg/kg dose of d-amphetamine, whereas those obtained from a commercial breeder (Charles River) did not. Additionally, using immediate early gene protein expression, we observed no significant effect of THC exposure following puberty onset in nucleus accumbens and dorsal hippocampus d-amphetamine-induced activation.

No effect of THC

Adolescent THC exposure did not potentiate the adult response to d-amphetamine. D-amphetamine increases dopaminergic tone when systemically administered^{44–47} and is highly rewarding^{48,49}. Given the premise that THC acts as a gateway drug, we assumed adolescent exposure to THC would potentiate reward circuitry, enhancing sensitivity to d-amphetamine.

Priming of amphetamine response by cannabinoids has been observed by some researchers^{50–52} and not others⁵³. Differences between among studies look for this effect include the dose, duration and starting age of exposure to THC as well as the timing of exposure to amphetamine, which one study reported

amphetamine-primed reward to be dependent on the time since exposure to THC⁵¹. However, our results should help mitigate many of these issues, as our dose of THC was relatively moderate, was given following the onset of puberty, which can be influenced by THC⁵⁴ and lasted throughout the adolescent period and in to early adulthood, all of which are reasonable analogues, given experimental constraints, to the human adolescent marijuana consumption experience. One possible explanation for the pattern of results obtained in the present study may be the use of CPP versus the self-administration paradigm. CPP is a standard metric for determining the rewarding properties of drugs of abuse and has been observed for multiple doses of drugs, including amphetamine^{33,55–57}. Future experiments should consider allowing animals to self-administer either THC or amphetamines, potentially looking at the correlations between self-administration of both drugs. Unfortunately, THC has proven problematic in self-administration paradigms in ^{55,58,59}.

Previous studies have demonstrated priming effects of THC to other drugs of abuse. Increased self-administration of heroin or other opiates has been observed^{52,60–62}, partially dependent on cannabinoid receptors⁶³. Thus, the endogenous opioid system is particularly sensitive to the long-term consequences of THC. Indeed, given the increased abuse of prescription opiates, research examining the interplay between the endogenous cannabinoid

and opioid systems could potentially prevent the transition of using marijuana to opiates. It is possible that priming of THC is specific to drugs targeting the opioid system.

Effect of rearing environment

LER females bred in house at the University of Lethbridge expressed CPP to a 0.7mg/kg dose of d-amphetamine whereas those purchased from Charles River did not. This was exclusively observed in LER females. Strain differences in response to amphetamine have been observed previously^{19,20,22,23}. This strain and sex specific effect in response to amphetamine may be the result of the interplay between the stress system and monoaminergic function, which has been posited to explain differences in response to amphetamine between two other strains of rats, Fisher 344 and Lewis^{64,65}. It is possible that differences in these systems may occur in LER females reared under different conditions^{66,67}. Regardless, further understanding of this fascinating effect of strain and rearing conditions should be explored as it is clear that genetic and differences in rearing correlate highly with drug abuse in adulthood^{21,32,68–70}.

Strain-dependent sex difference in response to amphetamines has been observed previously⁶⁴, although this effect was not observed in LER. Differential responses in one sex and not the other across strains are not uncommon (for example, 71), however most studies examining strain differences in response to drugs of abuse typically only use males (as discussed in 64). There is a tendency for females to be more sensitive to drugs of abuse, including amphetamine^{30,64,72–75}, which is partially mediated through the endogenous hormonal rhythms of females^{76–78}. Here, training days covered the extent of at least one full estrous cycle, and there was no significant effect of estrous cycle phase on CPP behaviour. Thus, we have identified that LER females are sensitive to rearing environment in relation to CPP behaviour in response to amphetamine. This kind of effect should not be underestimated as the implications of ignoring sex, strain, rearing

differences and their interactions in research are being increasingly recognized by granting agencies and scientific organizations to contribute to individual differences and reproducibility in current neuroscience research.

Conclusions

This study does not support a link between adolescent THC exposure and sensitivity to other drugs of abuse, where rats were tested for changes in sensitivity to d-amphetamine following long-term exposure of THC during adolescence. This is surprising, given the vulnerability of LER females to developmental perturbations (in this case, rearing environment) on d-amphetamine CPP. WR displayed stable behavioural profiles; neither rearing environment nor THC administration altered their response to a sub-threshold dose of d-amphetamine. Our previous research identified WR as resilient to the effects of adolescent THC exposure⁹. Further research into discovering the mechanisms behind resiliency in these groups may help identify mechanisms that can be protective for groups at-risk to the development of addiction.

Data availability

Dataset 1: Raw data associated with Figure 1–Figure 4. [10.5256/f1000research.14029.d196720](https://doi.org/10.5256/f1000research.14029.d196720)⁷⁹

Competing interests

The authors declare that there are no competing interests.

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Current Referee Status:  

Version 1

Referee Report 25 April 2018

doi:[10.5256/f1000research.15248.r32202](https://doi.org/10.5256/f1000research.15248.r32202)



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This report by Keeley et al., examines the effects of adolescent THC exposure on conditioned place preference for d-amphetamine. While no effects of adolescent THC exposure were found on the CPP or the neural correlates (c-fos expression), the authors' inclusion of important factors such as strain and sex make this an important study to consider for those interested in studying the effects of developmental cannabinoid exposure on future risk for substance use. Suggested changes would help to improve the impact of this paper.

Abstract:

- Last Sentence: include "sub-threshold dose of" before d-amphetamine

Introduction:

- The introduction begins with mentions of psychosis, anxiety and depression, whereas the central questions being asked in this study are not related to those topics. The authors should think about setting the research question in the opening paragraph.

Methods:

- Subjects: This section is unclear. Are the subjects being referred to here the parents of the rats used in the study?
- D-Amphetamine Doses: 1 mg/kg not listed here
- Statistical Analyses: Unclear whether dwell times were also compared within-animal pre- and post-training?

Results:

- Tables missing, but mentioned in the text.

Discussion:

- Since the authors chose to discuss psychosis in the introduction, maybe they can discuss it in the context of the null results of d-amphetamine on c-fos expression, since an accentuated response to amphetamine is something that is seen in patients with schizophrenia.

An excellent review by Chadwick and Hurd¹ might help to formulate some of these thoughts.

Rearing environment was not one of the research objectives/hypotheses and the relevance to this discussion needs to be strengthened.

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Is the work clearly and accurately presented and does it cite the current literature?

Yes

Is the study design appropriate and is the work technically sound?

Yes

Are sufficient details of methods and analysis provided to allow replication by others?

Yes

If applicable, is the statistical analysis and its interpretation appropriate?

Yes

Are all the source data underlying the results available to ensure full reproducibility?

Yes

Are the conclusions drawn adequately supported by the results?

Yes

Competing Interests: No competing interests were disclosed.

I have read this submission. I believe that I have an appropriate level of expertise to confirm that it is of an acceptable scientific standard.

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This article contributes valuable insight into issues that are commonly observed in literature regarding the impact of adolescent Δ^9 -tetrahydrocannabinol (THC) exposure on future drug-seeking behavior. Particularly, careful attention is directed towards the influence of sex (including estrous cycle) and *animal strain* on modulation of drug-seeking behavior by adolescent THC administration. These crucial factors likely contribute to conflicting findings reported on the influence of exogenous and endogenous cannabinoids on future drug consumption, and Keeley and colleagues illustrate the necessity of implementing sex and strain variables in experimental designs when addressing such research questions. As such, the authors' findings provide a more informative story that will be useful in guiding legislation on marijuana use policies. The introduction of this article provides clear explication of hypotheses and expected results as well as suitable background information that warrants investigation of the authors' research questions. However, there are some minor issues regarding the presentation of methods and

statistical results. Additionally, there are some aspects of the discussion that could benefit from citation of supporting literature.

Methods

In the statistical analysis section, the authors indicate that effect size and statistical power would be reported for all results (pg. 5), yet these data are not provided when presenting results in text. The analytical approaches utilized by the researchers are appropriate for the data; although, brief explanations of post-hoc statistical tests should be included.

It is stated in the *Subjects* section (pg. 3), that all rats were purchased and shipped from Charles River. However, in the first paragraph of the discussion, the authors allude to an effect of rearing environment on d-amphetamine sensitivity. Specifically, Long Evans females bred in-house showed CPP to the 0.7 mg/kg dose, but this was absent in Long Evans females obtained from a commercial breeder. These data are valuable and should be included, but the analyses do not appear anywhere in the manuscript.

Inclusion of an experimental timeline would increase the clarity of the researchers' methodological approach. It would also provide a useful model for future researchers interested in testing the effects of THC administration on the reinforcing effects of drugs of abuse.

Given that an observer scored dwell time in the CPP chambers, it would be beneficial to include the familiarity of the observer with animals' drug treatment group. In other words, were the observers blinded to treatment conditions?

The authors state that 3 doses of D-amphetamine were used, but only 2 doses are listed (0.5 and 0.7 mg/kg). We assume that the third dose is the vehicle (0 mg/kg), but that should be explicitly listed in the Methods (see pg. 3).

Results

The results are clearly and concisely stated, and the provided figures summarize the data well. However, Table 1 referenced in the "Experiment 1: Determination of a subthreshold dose of d-amphetamine" section and Table 2 referenced in the "Experiment 2: CPP to a sub-threshold dose of amphetamine" section are not provided in the article (see pg. 5).

In the caption for Figure 1, there is a space inserted into the word "Note". Also in this caption it is stated that mean and "SEE" are provided. Presumably this was supposed to read SEM.

The authors state that estrous cycle did not have any effect on the results (pg. 5). However, with only an N=8, it may simply be that there was not sufficient power to detect any differences. This should be briefly mentioned as a potential caveat.

Discussion

On pg. 8, the authors indicate that "this study does not support a link between adolescent THC exposure and sensitivity to other drugs of abuse...". Given that only sensitivity to d-amphetamine was assessed, it may be more reasonable to conclude from the provided results that adolescent THC exposure does not enhance sensitivity to d-amphetamine specifically.

On pg. 7 – 8, the authors suggest that "...given the increased abuse of prescription opiates, research examining the interplay between the endogenous cannabinoid and opioid systems could potentially prevent the transition of using marijuana to opiates." This is an exciting prospect for future research, and

some progress has been made in this regard. The authors may find it useful to include conclusions from Markos et al., (2017)¹ in which cannabidiol is shown to attenuate morphine CPP in mice.

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Is the work clearly and accurately presented and does it cite the current literature?

Yes

Is the study design appropriate and is the work technically sound?

Yes

Are sufficient details of methods and analysis provided to allow replication by others?

Yes

If applicable, is the statistical analysis and its interpretation appropriate?

Yes

Are all the source data underlying the results available to ensure full reproducibility?

Yes

Are the conclusions drawn adequately supported by the results?

Yes

Competing Interests: No competing interests were disclosed.

Referee Expertise: Endocannabinoids, cannabis, stress, reward, sex differences

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