RTICBM-74 is a Brain-Penetrant CB₁ Receptor Allosteric Modulator that Reduces Alcohol Intake in Rats

Dennis Lovelock,¹ Thuy Nguyen,³ Kalynn Van Voorhies,¹ Yanan Zhang,^{3} and Joyce
Besheer^{1,2*}*

¹ Bowles Center for Alcohol Studies, ² Department of Psychiatry, University of North Carolina -
Chapel Hill, Chapel Hill, NC, 27599-7178

³ Research Triangle Institute, Research Triangle Park, NC, 27719

*co-corresponding authors

Running Title: CB₁ NAM RTICBM-74 reduces alcohol intake in rats

Correspondence:

Joyce Besheer, Ph.D.

Professor

Bowles Center for Alcohol Studies

Department of Psychiatry

University of North Carolina at Chapel Hill

Chapel Hill, NC 27599-7171

Email: joyce_besheer@med.unc.edu

Yanan Zhang, Ph.D.

Sr. Research Chemist

Research Triangle Institute

Research Triangle Park, NC 27709

Email: yzhang@rti.org

Text pages: 20

Tables: 3

Figures: 4

References: 36

Abstract words: 190

Introduction words: 639

Discussion words: 953

Abbreviations:

ACN - acetonitrile

ADME - absorption, distribution, metabolism and excretion

AUC - area under the curve

AUD – alcohol use disorder

CB₁ – cannabinoid receptor subtype 1

GPCR – G-coupled protein receptor

HTD - high throughput dialysis

MDCK - Madin-Darby canine kidney

NAM – negative allosteric modulator

PAM – positive allosteric modulator

Papp - apparent permeability

PK – pharmacokinetics

TEER - Trans Epithelial Electric Resistance

Section assignment recommendation: Drug Discovery and Translational Medicine

ABSTRACT

The endocannabinoid system is implicated in the neuronal mechanisms of alcohol use disorder (AUD), with the cannabinoid receptor subtype 1 (CB₁) representing a promising target for AUD therapeutic interventions. We have previously shown negative allosteric modulators (NAMs) of the CB₁ receptor attenuated the reinstatement of other drugs of abuse including cocaine and methamphetamine in rats; however, their effects on alcohol-related behaviors have not been investigated. Here, we tested the pharmacokinetic properties of one such CB₁ NAM, RTICBM-74, and its effects on alcohol self-administration in rats. RTICBM-74 showed low aqueous solubility and high protein binding but had excellent half-life and low clearance against rat liver microsomes and hepatocytes, and excellent brain penetrance in rats. RTICBM-74 pretreatment specifically reduced alcohol intake across a range of doses in male or female Wistar or Long-Evans rats that were trained to self-administer alcohol. These effects were similar to the CB₁ antagonist/inverse agonist rimonabant which was tested as a positive control. Importantly, RTICBM-74 was effective at reducing alcohol intake at doses that did not affect locomotion or sucrose self-administration. Our findings suggest that CB₁ NAMs such as RTICBM-74 have promising therapeutic potential in treatment of AUD.

SIGNIFICANCE STATEMENT

The present work shows that a metabolically stable and brain penetrant CB₁ receptor negative allosteric modulator (NAM) reduces alcohol self-administration in rats without affecting locomotion or sucrose self-administration, suggesting potential therapeutic relevance for the treatment of AUD.

Keywords: alcohol; CB₁; allosteric modulation; self-administration; AUD

INTRODUCTION

Alcohol use disorder (AUD) is a chronic relapsing brain disease characterized by compulsive alcohol use despite adverse consequences (Koob, 2013). According to the 2015 National Survey on Drug Use and Health (NSDUH), 15.1 million adults ages 18 and older had AUD in the United States. Three medications, naltrexone (opioid antagonist), acamprosate (NMDA receptor agonist), and disulfiram (acetaldehyde dehydrogenase inhibitor), have been approved by the US Food and Drug Administration (FDA) to treat AUD; however, they have only modest efficacy (Heilig & Egli, 2006), illustrating the need for additional therapeutic agents. The cannabinoid type-1 receptor, a key member of the endocannabinoid system, plays a profound role in alcohol related behaviors and has been considered a promising target for alcohol addiction treatment.

The CB₁ receptor, well known for its role in addiction of multiple drugs (Parsons & Hurd, 2015), contributes to alcohol related behaviors including alcohol consumption, alcohol tolerance, alcohol dependence and addiction. CB₁ agonist CP55,940 dose-dependently increased ethanol consumption (Gallate et al., 1999), whereas CB₁ antagonist/inverse agonist rimonabant (SR141716) reduced ethanol intake in a number of preclinical models of ethanol consumption (Arnone et al., 1997; Colombo et al., 2007; Economidou et al., 2006; Hansen et al., 2008; Maccioni et al., 2009). CB₁ knock-out (KO) mice displayed diminished ethanol-induced conditioned place preference (Houchi et al., 2005). Acute administration of rimonabant completely abolished the alcohol deprivation effect (i.e., relapse-like drinking) and motivation to drink in alcohol-preferring rats (Colombo et al., 2004; Serra et al., 2002; Vinod et al., 2012), while chronic administration during alcohol exposure reduced severity of alcohol withdrawal-induced handling-induced convulsions in both alcohol-preferring and alcohol-avoiding mice (Vinod et al., 2008). Finally, rimonabant treatment prior to a reinstatement session reduced cue-

induced relapse (Cippitelli et al., 2005; Economidou et al., 2006; Economidou et al., 2007). Together, these results demonstrate the promise of CB₁ blockade as potential treatments of AUD. Unfortunately, after its initial approval in Europe for the treatment of obesity (Cahill & Ussher, 2007), rimonabant was eventually withdrawn from the market because of untoward side effects associated with its use in humans including depression, anxiety or even suicidal ideation (Beyer et al., 2010; Christensen et al., 2007; Moreira et al., 2009).

Several alternate approaches to modulate the CB₁ signaling that may avoid the psychiatric side-effects of CB₁ antagonists/inverse agonists have been pursued (Cinar et al., 2020; Nguyen et al., 2019), among which allosteric modulation has received particular attention. Allosteric modulators of G protein-coupled receptors (GPCRs) are molecules that bind at topographically distinct allosteric sites and either potentiate (positive allosteric modulators or PAMs) or inhibit (negative allosteric modulators or NAMs) the binding and/or signaling of an orthosteric ligand (Kenakin & Miller, 2010). Several CB₁ NAMs have been reported thus far (Nguyen et al., 2017b), including the two most extensively studied Org27569 and PSNCBAM-1. These NAMs were able to attenuate or block functional responses of CB₁ agonists such as CP55940 in several *in vitro* assays (Horswill et al., 2007; Price et al., 2005). We found that Org27569 and PSNCBAM-1, like rimonabant, resulted in a dose-related attenuation of cue- and/or drug-induced reinstatement of cocaine- and/or methamphetamine-seeking behavior (Jing et al., 2014; Nguyen et al., 2017a). Continued efforts from our team resulted in the identification of CB₁ NAM RTICBM-74, which displayed improved metabolic stability over PSNCBAM-1. Despite their similar *in vitro* potency, RTICBM-74 at 10 mg/kg produced the same degree of attenuation as that of 30 mg/kg of PSNCBAM-1 in prime-induced reinstatement of cocaine seeking behavior, suggesting improved *in vivo* potency (Nguyen et al., 2017a). However, the effects of

RTICBM-74 on alcohol reinforcement have yet to be studied.

Here, we first assessed the ADME (absorption, distribution, metabolism, and excretion) and pharmacokinetic properties of RTICBM-74 and then investigated its effects on alcohol self-administration in rats. To assess the reinforcer specificity of RTICBM-74, we also tested sucrose self-administration in parallel and tested rimonabant as a positive control.

MATERIALS AND METHODS

Drugs

RTICBM-74 was synthesized at Research Triangle Institute following previously published procedures (purity >98%) (Nguyen et al., 2017a). For behavioral experiments RTICBM-74 was suspended in a mixture of saline, tween-80 and DMSO of an 8:1:1 ratio. Rimonabant (National Institute on Drug Abuse Drug Supply Program) was dissolved in saline and 2 drops of Tween-80 per 5 ml of saline. All injections were administered at a volume of 1 mL/kg.

RTICBM-74 properties

Receptor Selectivity. The off-target selectivity of RTICBM-74 was assessed in radioligand binding assays against a panel of ~50 GPCRs, ion channels, and transporters at a single concentration of 10 μ M by NIMH Psychoactive Drug Screening Program (PDSP) according to their standard protocols. These targets included adrenergic (α_{1A} , α_{1B} , α_{1D} , α_{2A} , α_{2B} , α_{2C} , β_1 , β_2 , β_3), dopamine (D1 - D5), GABA_A, histamine (H1 - H4), muscarinic (M1 - M5), opioid (μ , κ , δ), serotonin (5-HT1A, 5-HT1B, 5-HT1D, 5-HT1E, 5-HT2A, 5-HT2B, 5-HT2C, 5-HT3, 5-HT5A, 5-HT6, 5-HT7A), sigma (σ_1 , σ_2), dopamine transporter (DAT), norepinephrine transporter (NET), serotonin transporter (SERT), and BZP Rat Brain Site. The percentage of inhibition was the average of two determinations. When significant displacement of radioligand was observed

(>50% inhibition at 10 μ M), complete concentration-dependent displacement curves (in duplicate) were constructed to generate IC_{50} values. IC_{50} values were determined by a nonlinear regression analysis. The equilibrium dissociation constant (K_i) was calculated with the Cheng–Prusoff equation using the observed IC_{50} of RTICBM-74, the concentration of radioligand, and the historical values of K_d of the ligand. The full methods and references can be found at: <http://pdspdb.unc.edu/pdspWeb>.

ADME Studies.

LC-MS/MS method used to quantify RTICBM-74 concentrations. RTICBM-74 bioanalytical samples were analyzed by LC-MS/MS using a CTC PAL autosampler, Thermo Accela UPLC and a Thermo Quantum triple quadrupole mass spectrometer. The $[M+H]^+$ adduct of RTICBM-74 and internal standards were monitored using positive mode electrospray ionization in multiple reaction monitoring (MRM) mode with precursor ion of 341.0 and product ion of 128.0. The lower and upper limits of quantification are 0.25 ng/mL and 1,000 ng/mL respectively. The analytes were injected onto a C18 column and chromatographed using a reverse phase gradient with 0.1% formic acid in water and 0.1% formic acid in isopropanol/acetonitrile (1:4). Peak integrations were performed using Thermo Xcalibur (v2.4) software.

Hepatocyte Stability. Plateable cryopreserved male Sprague Dawley rat hepatocytes (BioreclamationIVT, USA) were thawed in recovery media and diluted in plating media to a viable cell density of 1×10^6 cells/mL. Viability was determined by trypan blue exclusion. A 50 μ L suspension of cells was added to each well of a 96-well plate and pre-incubated for 10 minutes at 37°C, 5% CO_2 . At time zero, 50 μ L of test articles and reference compounds (testosterone and ethoxycoumarin) were added in duplicate to reach a final concentration of 1 μ M and to initiate the reaction. At selected time points (15, 30, 60, 120 and 240 min), the plate

was removed from the incubator and the wells were quenched with 150 μ L of (75/25) ice-cold acetonitrile (ACN) in water containing 2 % formic acid and internal standard (0.1 μ M glyburide and labetalol). The 0 min time points were spiked in the pre-quenched wells. Samples were then centrifuged and supernatants were transferred to a new plate for analysis by LC-MS/MS method described above. The half-life ($t_{1/2}$) was determined from a plot of the natural logarithm of the peak area ratio (remaining compound peak area/internal standard peak area) against time using Microsoft Excel. Intrinsic clearance was calculated by the below equation.

$$Cl_{int} \text{ (mL/min/kg)} = \frac{-\text{slope}}{0.5 \times 10^6 \text{ cells/mL}} \times \text{Species specific factor}$$

Where species specific factor = 1,800 for rats

Kinetic Solubility. The kinetic solubility of the test compound was measured in commercial PBS, which consisted of potassium phosphate monobasic 1 mM, sodium phosphate dibasic 3 mM and sodium chloride 155 mM. A 10 μ L of test compound stock solution (20 mM DMSO) was combined with 490 μ L of PBS buffer to reach a targeted concentration of 400 μ M. The solution was agitated on a VX-2500 multi-tube vortexer (VWR) for 2 hours at room temperature. Following agitation, the sample was filtrated on a glass-fiber filter (1 μ m) and the eluate was diluted 400-fold with a mixture of acetonitrile: water (1:1). On each experimental occasion, nicardipine and imipramine were assessed as reference compounds for low and high solubility, respectively. All samples were assessed in triplicate and analyzed by LC-MS/MS using electrospray ionization against standards prepared in the same matrix.

Plasma Protein Binding. Test compound and controls (1 μ M) were spiked in rat plasma (BioreclamationIVT, Westbury, NY, USA) and aliquoted in triplicate in a high throughput dialysis (HTD) 96-well plate, where the plasma and dialysate buffer were separated by a semi-

permeable cellulose membrane (12-14K MWCO). Once sealed, the HTD plate was incubated at 37°C and kept under light agitation for 6 hours, until equilibrium was reached. Plasma and buffer samples were then extracted along with their corresponding standard curve samples using ice-cold acetonitrile in methanol (1:1). After centrifugation, supernatants from both plasma- and buffer-containing samples were further diluted prior to being submitted to bioanalysis by LC-MS/MS. Acebutolol and warfarin served as low-bound and highly-bound controls, respectively. Determination of the percentage of plasma protein binding and recovery was achieved in Microsoft Excel.

Bidirectional MDCK-mdr1 permeability. MCK-mdr1 cells were seeded onto permeable polycarbonate supports in 12-well Costar Transwell plates and allowed to grow and differentiate for 3 days. On day 3, culture medium (DMEM supplemented with 10% FBS) was removed from both sides of the transwell inserts and cells were rinsed with warm HBSS. After the rinse step, the chambers were filled with warm transport buffer (HBSS containing 10 mM HEPES, 0.25% BSA, pH 7.4) and the plates were incubated at 37°C for 30 min prior to TEER (Trans Epithelial Electric Resistance) measurements. The buffer in the donor chamber (apical side for A-to-B assay, basolateral side for B-to-A assay) was removed and replaced with the working solution (10 μ M test article in transport buffer). The plates were then placed at 37°C under light agitation. At designated time points (30, 60 and 90 min), an aliquot of transport buffer from the receiver chamber was removed and replenished with fresh transport buffer. Samples were quenched with ice-cold ACN containing internal standard and then centrifuged to pellet protein. Resulting supernatants are further diluted with 50/50 ACN/H₂O (H₂O only for Atenolol) and submitted for LC-MS/MS analysis. Reported apparent permeability (P_{app}) values were calculated from single determination. Atenolol and propranolol were tested as low and moderate permeability

references. Bidirectional transport of digoxin was assessed to demonstrate P-gp activity/expression.

PK Studies. A PK study of RTICBM-74 was performed using male Sprague-Dawley rats (Paraza Pharma Inc., Montreal, Canada). RTICBM-74 was suspended in 5% Cremophor, 5% ethanol and 90% saline. On the morning of the PK study, animals were weighed and dosing formulation volumes were calculated accordingly (dose: 10 mg/kg, volume: 10 mL/kg). The compound was injected intraperitoneally (i.p.) to all animals at 10 mg/kg. At selected time points (0.5, 1, 4, 8, and 24 h post-dose), two animals were anesthetized to perform a cardiac puncture to collect blood for plasma analysis, followed by whole body perfusion with phosphate saline buffer (pH 7.4) to wash out any remaining blood from the organs. Brains were harvested and homogenized by polytron 1:4 (w/v) in 25% 2-propanol in water. Brain homogenates were extracted by protein precipitation for drug quantification of LC-MS/MS. Specifically, 25 μ L of brain homogenate and 25 μ L of spiking solution or blank were mixed with 150 μ L of internal standard working solution (0.1 μ M Glyburide/Labetalol in methanol/acetonitrile). The samples were vortexed and centrifuged at 4000 *g* for 15 min at 4 °C. 100 μ L of supernatant was transferred to a 2 mL deepwell plate and diluted with 100 μ L of water. The plates were capped and vortexed. 5 μ L of samples were injected on LC/MS/MS as described above. Plasma samples, standard curve and quality control samples were all treated in the same manner. Pharmacokinetic parameters were calculated using Phoenix WinNonlin (version 8.1, Certana, NJ, US) and the built-in noncompartmental analysis tool. AUC was calculated using linear trapezoidal linear interpolation in WinNonlin.

Self-administration Experiments.

Subjects. Adult male Wistar rats (Jackson Laboratories) and adult male and female Long Evans

rats (Envigo) were used in this study. Rats were pair housed under a 12h light/dark cycle (7am/pm). All experiments were conducted during the light cycle. Animals were continuously monitored and cared for by the veterinary staff of the UNC-Chapel Hill Division of Comparative Medicine. All procedures were carried out in accordance with the NIH Guide for Care and Use of Laboratory Animals and institutional guidelines. All protocols were approved by the UNC Institutional Animal Care and Use Committee (IACUC). UNC-Chapel Hill is accredited by the Association for Assessment and Accreditation of Laboratory Animal Care (AAALAC).

Apparatus. Self-administration was conducted in operant chambers (Med Associates, Georgia, VT) located within sound-attenuating cabinets equipped with an exhaust fan to provide ventilation and mask outside noise. Chambers were equipped with two retractable levers on opposite sides of the chamber (left and right), and a cue light was located above each lever. When the response requirement was met on the left (active) lever, a cue light (directly above the lever) and a stimulus tone were presented for the duration of the alcohol reinforcer delivery (0.1mL of solution into a well on the left side of the chamber across 1.66s). Responding during reinforcer delivery and on the right (inactive) lever was recorded, but had no programmed consequences. Chambers were also equipped with 4 parallel infrared beams across the bar floor to measure general locomotor activity throughout the session. The number of beam breaks for the entire session was collected and this total was divided by the session length (30 min) to represent the locomotor rate (beam breaks/min).

Alcohol self-administration training

Rats were trained to self-administer a 15% (v/v) alcohol on a fixed ratio 2 (FR2) schedule of reinforcement in 30-minute sessions, five days a week (M-F) via sucrose fading as described in (Makhijani et al., 2020; Ornelas et al., 2021; Randall et al., 2017; Randall et al., 2019). Sucrose

fading began with self-administration of 10% sucrose and 2% alcohol (2A/10S), then 5A/10S, 10A/10S, 10A/5S, 15A/5S, 15A/2S on subsequent sessions, and then remained at 15A for the duration of training. Rats had approximately 4 months of self-administration training prior to testing and rats in SA2 and SA3 were used in an unrelated drug test study that concluded a week prior to the initiation of this study.

Self-administration Experiment 1 (SA1): Effect of RTICBM-74 on alcohol and sucrose self-administration in female Long-Evans rats

To measure the effect of RTICBM-74 on maintenance of alcohol and sucrose self-administration, female Long-Evans rats received RTICBM-74 (0, 5, 7.5, 10 mg/kg, IP) 20 minutes prior to a self-administration session. A within subject design was used such that each rat ($n = 11$) received each treatment in a random order. Test days were on Tuesdays and Thursdays with standard self-administration sessions on Monday, Wednesday and Friday. Alcohol lever responses had to be at least 80% of baseline self-administration (average responding in the 2 sessions preceding the test) for a rat to be tested and all rats met this criterion. 1 week after conclusion of testing, the same cohort of rats was trained to self-administer 0.8% (w/v) sucrose solution on a FR2 schedule. Rats ($n = 11$) received the same doses of RTICBM-74 using the same testing procedure and within subject design as when tested with alcohol self-administration.

SA2: Effect of RTICBM-74 on alcohol self-administration in male Wistar rats

To measure the effect of RTICBM-74 on maintenance of alcohol self-administration in different strains or sexes, male Wistar rats received RTICBM-74 (0, 7.5, 10 mg/kg, IP) 20 minutes prior to a self-administration session. A within subject design was used such that each rat ($n = 11$)

received each treatment in a random order. The testing schedule was the same as described in Experiment 1 (SA1).

SA3: Effect of RTICBM-74 on sucrose self-administration in male Wistar rats

A separate cohort of male Wistar rats (n=11) were trained to self-administer 0.5% (w/v) sucrose solution on an FR2 schedule and received the same doses and treatment schedule of RTICBM-74 as described in Experiment 2. A lower concentration of sucrose was used compared to Experiment 1 in order to more closely match total lever presses between alcohol (SA2) and this sucrose self-administration group.

SA4: Effect of rimonabant on alcohol self-administration in Long-Evans rats

To assess the effects of rimonabant on alcohol self-administration, male and female Long-Evans rats received rimonabant (0, 1, 3 mg/kg, IP) 20 minutes prior to an alcohol self-administration session. A within subject design was used such that each rat (n = 12/sex) received each treatment in a random order. The testing schedule was the same as described in Experiment 1.

Self-administration data analysis. Alcohol or sucrose lever responses, inactive lever responses, alcohol or sucrose intake (g/kg or mL/kg, respectively; estimated from the number of reinforcers received) and locomotor rate were analyzed with one-way repeated measures analysis of variance (RM- ANOVA) with RTICBM-74 dose as a factor. Alcohol and sucrose intake data for all experiments, as calculated from number of reinforcers received and body weight, are included in Table 3. Cumulative alcohol lever responses within each session were analyzed by two-way RM-ANOVA with RTICBM-74 dose and session time as factors. In all cases, post-hoc analysis (Tukey) was used to determine specific differences from the vehicle condition. For all analyses, significance was set at $p \leq 0.05$.

RESULTS

ADME Studies. The metabolic stability of RTICBM-74 in rat liver microsomes has previously been reported by us (Nguyen et al., 2017a). In hepatocytes, RTICBM-74 showed a clearance of 14.5 mL/min/kg, with a half-life ($t_{1/2}$) of 461 minutes (Table 1). RTICBM-74 had kinetic solubility of 1.09 μ M in the phosphate-buffered saline (PBS) solution at pH = 7.4. When tested using the MDCK-mdr1 cell line expressing the human P-glycoprotein efflux transporter, RTICBM-74 showed apparent permeability P_{app} values of 0.3 (A-to-B) and 0.2 10^{-6} cm/s (B-to-A), respectively. The efflux ratio was 0.7.

Receptor Selectivity. RTICBM-74 at 10 μ M showed <50% inhibition of all the GPCRs, enzymes or transporters tested.

PK Study. As seen in Table 2 and Figure 1, following an intraperitoneal (i.p.) dose of 10 mg/kg in rats, the concentrations of RTICBM-74 in the plasma decreased between the first and second time points (30 minutes and 1 hour, respectively) and then increased, reaching peak plasma concentration (C_{max} = 70 ng/mL) between 4-8 h post-dose. The brain concentration peaked at 1h (226.1 ng/mL), and decreased and then increased, reaching another high concentration of 109.5 ng/mL at 8 h post dose (Table 2). Higher concentrations in the brain than plasma were observed at all data points tested (0.5, 1, 4, 8, 24h). RTICBM-74 was eliminated from the brain with a clearance of 68.4 mL/min/kg and from the plasma with a clearance of 119.4 mL/min/kg. The overall brain to plasma ratio (K_p) was 1.7, as determined by the AUC_{inf} ratio of brain and plasma.

Self-administration experiments

SA1: The effects of RTICBM-74 on alcohol self-administration in female Long-Evans rats.

A one-way RM ANOVA found a main effect of RTICBM-74 on alcohol lever responses [$F(3, 30)=7.90$, $p<0.0001$] with significant reductions at all three doses (p 's <0.05 ; figure 2A). Across

5 minutes bins (figure 2B), there were main effects of dose [$F(5,50)=81.73$, $p<0.0001$] and time bin [$F(3,30)=7.37$, $p<0.001$] as well as an interaction [$F(15,150)=5.43$, $p<0.0001$]. Post-hoc comparisons revealed that the 7.5 and 10 mg/kg doses reduced responding in the first 5 minutes ($p<0.01$, $p<0.0001$), while all three doses reduced active lever responses from the second time bin onward (p 's <0.0001). Very few inactive lever presses were seen and no significant differences were found (vehicle: 0.17 ± 0.17 ; 5 mg/kg: 0.25 ± 0.13 ; 7.5 mg/kg: 0.08 ± 0.08 ; 10 mg/kg: 0.92 ± 0.92). In this experiment, SA1 (sucrose) and SA2-4, self-administration returned to baseline levels on the intervening days indicating no lasting effects of the drug treatments. In locomotor activity there was an effect of treatment [$F(3,30)=14.60$, $p<0.0001$] with a reduction found only with the highest dose ($p<0.0001$; figure 1C), indicating a possible motor impairment at the highest dose. This finding suggests that the reduction in self-administration at this 10 mg/kg dose may be a nonspecific reduction related to a motor impairment.

SA1: The effects of RTICBM-74 on sucrose self-administration in female Long-Evans rats.

In sucrose self-administration there was a main effect of drug treatment [$F(3,30)=5.19$, $p<0.01$] with a reduction found only in the 10 mg/kg group ($p<0.05$; figure 2D). A similar result was found with locomotor activity [$F(3,30)=65.46$, $p<0.01$] as again a reduction was found with treatment of the highest dose ($p<0.0001$; figure 2F), confirming a likely motor impairment at this dose. Further, the lack of a reduction of sucrose self-administration at the 5 or 7.5 mg/kg dose suggests that the reductions in alcohol self-administration at those doses are specific to alcohol reinforcement. At all doses in this experiment no inactive lever responses were recorded which may have been due to a program error, however in all other experiments inactive lever responses counts were very low and did not differ between groups.

SA2: The effects of RTICBM-74 on alcohol self-administration in male Wistar rats. A one-way RM ANOVA found that RTICBM-74 reduced lever responses [$F(2, 20)=4.79$, $p<0.05$] at the 7.5 and 10 mg/kg doses (p 's < 0.05 ; figure 3A). Across time (figure 3B) there was a main effect of time bin [$F(5,165)=64.83$, $p<0.0001$], a trend for an effect of dose ($p=0.085$), and a significant interaction [$F(10,65)=2.41$, $p<0.05$]. Post-hoc comparisons revealed that the 7.5 mg/kg dose was lower than vehicle at the 15 min time bin ($p<0.05$), while the 10 mg/kg dose reduced responding at 20, 25, and 30 min (p 's <0.05). RTICBM-74 did not affect locomotor rate but approached a statistical trend ($p=0.079$; figure 3C). No significant differences in inactive lever presses were found (vehicle: 1.92 ± 0.54 ; 7.5 mg/kg: 3.00 ± 0.74 ; 10 mg/kg: 2.50 ± 0.72).

SA3: The effects of RTICBM-74 on sucrose self-administration in male Wistar rats.

RTICBM-74 had no significant effects on behaviors in sucrose self-administration or locomotion (figures 3D-F), suggesting specificity of the reduction to the alcohol reinforcer. No significant differences in inactive lever presses were found (vehicle: 2.55 ± 0.90 ; 7.5 mg/kg: 1.72 ± 0.94 ; 10 mg/kg: 1.82 ± 1.07).

SA4: The effects of rimonabant on alcohol self-administration in male and female Long Evans rats. In males, rimonabant reduced alcohol lever responding [$F(2, 22)=6.22$, $p<0.01$] at 1 and 3 mg/kg (p 's <0.05 ; figure 4A). A two-way RM ANOVA analyzing lever responses in 5 minute bins (figure 4B) found main effects of rimonabant [$F(2,33)=5.35$, $p<0.01$] and time bin [$F(5,165)=44.23$, $p<0.0001$] with a significant interaction [$F(10,165)=2.33$, $p<0.05$]. Post-hoc analyses indicated that both doses of rimonabant reduced the total number of lever responses starting with the third time bin (p 's < 0.05). In females, rimonabant also reduced lever pressing for alcohol [$F(2,22)=15.13$, $p<0.0001$] at the 1 ($p<0.01$) and 3 ($p<0.0001$) mg/kg doses (figure

4D). When analyzed across the session (figure 4E) there were main effects of rimonabant [$F(2,22)=18.06$, $p<0.0001$], time bin [$F(5,55)=21.66$, $p<0.0001$], and a significant interaction [$F(10,110)=2.63$, $p<0.01$]. Post-hoc comparisons found that both doses of rimonabant reduced lever pressing at every time bin compared to vehicle (p 's <0.01). There was a main effect of rimonabant on locomotor activity [$F(2,22)=8.92$, $p<0.01$] where the 3 mg/kg dose increased the number of beam breaks/min ($p<0.01$; figure 4F). No significant differences in inactive lever presses were found in both males (veh: 1.33 ± 0.43 ; 1 mg/kg: 0.58 ± 0.040 ; 3 mg/kg: 1.50 ± 0.65) and females (veh: 1.50 ± 0.86 ; 1 mg/kg 0.83 ± 0.52 ; 3 mg/kg 1.00 ± 0.43). In sum, rimonabant pretreatment, which was used as a positive control, induced reductions in alcohol self-administration in males and females consistent with results previously found in males (Colombo et al., 2007; Freedland et al., 2001; Maccioni et al., 2009).

DISCUSSION

The present report evaluated the pharmacokinetic and behavioral characteristics of the potent CB₁ NAM, RTICBM-74. RTICBM-74 showed good in vitro potency at the CB₁ receptor, with IC₅₀ values of 23 nM in calcium mobilization assay and 153 nM in [³⁵S]GTPγS assay (Nguyen et al., 2017a). In rat liver microsomes, RTICBM-74 displayed excellent metabolic stability with long half-life (>300 min) and low clearance (<4.6 μL/min/mg) (Nguyen et al., 2017a). In hepatocytes, similar high stability was observed with RTICBM-74, with half-life of 461 min and low clearance (14.5 μL/min/mg). However, poor aqueous solubility (1.09 μM) and high plasma protein binding (99.9%) were observed. These are consistent with the high lipophilicity and low solubility of many cannabinoid ligands including rimonabant (solubility of 2.6 μM at pH 7.4) (Alelyunas et al., 2010). The high protein binding may have, in part,

contributed to the observed good metabolic stability of RTICBM-74, with only small fractions of the compounds being unbound and available to be metabolized. This low unbound drug concentration may have also resulted in the modest apparent permeability values (P_{app} of 0.3 (A-to-B) and 0.2×10^{-6} cm/s (B-to-A)) observed in the MDCK-mdr1 assay. The efflux ratio of 0.7 suggests that RTICBM-74 was not a P-glycoprotein substrate.

The low P_{app} values in the MDCK-mdr1 assay may suggest low brain penetration for RTICBM-74; however, higher concentrations in brain than plasma at all time points and a significant higher brain C_{max} than plasma (272.0 ng/mL vs. 70.2 ng/mL) were observed in the PK study in rats. In addition, a brain/plasma ratio ($K_p = AUC_{brain}/AUC_{plasma}$) of 1.7 was obtained. These results support good brain penetration for RTICBM-74. Brain concentrations were at high levels at the early time points (0.5 and 1 h) and remained high through the testing period (>10 ng/mL in brain at 24h), confirming the excellent in vivo metabolic stability of RTICBM-74. Interestingly, following the high concentrations at 1h, the brain concentrations decreased but then increased, reaching another high concentration at 8h, rendering the large T_{max} value and variance (4.5 ± 3.5 h). This is again likely because of the high protein binding of RTICBM-74 and the resulting low unbound drug levels in both plasma and brain, which is also consistent with the relatively low plasma concentrations. These results also suggest possible sequestration into adipose tissue, a known phenomenon with the generally lipophilic cannabinoid compounds such as Δ^9 -tetrahydrocannabinol (Johansson et al., 1989; Kreuz & Axelrod, 1973). RTICBM-74 was cleared from the brain and plasma at moderate or high rate at 68.4 and 119.4 mL/min/kg, respectively. Together, these results indicate RTICBM-74 is a brain-penetrant CB₁ allosteric modulator, albeit with low penetration rate. The ADME and PK studies were performed in rats to be consistent with the behavioral studies that were also conducted in rats. It has been shown that

in vivo human and rat pharmacokinetic data are highly correlated (Caldwell et al., 2004). Only male Sprague-Dawley rats were used in the PK studies; it is possible that female rats might exhibit a slightly different profile.

The off-target screen using orthosteric radioligands showed that RTICBM-74 is highly selective to CB₁. Radioligand binding studies are commonly conducted for target selectivity determination for orthosteric ligands as well as allosteric modulators, which potentiate or attenuate binding affinity of orthosteric agonists. However, signal changes caused by allosteric modulators are sometimes less pronounced (e.g. <50%, a cut-off often used in binding studies for orthosteric ligands), and therefore they may not be identified in these radioligand binding assays. In addition, it is possible that RTICBM-74 may act as an allosteric agonist at other targets, which bind to the allosteric sites and activate the receptors by themselves, and functional assays will be required for such determination.

RTICBM-74 administered 20 minutes prior to alcohol self-administration attenuated alcohol intake. All three doses tested (5, 7.5, 10 mg/kg) reduced alcohol self-administration in Long-Evans female rats (figure 2A), and both the highest doses reduced initial intake in the first 5 minute of the session (figure 2B), which could be interpreted as a reduction in initiation to drink. However, once the session progressed and alcohol was consumed, a greater reduction in ongoing alcohol self-administration was observed, suggesting that RTICBM-74 is more effective at reducing ongoing drinking. General locomotor activity was decreased only at the highest dose which may have contributed to the reduction in self-administration at that dose (figure 2C). However, as locomotor activity was not reduced at lower doses (5 and 7.5 mg/kg) the reduced consumption was likely not due to a non-specific motor reduction. The effects of RTICBM-74 were tested on sucrose self-administration to test for reinforcer specificity, and a reduction in

sucrose intake was only seen at the highest dose (10 mg/kg; figure 2D), which again induced a concordant reduction in locomotor activity that suggests general motor impairments (figure 2F).

To examine replicability and generalizability of these effects across sex and strain, we tested the two highest doses of RTICBM-74 using the same self-administration procedures in male Wistar rats. Again, all doses reduced alcohol self-administration (figure 3A). In male Wistar rats the effect was not immediately detected but emerged after 15 (10 mg/kg) or 20 (7.5 mg/kg) minutes of self-administration (figure 3B), again supporting that RTICBM-74 is more effective at reducing ongoing drinking. Notably, there was no reduction in locomotor activity at the highest dose (10 mg/kg) as was seen in the Long-Evans females (figure 3C), suggesting species differences. Likewise, RTICBM-74 had no effect on sucrose self-administration (figure 3D-F). The Wistar male rats were less sensitive to the locomotor reduction at the highest dose as compared to the Long-Evans females, but notably the 7.5 mg/kg dose effectively and specifically reduced alcohol self-administration in both groups.

Rimonabant has been shown to reduce operant alcohol-self administration across rat strains (Colombo et al., 2004; Freedland et al., 2001; Hansson et al., 2007; Malinen & Hyttiä, 2008). We therefore tested rimonabant as the positive control experiment on alcohol self-administration. As expected, both low and high (1 and 3 mg/kg) doses of rimonabant effectively reduced alcohol lever responses (figure 4A,D) in male and female Long Evans rats. Across time a similar pattern was seen as in prior experiments where the reduction became significant over time in males but was immediately apparent in females (figure 4B,E). Rimonabant did not reduce locomotor activity at either dose, although the high dose increased locomotor activity in females. This increase was unexpected as few studies have tested females, but the increase in generalized activity did not affect the expected reduction in alcohol intake. Overall, we found that RTICBM-

74 effectively reduced alcohol intake as effectively as rimonabant across multiple rat strains and in both sexes, and showed that the reduction effect was specific to alcohol.

In summary, the present studies were designed to characterize RTICBM-74 towards the goal of developing a novel CB₁ NAM-based treatment for AUD. RTICBM-74 was found to have good *in vitro* potency at the CB₁ receptor, excellent metabolic stability, and effective CNS penetration, all of which suggest efficacy as a centrally-acting peripherally-administered treatment. IP administration of RTICBM-74 effectively and specifically reduced alcohol self-administration in both male and female rats across multiple strains, without affecting locomotor activity or sucrose intake. Altogether, the findings suggest that CB₁ NAMs such as RTICBM-74 may have therapeutic potential in treatment of AUD.

Conflict of interest: none.

Acknowledgements

We are grateful to the NIMH Psychoactive Drug Screening Program (PDSP) for the receptor selectivity testing and to Brayden Fortino for technical assistance with the self-administration studies.

Authorship Contributions

Participated in research design: Zhang, Besheer.

Conducted experiments: Lovelock, Nguyen, Voorhies.

Performed data analysis: Lovelock, Nguyen, Zhang, Besheer.

Wrote or contributed to the writing of the manuscript: Lovelock, Nguyen, Zhang, Besheer.

References:

- Alelyunas YW, Empfield JR, McCarthy D, Spreen RC, Bui K, Pelosi-Kilby L, & Shen C (2010). Experimental solubility profiling of marketed CNS drugs, exploring solubility limit of CNS discovery candidate. *Bioorg Med Chem Lett* 20: 7312-7316.
- Arnone M, Maruani J, Chaperon F, Thiebot MH, Poncelet M, Soubrie P, & Le Fur G (1997). Selective inhibition of sucrose and ethanol intake by SR 141716, an antagonist of central cannabinoid (CB1) receptors. *Psychopharmacology (Berl)* 132: 104-106.
- Beyer CE, Dwyer JM, Piesla MJ, Platt BJ, Shen R, Rahman Z, Chan K, Manners MT, Samad TA, Kennedy JD, Bingham B, & Whiteside GT (2010). Depression-like phenotype following chronic CB1 receptor antagonism. *Neurobiol Dis* 39: 148-155.
- Cahill K, & Ussher M (2007). Cannabinoid type 1 receptor antagonists (rimonabant) for smoking cessation. *Cochrane Database Syst Rev*: CD005353.
- Caldwell GW, Masucci JA, Yan Z, & Hageman W (2004). Allometric scaling of pharmacokinetic parameters in drug discovery: can human CL, Vss and t1/2 be predicted from in-vivo rat data? *Eur J Drug Metab Pharmacokinet* 29: 133-143.
- Christensen R, Kristensen PK, Bartels EM, Bliddal H, & Astrup A (2007). Efficacy and safety of the weight-loss drug rimonabant: a meta-analysis of randomised trials. *Lancet* 370: 1706-1713.
- Cinar R, Iyer MR, & Kunos G (2020). The therapeutic potential of second and third generation CB1R antagonists. *Pharmacol Ther*: 107477.
- Cippitelli A, Bilbao A, Hansson AC, del Arco I, Sommer W, Heilig M, Massi M, Bermudez-Silva FJ, Navarro M, Ciccocioppo R, de Fonseca FR, & European TC (2005). Cannabinoid CB1 receptor antagonism reduces conditioned reinstatement of ethanol-seeking behavior in rats. *Eur J Neurosci* 21: 2243-2251.
- Colombo G, Orrù A, Lai P, Cabras C, Maccioni P, Rubio M, Gessa GL, & Carai MA (2007). The cannabinoid CB 1 receptor antagonist, rimonabant, as a promising pharmacotherapy for alcohol dependence: Preclinical evidence. *Molecular neurobiology* 36: 102-112.
- Colombo G, Vacca G, Serra S, Carai MA, & Gessa GL (2004). Suppressing effect of the cannabinoid CB1 receptor antagonist, SR 141716, on alcohol's motivational properties in alcohol-preferring rats. *Eur J Pharmacol* 498: 119-123.
- Economidou D, Mattioli L, Cifani C, Perfumi M, Massi M, Cuomo V, Trabace L, & Ciccocioppo R (2006). Effect of the cannabinoid CB1 receptor antagonist SR-141716A on ethanol self-administration and ethanol-seeking behaviour in rats. *Psychopharmacology (Berl)* 183: 394-403.

Economidou D, Mattioli L, Ubaldi M, Lourdasamy A, Soverchia L, Hardiman G, Campolongo P, Cuomo V, & Ciccocioppo R (2007). Role of cannabinoidergic mechanisms in ethanol self-administration and ethanol seeking in rat adult offspring following perinatal exposure to Delta9-tetrahydrocannabinol. *Toxicol Appl Pharmacol* 223: 73-85.

Freedland C, Sharpe A, Samson H, & Porrino L (2001). Effects of SR141716A on ethanol and sucrose self-administration. *Alcoholism: Clinical and Experimental Research* 25: 277-282.

Gallate JE, Saharov T, Mallet PE, & McGregor IS (1999). Increased motivation for beer in rats following administration of a cannabinoid CB1 receptor agonist. *Eur J Pharmacol* 370: 233-240.

Hansen HH, Krutz B, Sifringer M, Stefovská V, Bittigau P, Pragst F, Marsicano G, Lutz B, & Ikonomidou C (2008). Cannabinoids enhance susceptibility of immature brain to ethanol neurotoxicity. *Ann Neurol* 64: 42-52.

Hansson AC, Bermúdez-Silva FJ, Malinen H, Hyytiä P, Sanchez-Vera I, Rimondini R, de Fonseca FR, Kunos G, Sommer WH, & Heilig M (2007). Genetic impairment of frontocortical endocannabinoid degradation and high alcohol preference. *Neuropsychopharmacology* 32: 117-126.

Heilig M, & Egli M (2006). Pharmacological treatment of alcohol dependence: target symptoms and target mechanisms. *Pharmacol Ther* 111: 855-876.

Horswill JG, Bali U, Shaaban S, Keily JF, Jeevaratnam P, Babbs AJ, Reynet C, & Wong Kai In P (2007). PSNCBAM-1, a novel allosteric antagonist at cannabinoid CB1 receptors with hypophagic effects in rats. *Br J Pharmacol* 152: 805-814.

Houchi H, Babovic D, Pierrefiche O, Ledent C, Daoust M, & Naassila M (2005). CB1 receptor knockout mice display reduced ethanol-induced conditioned place preference and increased striatal dopamine D2 receptors. *Neuropsychopharmacology* 30: 339-349.

Jing L, Qiu Y, Zhang Y, & Li JX (2014). Effects of the cannabinoid CB(1) receptor allosteric modulator ORG 27569 on reinstatement of cocaine- and methamphetamine-seeking behavior in rats. *Drug and Alcohol Dependence* 143: 251-256.

Johansson E, Noren K, Sjoval J, & Halldin MM (1989). Determination of delta 1-tetrahydrocannabinol in human fat biopsies from marihuana users by gas chromatography-mass spectrometry. *Biomed Chromatogr* 3: 35-38.

Kenakin T, & Miller LJ (2010). Seven transmembrane receptors as shapeshifting proteins: the impact of allosteric modulation and functional selectivity on new drug discovery. *Pharmacol Rev* 62: 265-304.

Koob GF (2013). Theoretical frameworks and mechanistic aspects of alcohol addiction: alcohol addiction as a reward deficit disorder. *Curr Top Behav Neurosci* 13: 3-30.

Kreuz DS, & Axelrod J (1973). Delta-9-tetrahydrocannabinol: localization in body fat. *Science* 179: 391-393.

Maccioni P, Fantini N, Carai MA, Gessa GL, & Colombo G (2009). Suppressing effect of the cannabinoid CB1 receptor antagonist, rimonabant, on alcohol self-administration in alcohol-preferring rats. *The Open Neuropsychopharmacology Journal* 2.

Makhijani VH, Irukulapati P, Van Voorhies K, Fortino B, & Besheer J (2020). Central amygdala mineralocorticoid receptors modulate alcohol self-administration. *Neuropharmacology* 181: 108337.

Malinen H, & Hyttiä P (2008). Ethanol self-administration is regulated by CB1 receptors in the nucleus accumbens and ventral tegmental area in alcohol-preferring AA rats. *Alcoholism: Clinical and Experimental Research* 32: 1976-1983.

Moreira FA, Grieb M, & Lutz B (2009). Central side-effects of therapies based on CB1 cannabinoid receptor agonists and antagonists: focus on anxiety and depression. *Best Pract Res Clin Endocrinol Metab* 23: 133-144.

Nguyen T, German N, Decker AM, Langston TL, Gamage TF, Farquhar CE, Li JX, Wiley JL, Thomas BF, & Zhang Y (2017a). Novel Diarylurea Based Allosteric Modulators of the Cannabinoid CB1 Receptor: Evaluation of Importance of 6-Pyrrolidinylpyridinyl Substitution. *J Med Chem* 60: 7410-7424.

Nguyen T, Li JX, Thomas BF, Wiley JL, Kenakin TP, & Zhang Y (2017b). Allosteric Modulation: An Alternate Approach Targeting the Cannabinoid CB1 Receptor. *Med Res Rev* 37: 441-474.

Nguyen T, Thomas BF, & Zhang Y (2019). Overcoming the Psychiatric Side Effects of the Cannabinoid CB1 Receptor Antagonists: Current Approaches for Therapeutics Development. *Curr Top Med Chem* 19: 1418-1435.

Ornelas LC, Tyler RE, Irukulapati P, Paladugu S, & Besheer J (2021). Increased alcohol self-administration following exposure to the predator odor TMT in active coping female rats. *Behavioural Brain Research* 402: 113068.

Parsons LH, & Hurd YL (2015). Endocannabinoid signalling in reward and addiction. *Nat Rev Neurosci* 16: 579-594.

Price MR, Baillie GL, Thomas A, Stevenson LA, Easson M, Goodwin R, McLean A, McIntosh L, Goodwin G, Walker G, Westwood P, Marrs J, Thomson F, Cowley P, Christopoulos A, Pertwee RG, & Ross RA (2005). Allosteric modulation of the cannabinoid CB1 receptor. *Mol Pharmacol* 68: 1484-1495.

Randall PA, Stewart RT, & Besheer J (2017). Sex differences in alcohol self-administration and relapse-like behavior in Long-Evans rats. *Pharmacol Biochem Behav* 156: 1-9.

Randall PA, Vetreno RP, Makhijani VH, Crews FT, & Besheer J (2019). The toll-like receptor 3 agonist poly (I: C) induces rapid and lasting changes in gene expression related to glutamatergic function and increases ethanol self-administration in rats. *Alcoholism: Clinical and Experimental Research* 43: 48-60.

Serra S, Brunetti G, Pani M, Vacca G, Carai MA, Gessa GL, & Colombo G (2002). Blockade by the cannabinoid CB(1) receptor antagonist, SR 141716, of alcohol deprivation effect in alcohol-preferring rats. *Eur J Pharmacol* 443: 95-97.

Vinod KY, Maccioni P, Garcia-Gutierrez MS, Femenia T, Xie S, Carai MA, Manzanares J, Cooper TB, Hungund BL, & Colombo G (2012). Innate difference in the endocannabinoid signaling and its modulation by alcohol consumption in alcohol-preferring sP rats. *Addict Biol* 17: 62-75.

Vinod KY, Yalamanchili R, Thanos PK, Vadasz C, Cooper TB, Volkow ND, & Hungund BL (2008). Genetic and pharmacological manipulations of the CB(1) receptor alter ethanol preference and dependence in ethanol preferring and nonpreferring mice. *Synapse* 62: 574-581.

Footnotes

This work was supported in part by the National Institute on Alcohol Abuse and Alcoholism [AA011605, JB and AA026537, JB]; National Institute on Drug Abuse [DA040693, YZ] and the UNC Bowles Center for Alcohol Studies

The authors declare they have no conflicts of interest with the contents of this article.

Legends for Figures

Figure 1: Brain and plasma concentrations of RTICBM-74 over time following 10 mg/kg intraperitoneal injection to male Sprague-Dawley rats. Please note log scale is used on the Y-axis.

Figure 2: Experiment SA1 found that RTICBM-74 reduced alcohol lever responses in female Long-Evans rats at all doses tested (A,B) while locomotor rate was only reduced by the highest dose (C). Only the highest dose reduced lever responding for a sucrose reward (D,E), which is likely explained by the generalized reduction in motor activity at this dose (F). * $p < 0.05$

Figure 3: In Experiment SA2, RTICBM-74 reduced lever responses for alcohol in Male Wistar rats (A,B) and had no effect on locomotor rate during the test session (C). Experiment SA3 examined the effect of RTICBM-74 on self-administration for sucrose finding no effects (D-F), suggesting the results from Experiment SA2 are reinforcer-specific. * $p < 0.05$

Figure 4: Experiment SA4 tested rimonabant as a positive control experiment in male and female Long-Evans rats. As expected, rimonabant reduced lever responding in both males (A,B) and females (D,E). The high dose of rimonabant increased locomotor activity during the test session in females (F), though this did not result in a concordant increase in lever responses. * $p < 0.05$

Tables

Table 1. ADME properties of RTICBM-74

Rat Liver Microsome ^a		Hepatocyte		Kinetic solubility (mM)	%Protein bound	MDCK-mdr1		
t _{1/2} (min)	Cl _{int} (mL/min/kg)	t _{1/2} (min)	Cl _{int} (mL/min/kg)			P _{app} (A to B) (10 ⁻⁶ cm/s)	P _{app} (B to A) (10 ⁻⁶ cm/s)	efflux ratio
>300	<4.6	461 ± 38	14.5 ± 1.2	1.09	99.9% ± 0.03	0.3 ± 0.05	0.2 ± 0.05	0.7

^a Reported previously in (Nguyen et al., 2017a).

Data are shown mean ± standard error of the mean.

Table 2. Pharmacokinetic analysis of RTICBM-74 in rats (i.p. 10 mg/kg)

	C_{max} (ng/mL)	t_{max} (h)	CL_F (mL/min/kg)	AUC_{inf} (ng/mL*h)
Plasma	70.2 ± 1.9	6.0 ± 2.0	119.4 ± 27.9	1476.2 ± 344.9
Brain	272.0 ± 126.9	4.5 ± 3.5	68.4 ± 9.2	2481.8 ± 332.1
Brain/plasma ratio				1.7

Data are shown as mean ± standard error of the mean.

Table 3. Alcohol intake from self-administration experiments.

RTICBM-74	Self-admin	Vehicle	5 mg/kg	7.5 mg/kg	10 mg/kg	Statistics
SA1: Long-Evans females	♀ Alcohol	0.53 ± 0.10	0.40 ± 0.07*	0.42 ± 0.06*	0.17 ± 0.07*	F(3,30) = 9.18, p<0.001
	♀ Sucrose	7.90 ± 2.83	8.21 ± 2.65	8.48 ± 3.81	5.38 ± 1.79*	F(3,30)=5.52, p<0.01
SA2: Wistar males	♂ Alcohol	0.65 ± 0.66	N/A	0.43 ± 0.94*	0.44 ± .088*	F(2,20) = 4.07, p<0.01
SA3: Wistar males	♂ Sucrose	12.28 ± 2.08	N/A	10.88 ± 1.80	8.37 ± 3.02	F(2,20) = 1.96, p>0.05

Rimonabant	Self-admin	Vehicle	1 mg/kg	3 mg/kg		Statistics
SA4: Long-Evans males & females	♀ Alcohol	0.82 ± 0.08	0.52 ± 0.76*	0.39 ± 0.68*		F(2,22) = 17.35, p<0.0001
	♂ Alcohol	0.73 ± 0.11	0.45 ± 0.05*	0.38 ± 0.07*		F(2,22) = 6.96, p<0.01

Data are shown as g/kg (alcohol) or ml/kg (sucrose) ± standard error of the mean. Bold in statistics column indicates a main effect of drug treatment, * p<0.05 difference from vehicle.

Figures

Figure 1

Figure 2

Figure 3

Figure 4