

Teaching an Old Disorder New Tricks: Endocrine Pharmacotherapies for Alcohol Use Disorder

Rani Richardson, B.A.

Stephanie T. Weiss, M.D., Ph.D.

Andras Leko, M.D., Ph.D.

Mehdi Farokhnia, M.D., M.P.H.

**Translational Addiction Medicine Branch, Intramural Research Program
National Institute on Drug Abuse, National Institutes of Health**



Disclosure Information

- ◆ **Presenter 1: Rani Richardson, B.A.**
 - ◆ **No Disclosures**
- ◆ **Presenter 2: Stephanie T. Weiss, M.D., Ph.D.**
 - ◆ **No Disclosures**
- ◆ **Presenter 3: Andras Leko, M.D., Ph.D.**
 - ◆ **No Disclosures**
- ◆ **Presenter 4: Mehdi Farokhnia, M.D., M.P.H.**
 - ◆ **No Disclosures**

Learning Objectives

- ◆ Review AUD treatments currently used in the U.S.
- ◆ Describe ongoing, cutting-edge translational and clinical research to develop novel AUD therapies
- ◆ Illustrate the early-phase clinical trial process using case examples
- ◆ Identify barriers and complexities in the AUD medication development process

Current Therapies for AUD



Alcohol Use Disorder (AUD)

- ◆ **15 million American adults living with AUD**
- ◆ **High burden of morbidity and mortality, comorbidities**
- ◆ **Only 3 FDA-approved treatments for AUD**
 - ◆ **Heterogeneity in the AUD population**



Litten RZ, Handb Exp Pharmacol 2018, Grant BF, JAMA Psychiatry 2015; National Survey on Drug Use and Health Koob GF, Pharmacol Rev 2021; The Healthcare Professional's Core Resource on Alcohol (NIAAA)

NIAAA Clinician Guidelines

“All approved drugs have been shown to be effective adjuncts to the treatment of alcohol dependence. Thus, consider adding medication whenever you are treating someone with active alcohol dependence or someone who has stopped drinking in the past few months but is experiencing problems such as craving or slips.”



NIAAA NIH Government Publications.
*Helping Patients Who Drink Too Much: A
Clinician's Guide.* 2005, 2007 Edition.

FDA-Approved AUD Pharmacotherapies

◆ Disulfiram

- ◆ Aldehyde dehydrogenase inhibitor
- ◆ Causes nausea, dizziness, headache, flushing when taken with alcohol
- ◆ Approved by FDA in 1948

◆ Naltrexone

- ◆ Opioid antagonist
- ◆ Binds to opioid receptors, thus blocking alcohol reward pathways
- ◆ Approved by FDA in 1994

◆ SR-Naltrexone

- ◆ Sustained-release naltrexone
- ◆ Approved by FDA in 2006

◆ Acamprosate

- ◆ Glutamate receptor modulator (?)
- ◆ Helps maintain complete abstinence during post-acute withdrawal
- ◆ Approved by FDA in 2004

Acamprosate

- ◆ Abstinence, relapse prevention
- ◆ Drug metabolism
- ◆ Compliance

Naltrexone

- ◆ Multiple outcomes
 - ◆ Craving
 - ◆ Number of heavy drinking days
 - ◆ Return to drinking
 - ◆ Binge drinking

Disulfiram

- ◆ Complete abstinence, not drinking reduction
- ◆ Supervised administration
- ◆ Difficulty in study design

- ◆ Liver considerations

Promising Medications for AUD (not approved by the FDA)

Topiramate

- ◆ Increases GABA-A activity
- ◆ Approved by FDA for epilepsy, migraine and obesity

Gabapentin

- ◆ GABA analog, voltage-gated calcium channels
- ◆ Approved by FDA for epilepsy and postherpetic neuralgia

Baclofen

- ◆ GABA-B receptor agonist
- ◆ Approved by FDA for muscle spasticity

Topiramate

- ◆ Reduction of craving
- ◆ Reduction of heavy drinking days, drinks per day



Miranda R, Jr. *Addict Biol* 2016; Johnson BA, *JAMA* 2007; Johnson BA, *Lancet* 2003, Palpacuer C, *Addiction*, 2018; Wetherill RR, *Neuropsychopharmacology* 2021; Kranzler HR, *Addict Biol.* 2022

Gabapentin

- ◆ Heavy drinking days
- ◆ Responders: withdrawal symptoms

Baclofen

- ◆ Craving, abstinence, time to relapse
- ◆ High drinking individuals
- ◆ Comorbid liver disease

Varenicline

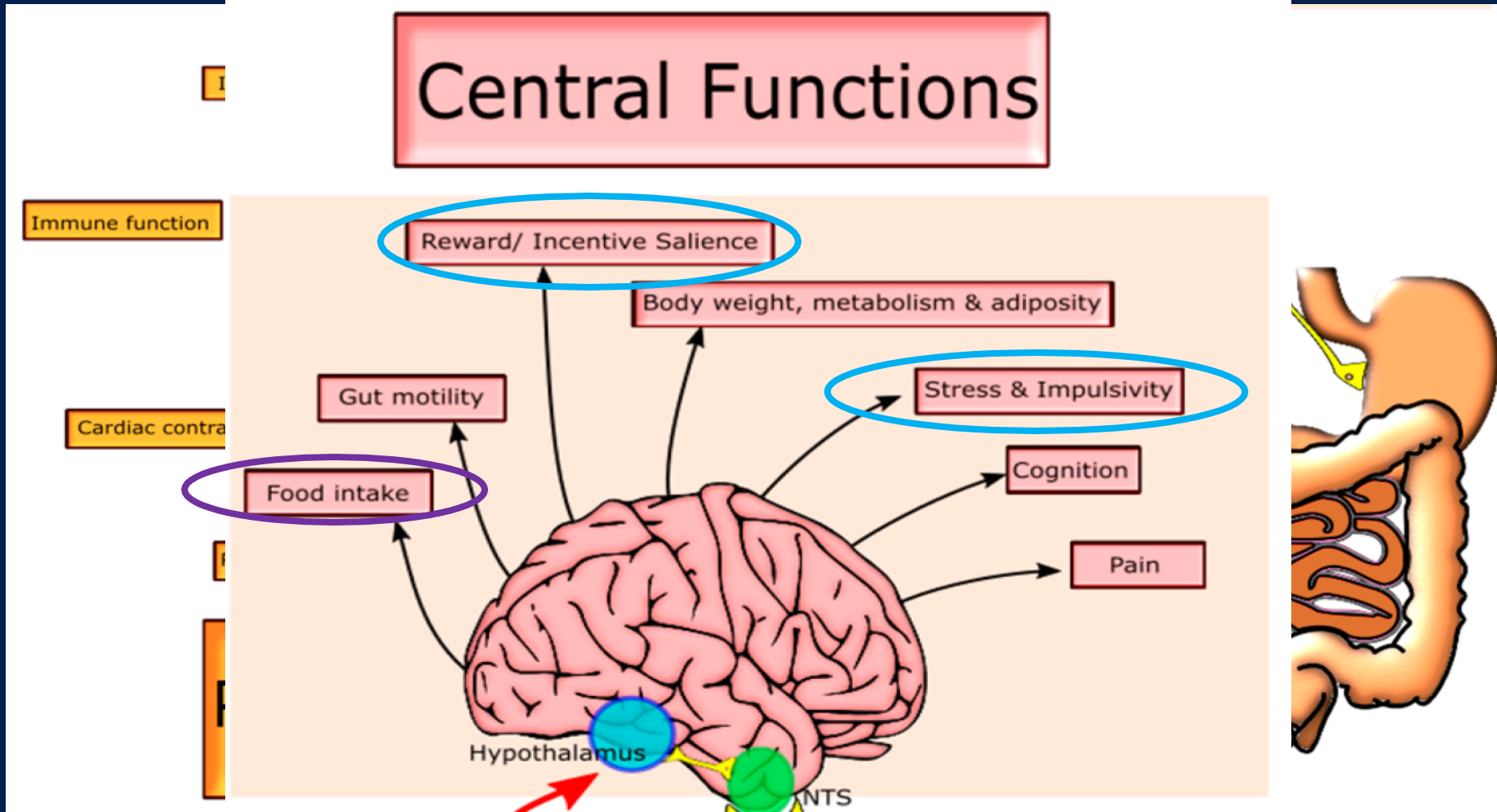
- ◆ Craving
- ◆ Heavy Drinking Days
- ◆ Comorbid smokers (nicotine)

Drug	Advantages	Considerations	References
Acamprosate	Abstinence, relapse prevention, no hepatic metabolism	3x daily admin (compliance)	Anton RF, JAMA 2006; Mann K, Addict Biol 2013; Cheng HY, Bmj. 2020; Maisel NC, Addiction 2013; Jonas DE, JAMA 2014
Naltrexone	Craving, # heavy drinking days, binge drinking, return to drinking	Requires abstinence from opioids prior to initiation	Anton RF, JAMA 2006; Mann K, Addict Biol. 2013; Cheng HY, BMJ. 2020; Maisel NC, Addiction 2013; Jonas DE, JAMA 2014
Disulfiram	Complete abstinence, not reduction	Supervised admin, study design difficult, potential hepatotoxicity	Jørgensen CH, Alcohol Clin Exp Res 2011; Skinner MD, PLoS One 2014
Topiramate	Craving, heavy drinking days, drinks per day	Potential cognitive dysfunction	Miranda R, Jr. Addict Biol 2016; Johnson BA, JAMA 2007; Johnson BA, Lancet 2003, Palpacuer C, Addiction, 2018; Wetherill RR, Neuropsychopharmacology 2021; Kranzler HR, Addict Biol. 2022
Gabapentin	Withdrawal symptoms (responders), heavy drinking days	Potential sedation, drowsiness	Furieri FA, J Clin Psychiatry; Karam-Hage M, Am J Psychiatry; Myrick H, Am J Psychiatry. 1998; Karam-Hage M, 2003; Mason BJ, JAMA Intern Med, 2014; Anton RF, JAMA Intern Med 2020; Falk DE, ACER 2019
Baclofen	Time to relapse, high drinking, comorbid liver disease, dose dependent	Potential sedation, drowsiness	Addolorato G, Alcohol Alcohol 2002; Leggio L, CNS Neurol Disord Drug Targets 2010; Agabio R, 2018; Pierce M, Eur Neuropsychopharmacol 2018; Rombouts SA, Alcohol Alcohol 2019; Reynaud M, Alcohol Alcohol 2017
Varenicline	Craving, heavy drinking days, comorbid nicotine		Litten RZ, J Addict Med. 2013; Falk DE, J Addict Med. 2015; Donato S, Alcohol Clin Exp Res. 2021; de Bejczy A, Alcohol Clin Exp Res. 2015; Gandhi KD, J Clin Psychiatry. 2020

Development of the GLWL-01 Investigational New Drug

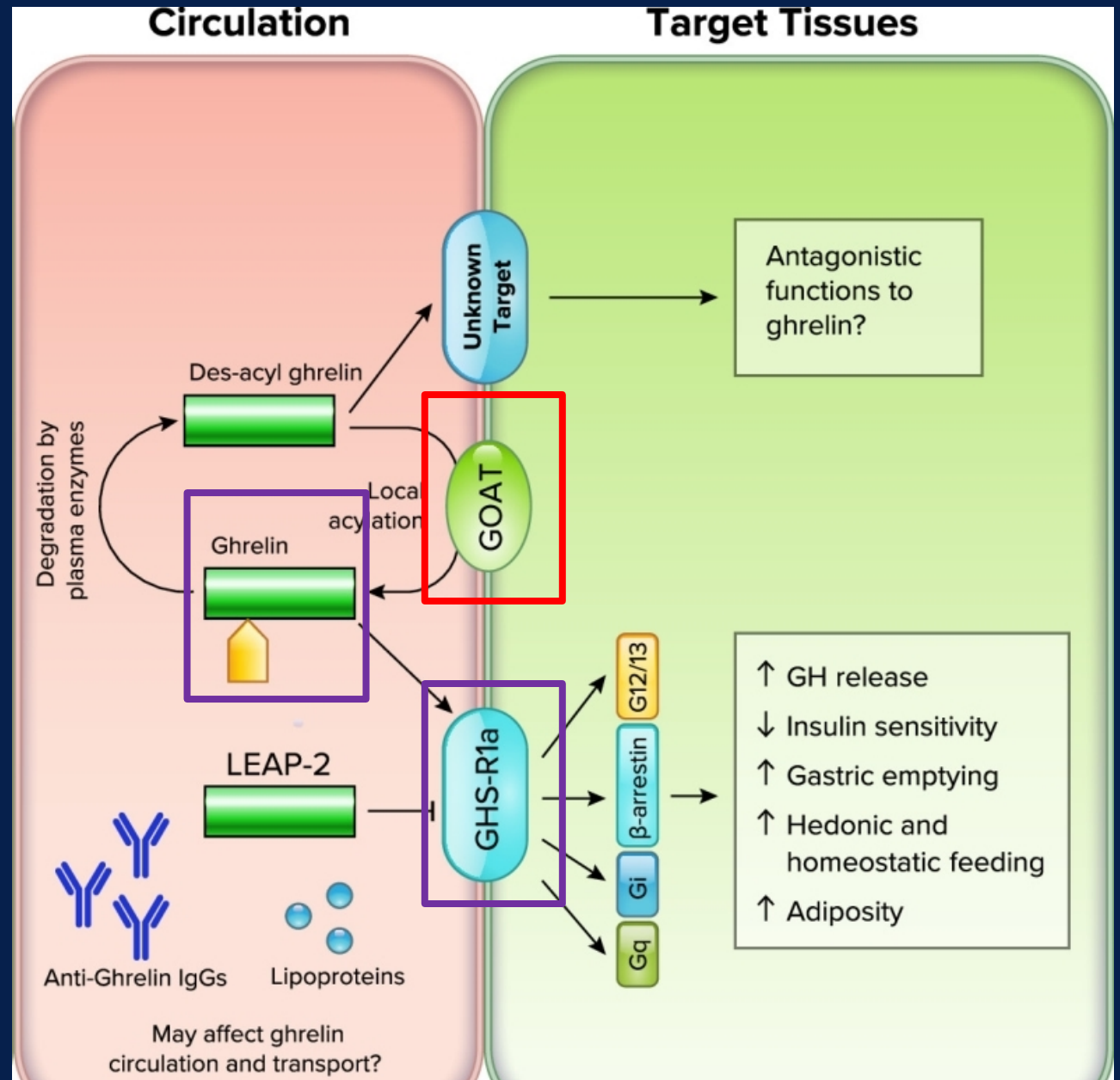


Ghrelin Functions



Howick, K., Griffin, B.T., Cryan, J.F. and Schellekens, H., 2017. From belly to brain: targeting the ghrelin receptor in appetite and food intake regulation. *International journal of molecular sciences*, 18(2), p.273.

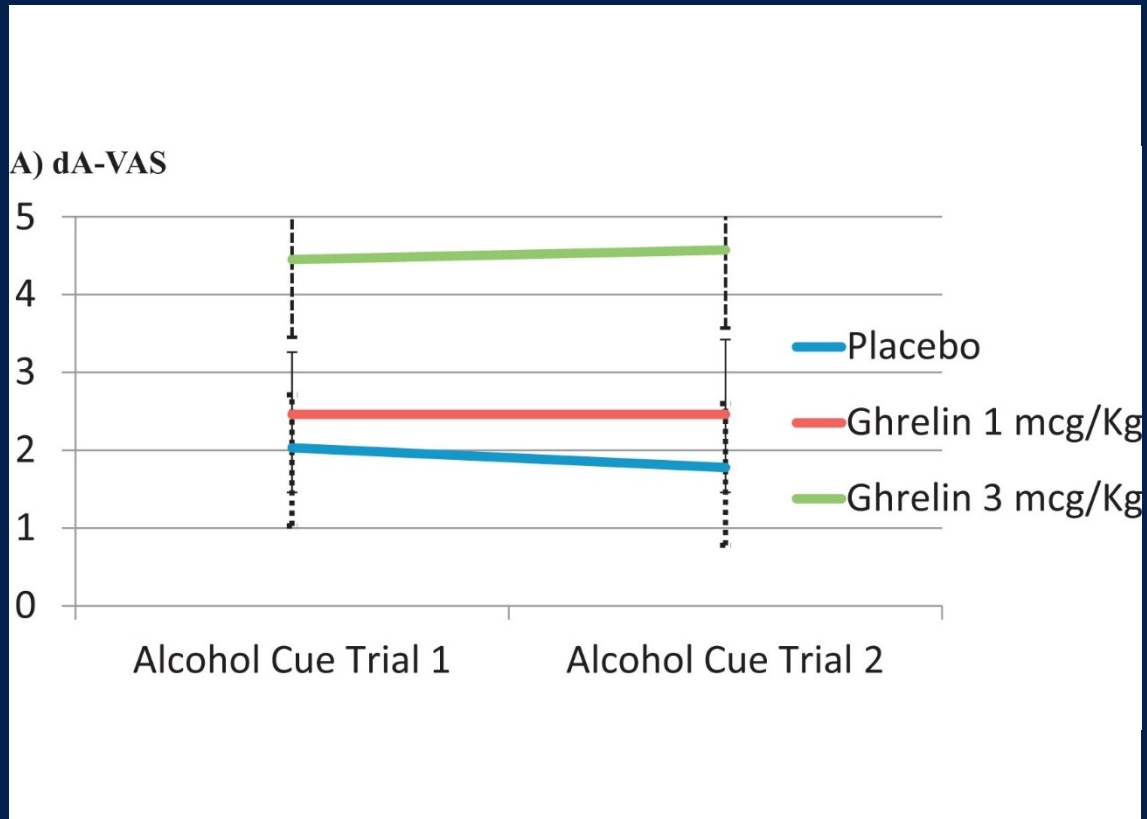
Overview of Ghrelin Physiology



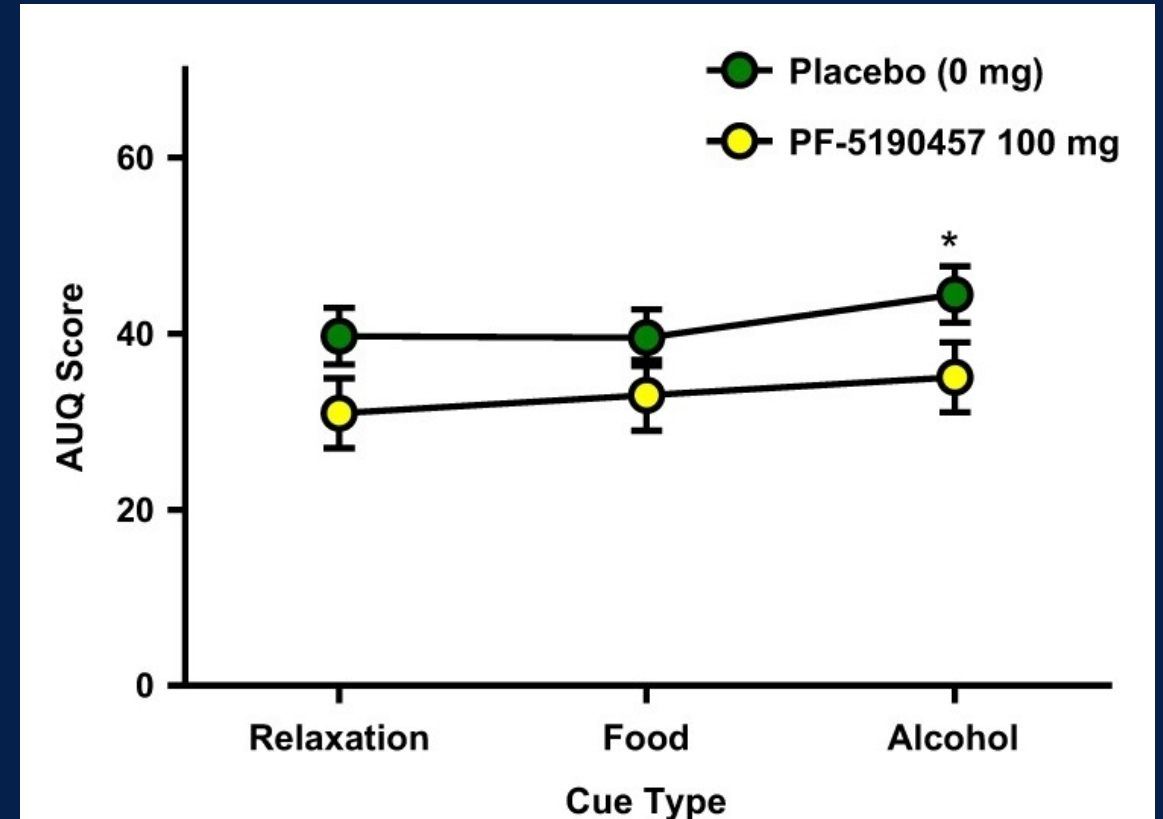
Deschaine, S.L. and Leggio, L., 2022. From “hunger hormone” to “it’s complicated”: ghrelin beyond feeding control. *Physiology*, 37(1), pp.5-15.

Ghrelin as a Target for AUD

Ghrelin Effect on Alcohol Urge



Alcohol Craving in Cue Reactivity



Leggio, L., et al., 2014. Intravenous ghrelin administration increases alcohol craving in alcohol-dependent heavy drinkers: a preliminary investigation. *Biological psychiatry*, 76(9), pp.734-741.; Lee, M.R., et al., 2020. The novel ghrelin receptor inverse agonist PF-5190457 administered with alcohol: preclinical safety experiments and a phase 1b human laboratory study. *Molecular psychiatry*, 25(2), pp.461-475..

Manipulating Ghrelin Signaling Via GOAT Inhibition in Alcohol Use Disorder

◆ Objectives:

- ◆ Evaluate the safety of the IND GLWL-01 in people with AUD
- ◆ Provide preliminary evidence of its efficacy to decrease drinking
- ◆ Evaluate its effect on food choices

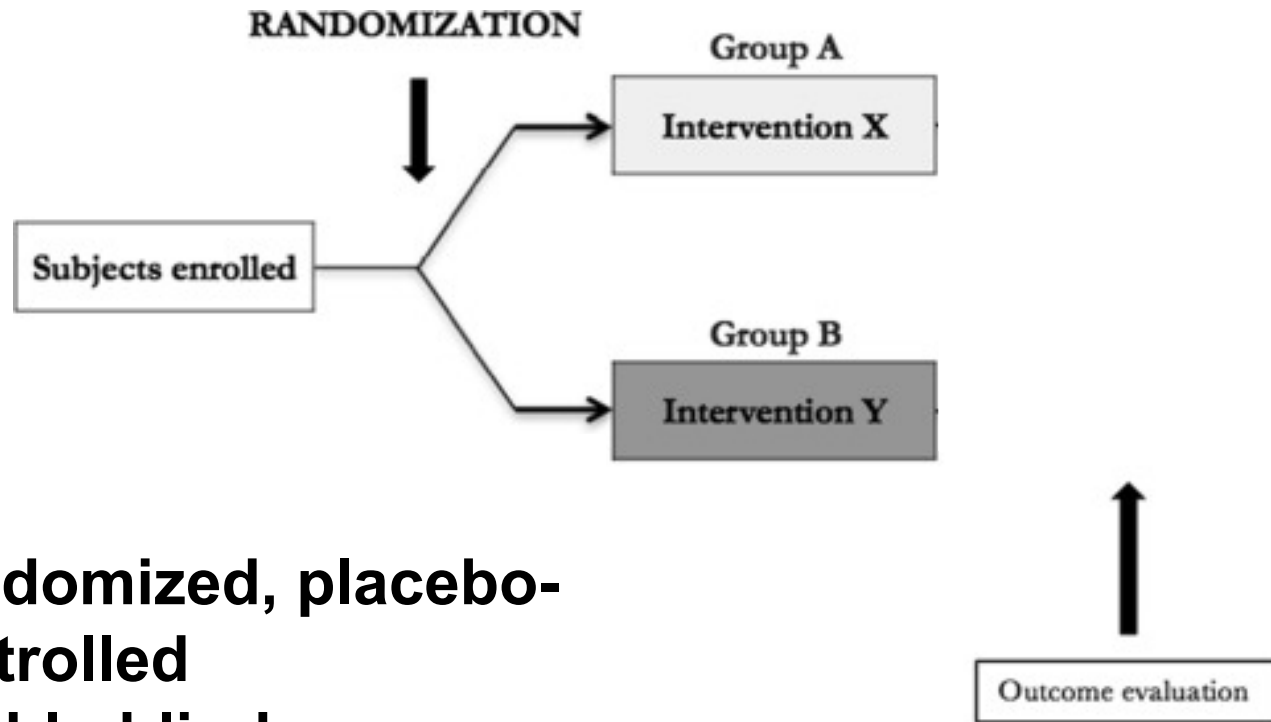
◆ Population:

- ◆ People ages 18-70 with AUD (34 completers)
- ◆ Not physically dependent on alcohol/benzodiazepines
- ◆ No other health conditions that would make participation unsafe

<https://clinicaltrials.gov/ct2/show/NCT03896516>



Study Design



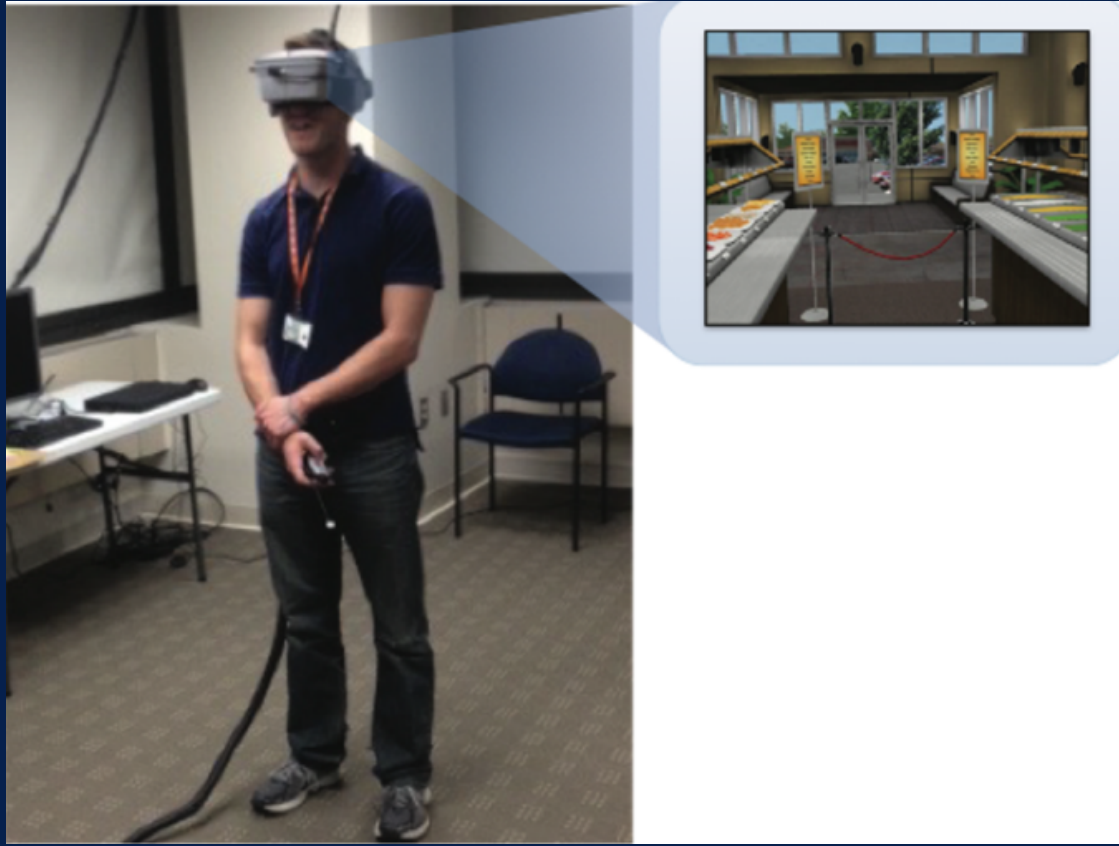
- **Randomized, placebo-controlled**
- **Double-blind**
- **Within-subject (crossover)**
- **Counterbalanced**

GOAT Study Schedule

DAY 1
Enrollment

Study Procedures

Virtual Reality Buffett



Cue Reactivity in the Mock Bar



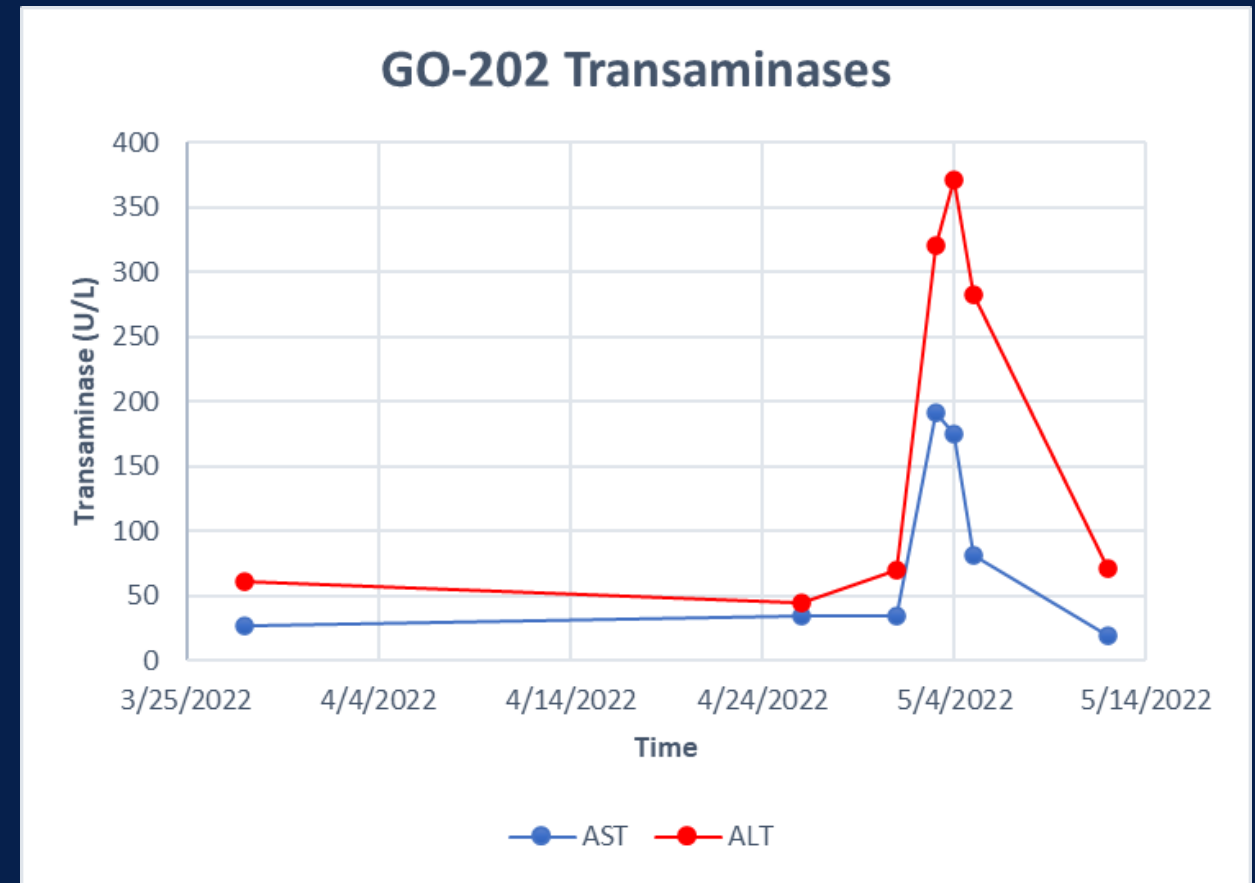
Known Unknowns

- ◆ “Surprising” side effects?
 - ◆ Green poop!
 - ◆ Decreased tobacco smoking in some patients



Known Unknowns

- ◆ **Hepatic Toxicity?**
 - ◆ **Stricter enrollment criteria (LFTs <1.5x ULN)**
 - ◆ **Liver fibrosis risk calculation (FIB-4)**



Unknown Unknowns

- ◆ Drug withdrawal
- ◆ Caffeine
- ◆ Cannabis/THC



Supplement Facts

Serving Size 8.0 fl oz (240 mL)

Servings Per Container: 2

Amount Per Serving	% Daily Value	
Calories	100	
Total Carb	27g	9%*
Sugars	27g	†
Riboflavin Vit B2	1.7mg	100%
Niacin Vit B3	20mg	100%
Vitamin B6	2mg	100%
Vitamin B12	6mcg	100%
Sodium	180mg	8%
Taurine	1000mg	†
Panax Ginseng	200mg	†
Energy Blend	2500mg	†
L-Carnitine, Glucose, Caffeine, Guarana, Inositol, Glucuronolactone, Maltodextrin		

*Percent Daily Values are based on a 2000 calorie diet. † Daily Value not established.

<https://shop.mypricechopper.com/product/monster-energy-drink-ultra-paradise-16-oz-00070847033080>



Unknown Unknowns

- ◆ Clinically normal protocol-based patient withdrawal
 - ◆ Threshold QTcF: <470 ms
 - ◆ Baseline QTcF: 377 ms
 - ◆ Post-Dosing QTcF: 439 ms

- **3a. Criteria for individual subject withdrawal:**
 -If there is prolongation of the QTcF interval to >500 ms or >60 ms over that on Day 1 ECG.....

Unknown Unknowns

- ◆ **Other logistics**
- ◆ **Food issues**
- ◆ **Managing patient expectations**



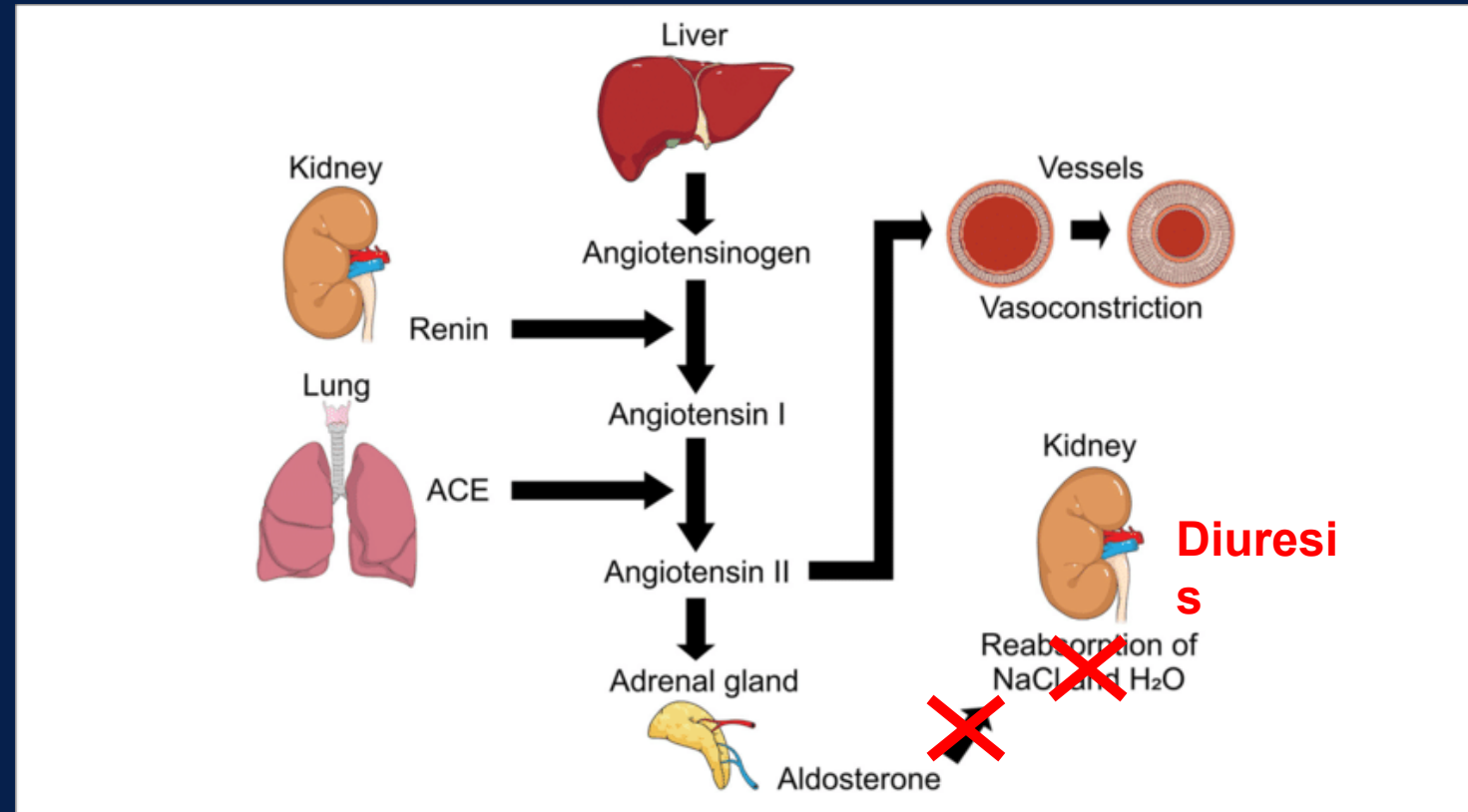
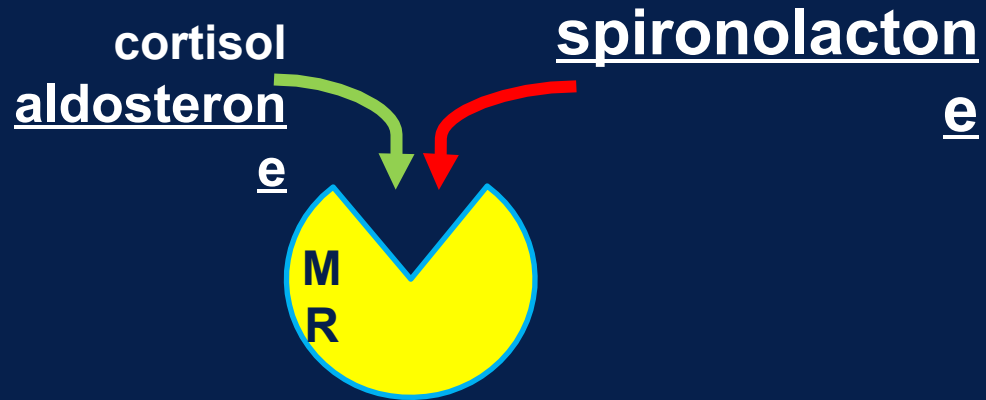
<https://easybudgetrecipes.com/ground-turkey-chili/>

Repurposing of Spironolactone

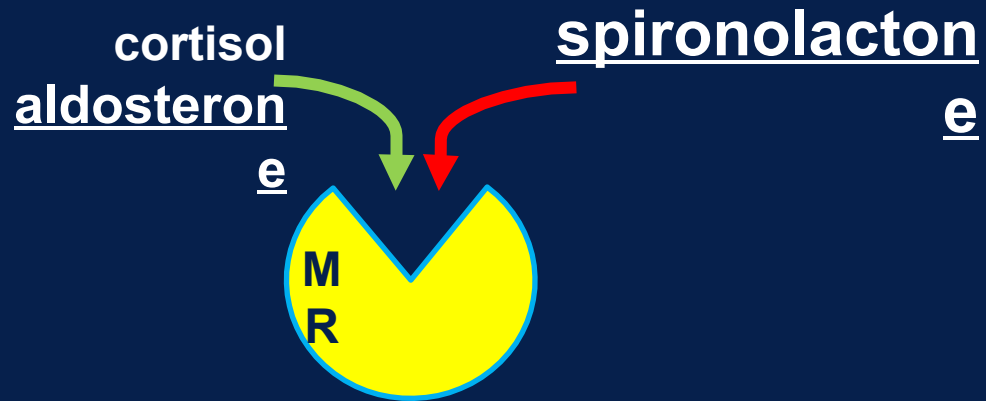
A new indication for a “good old drug”?



What is spironolactone?



What is spironolactone?



FDA approved in 1960

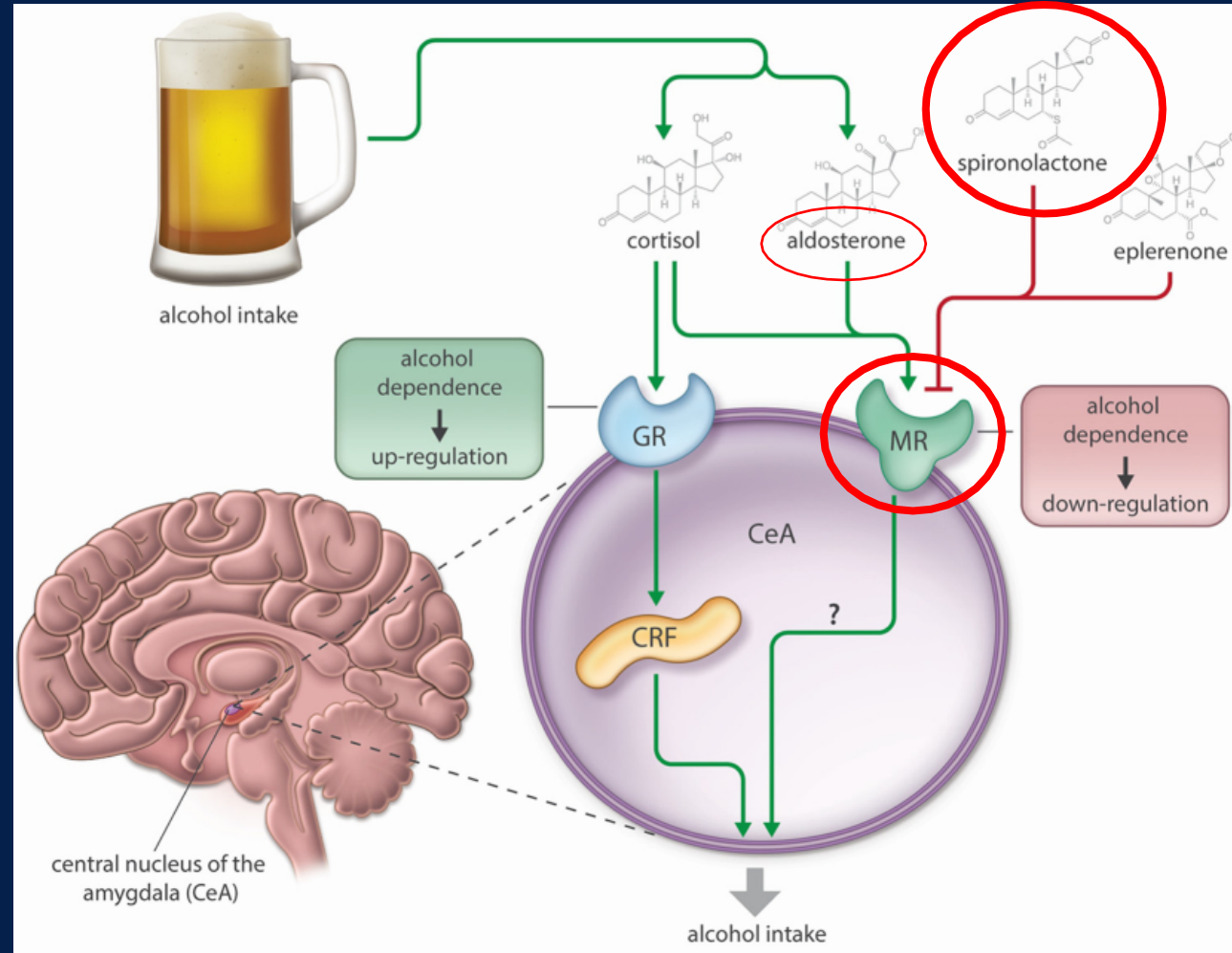
Hypertension: 25-100 mg/day

Heart failure: 25-50 mg/day

**Edema due to cirrhosis: 25-200
mg/day**

Hyperaldosteronism: 100-400 mg/day

Why Spironolactone and the Mineralocorticoid Receptor (MR)?

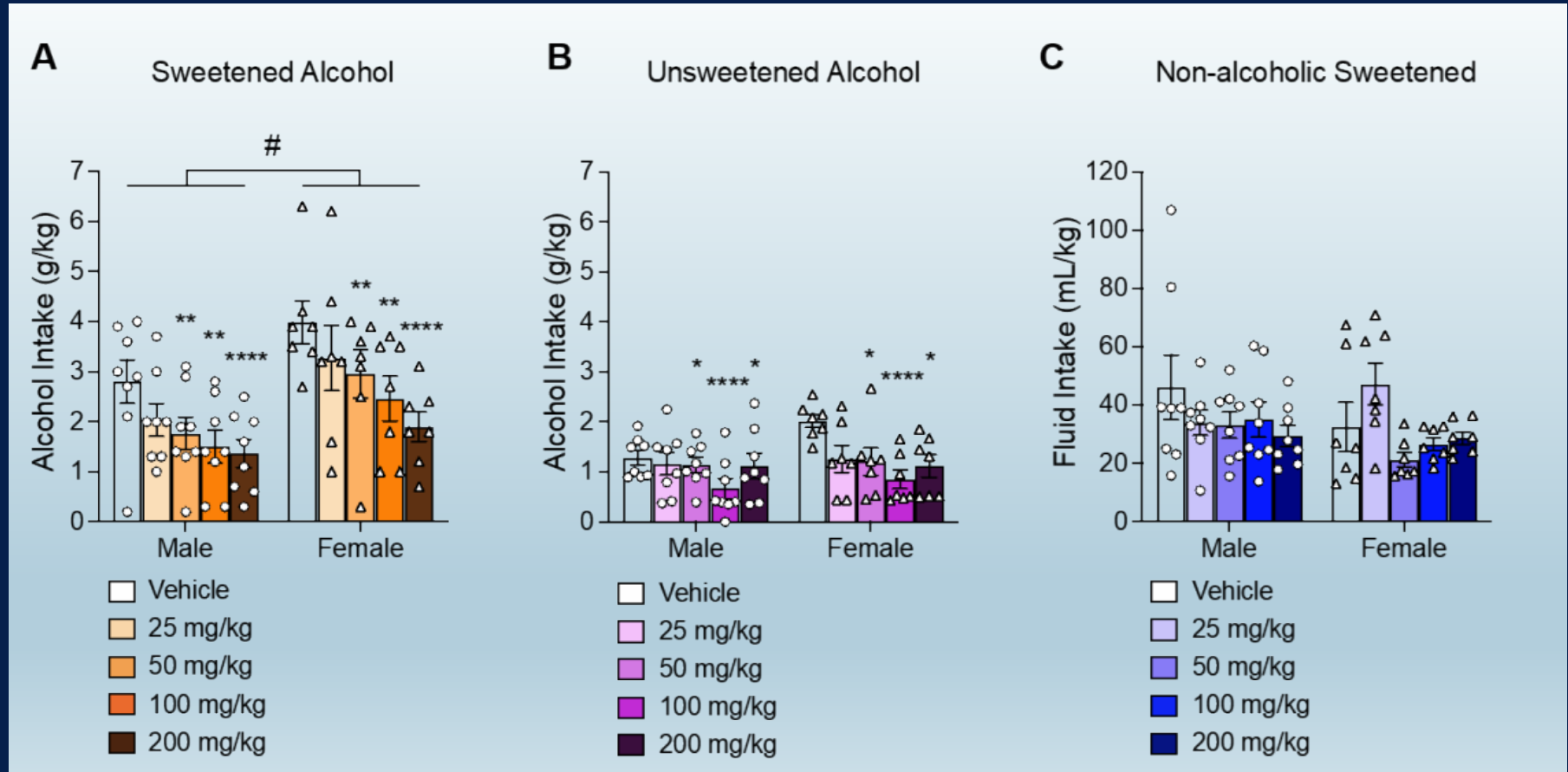
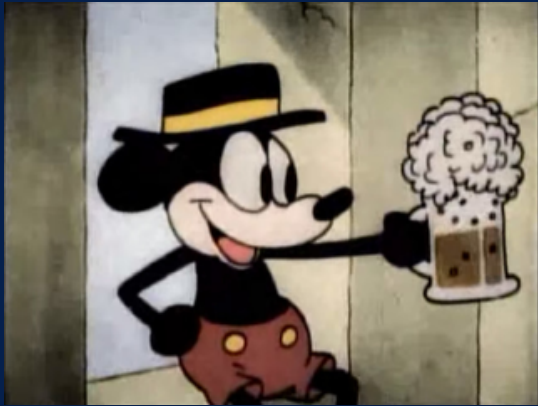


Lékó AH et al., ACS Chem Neurosci, 2022

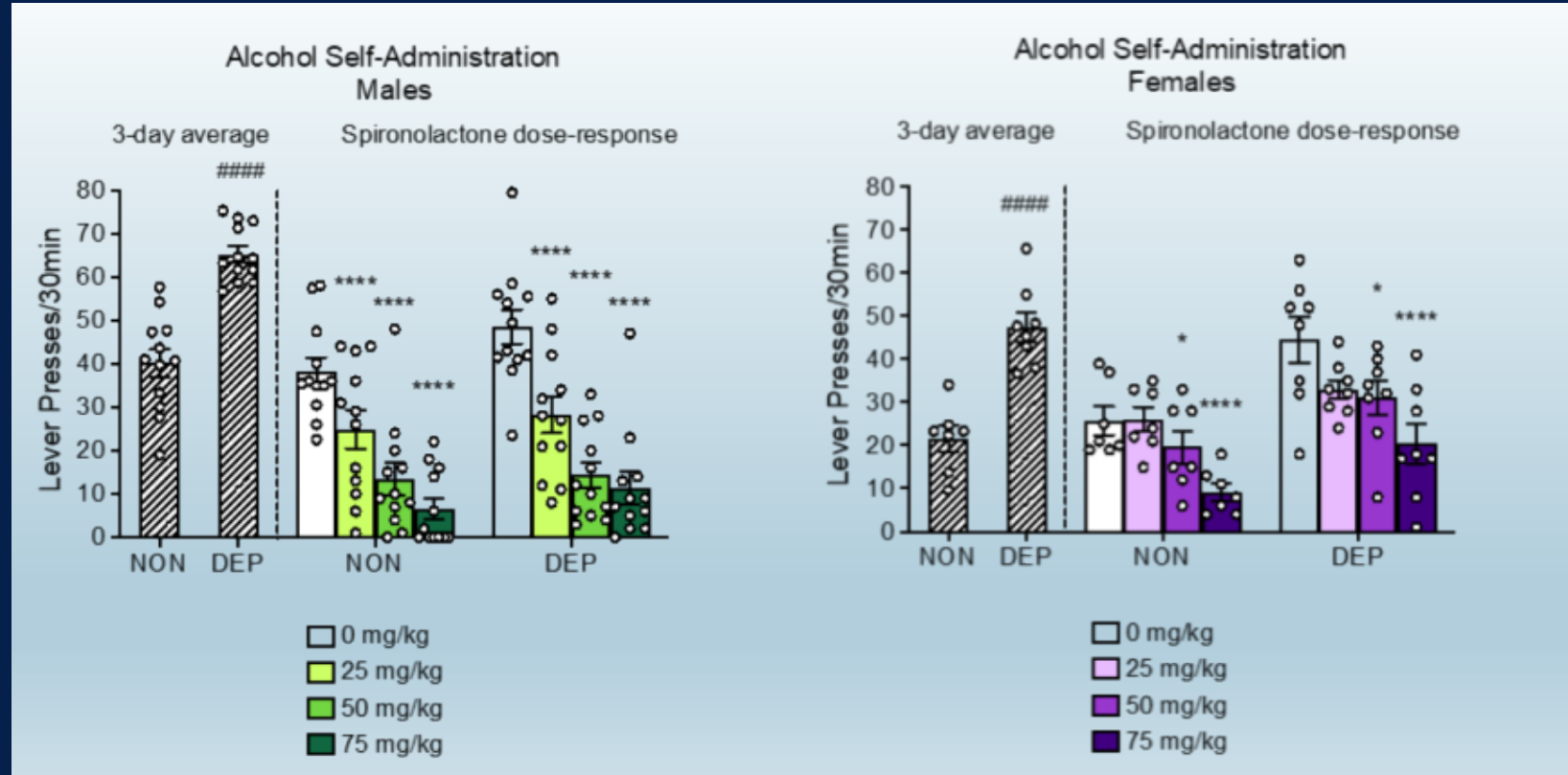
Leggio L et al., Alcohol, 2008

Aoun et al., Mol Psychiatry, 2018

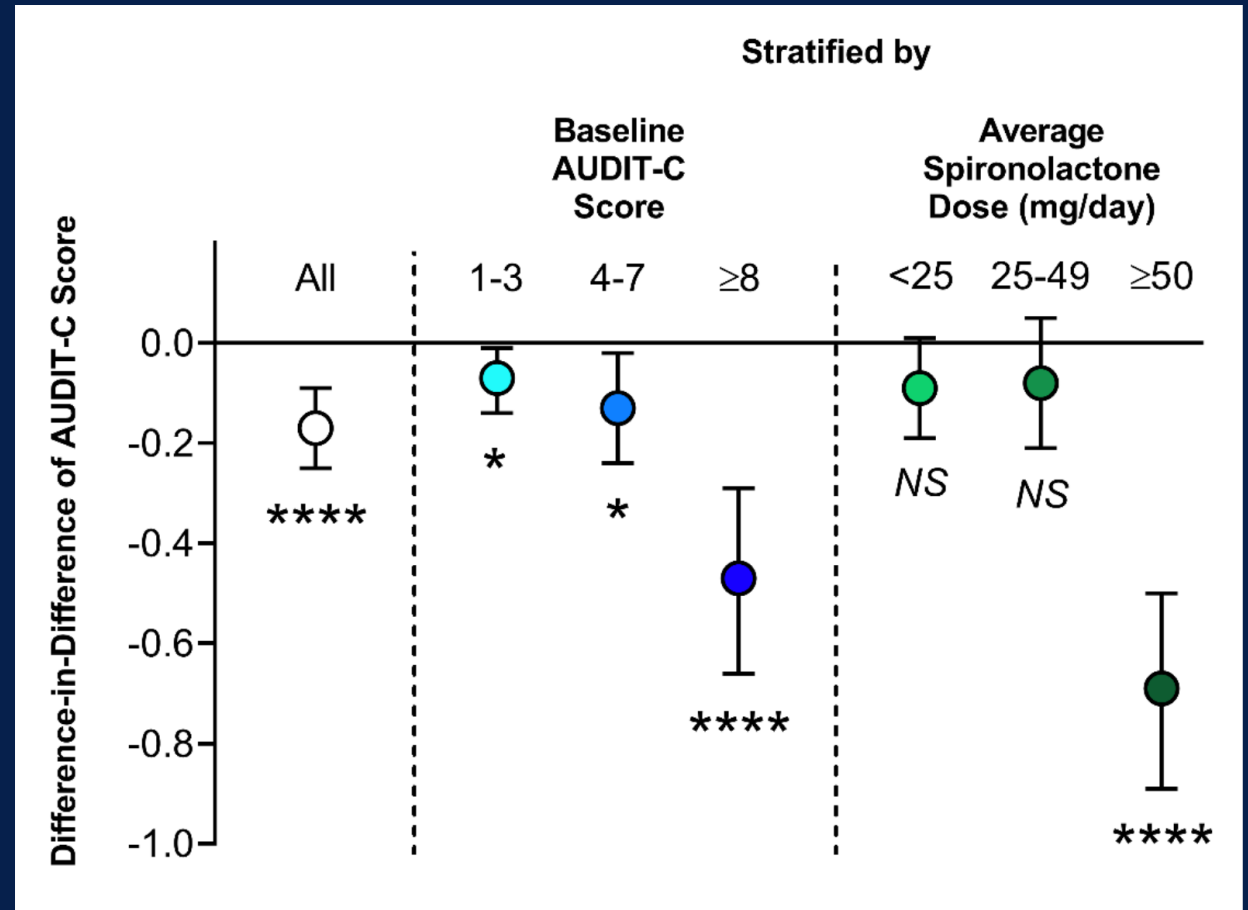
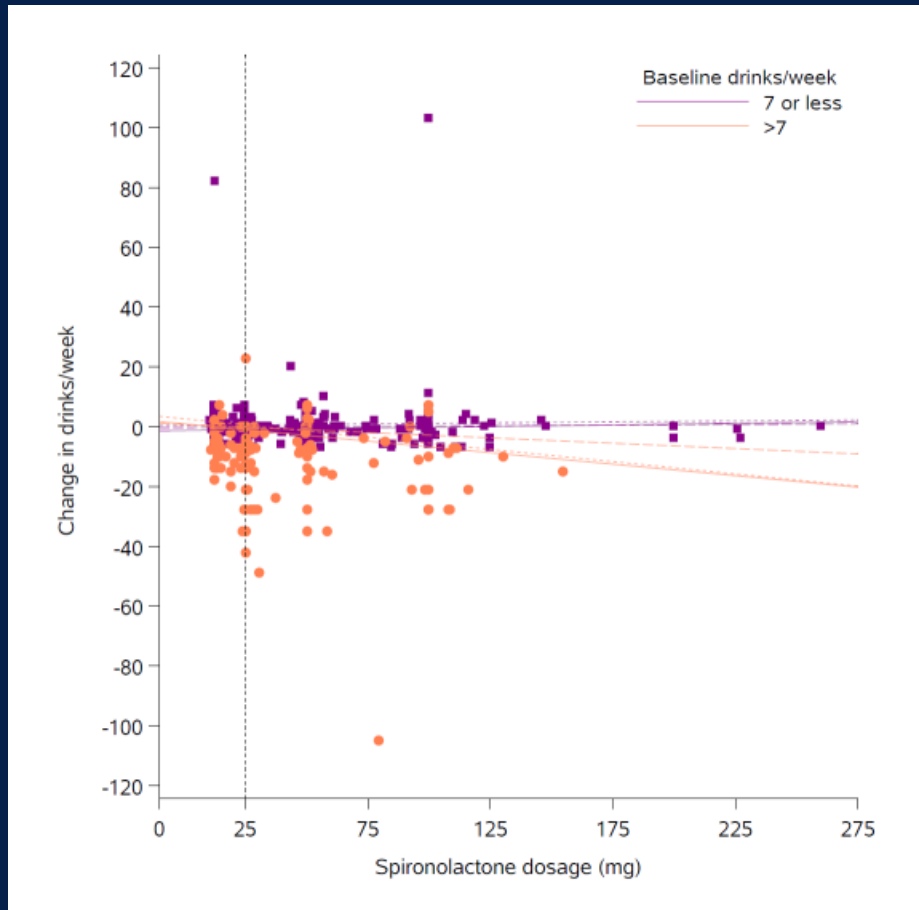
Spironolactone affects alcohol intake



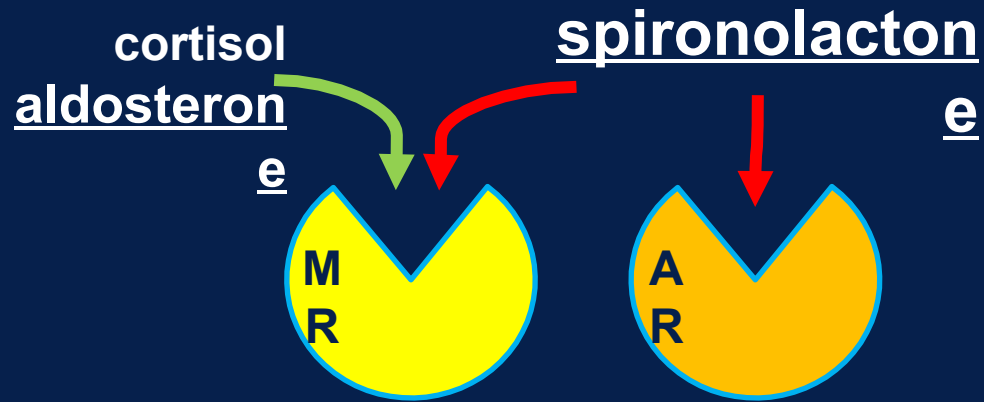
Spironolactone affects alcohol intake



OK, but what about humans?



Spironolactone – side effects



Hyperkalemia

Hypotension

Gynecomastia

Worsening renal function

Hyponatremia

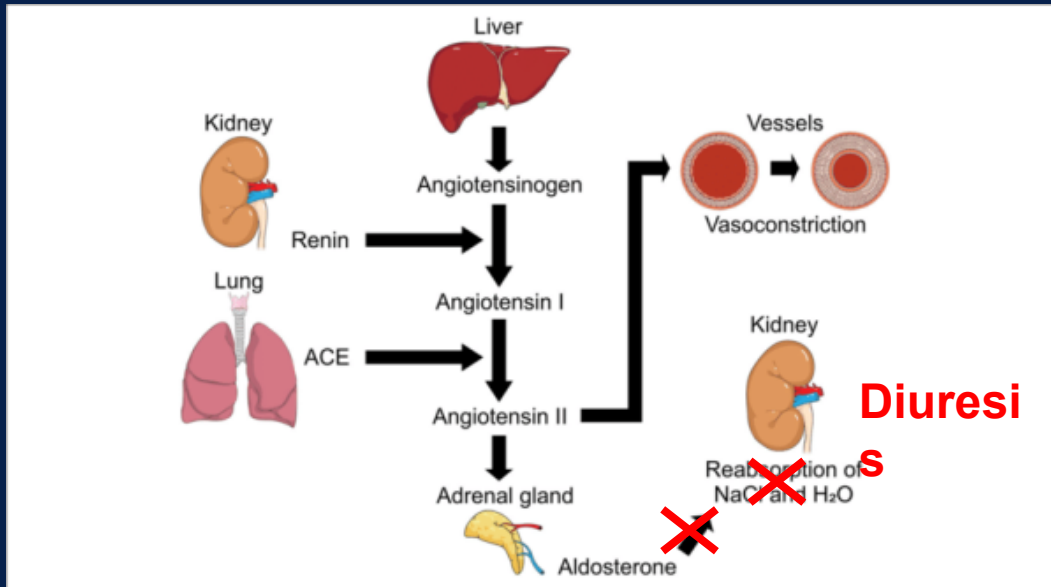
Hypomagnesemia

Hypocalcemia

Hypochloremic alkalosis

Hyperglycemia

Hyperuricemia
(asymptomatic)



Spiroonolactone in Alcohol Use Disorder (SAUD)

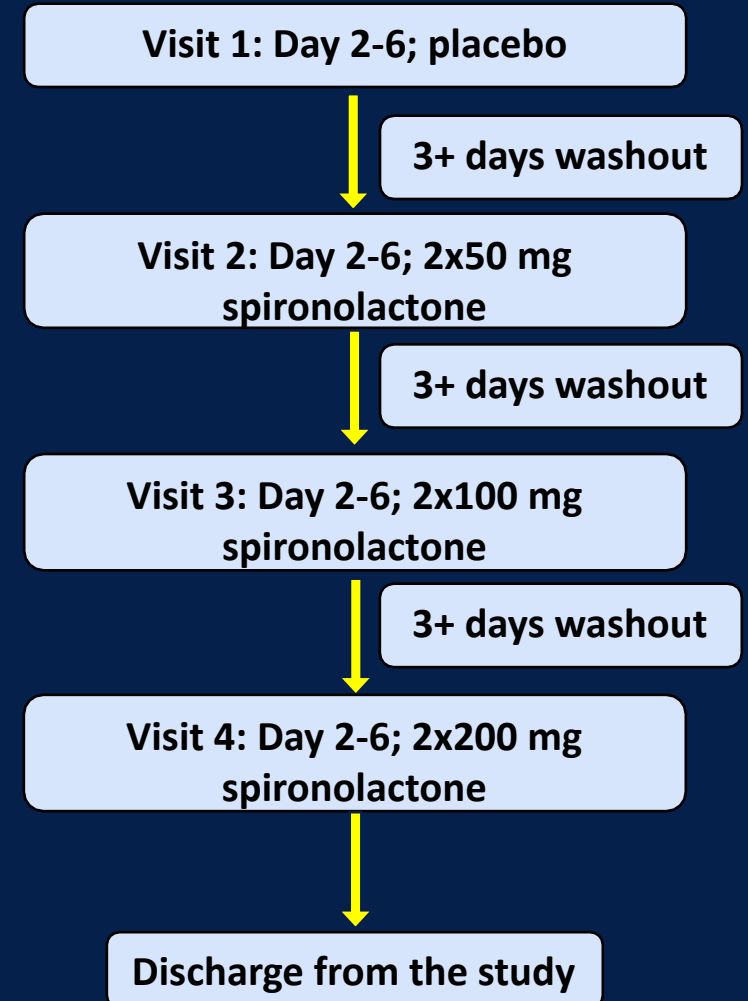
Phase 1b study under review

◆ Features

- ◆ Inpatient, within-subject, double-blind, placebo-controlled
- ◆ Heavy-drinking individuals with AUD (N=12)
- ◆ Day 6: Spiroonolactone + Alcohol

◆ Outcomes:

- ◆ Spiroonolactone Pharmacokinetics
- ◆ Safety/tolerability
- ◆ Alcohol Pharmacokinetics
- ◆ Subjective and cognitive effects of alcohol



“Alcohol Challenge”

Day 6: morning dose
(spironolactone/placebo)

90 min

Oral alcohol 0.5 g / kg fat-free
mass (based on DXA scan)

BAC 0.08%

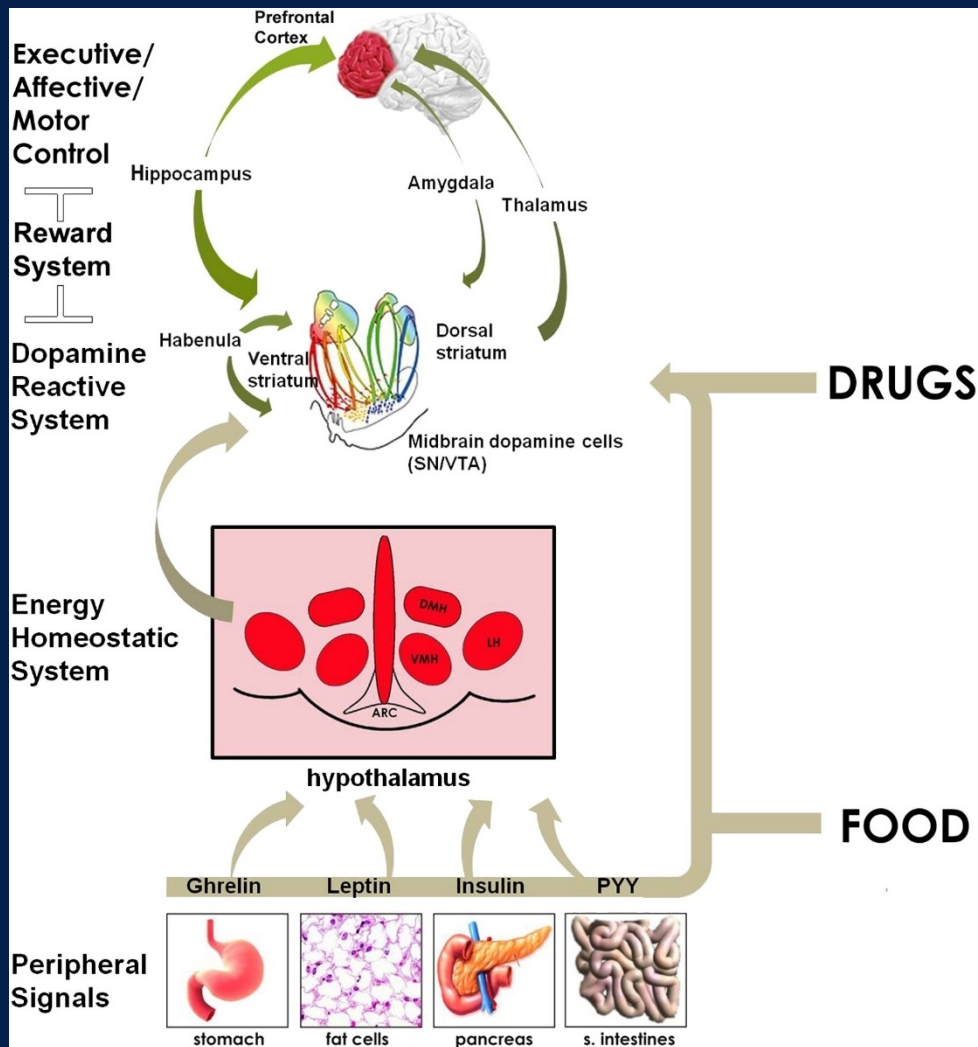
Spironolactone PK
Alcohol PK (BAC)
Subjective and cognitive effects
of alcohol



GLP-1 Analogues for AUD: Bench-to-Bedside and Back



Feeding/Metabolic Neuropeptides



GLP-1

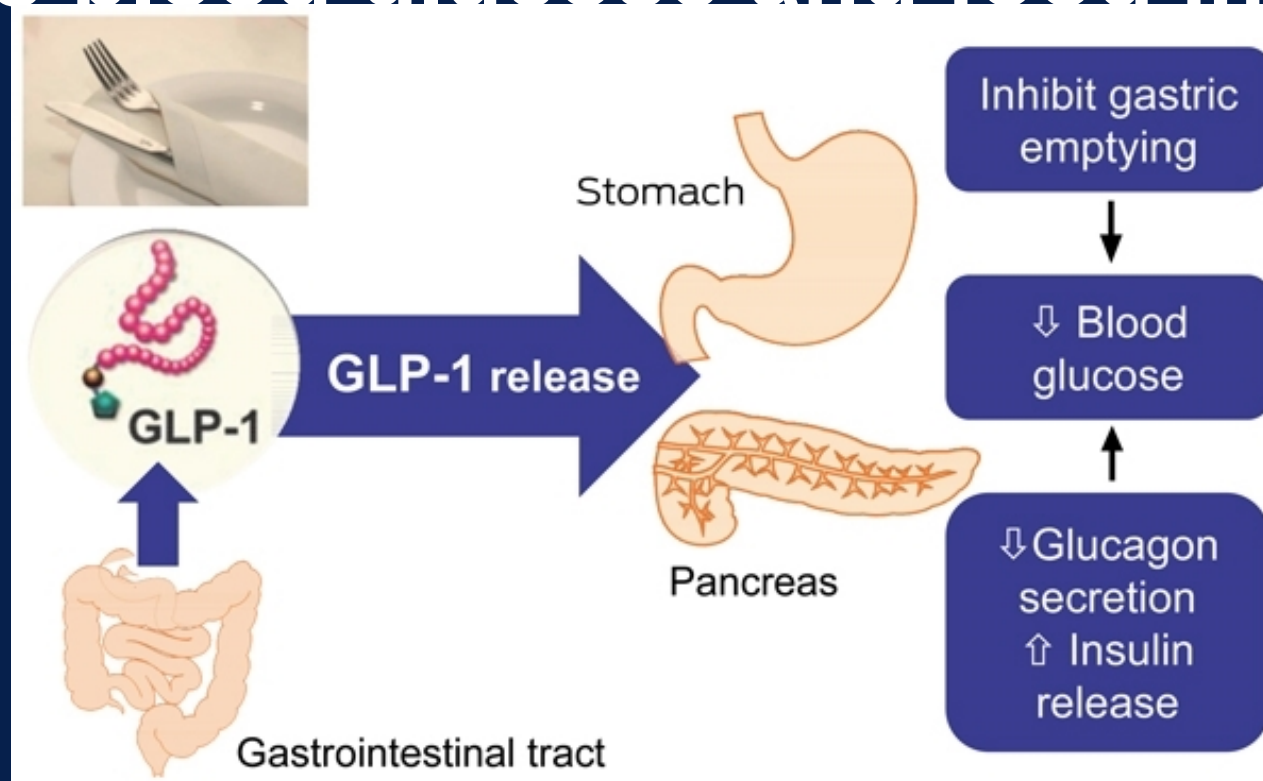


Volkow ND, et. al. The addictive dimensionality of obesity. Biol Psychiatry 2013



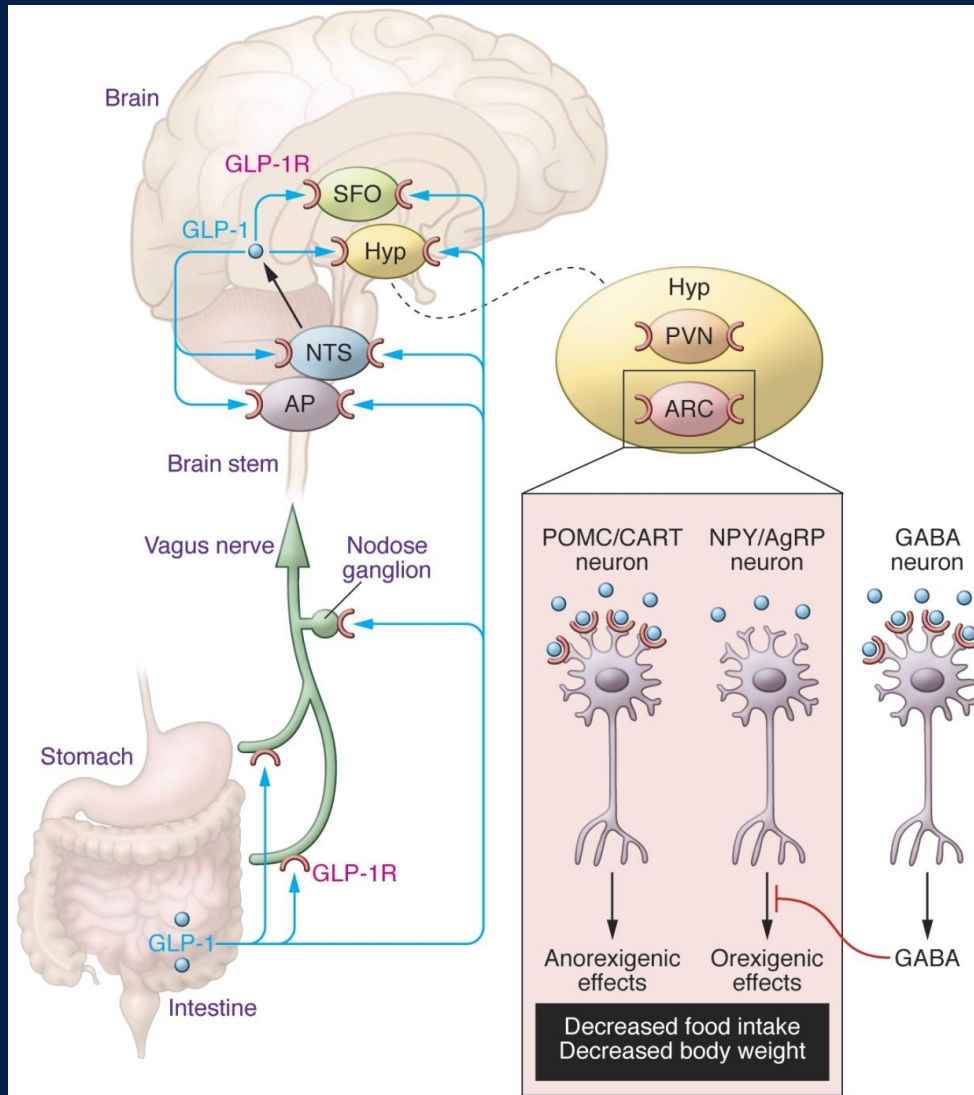
Glucagon-Like Peptide-1 (GLP-1)

- ◆ A peptide with 30 amino acids
- ◆ Produced in the intestinal mucosa and pancreas
- ◆ Regulates blood glucose and food intake

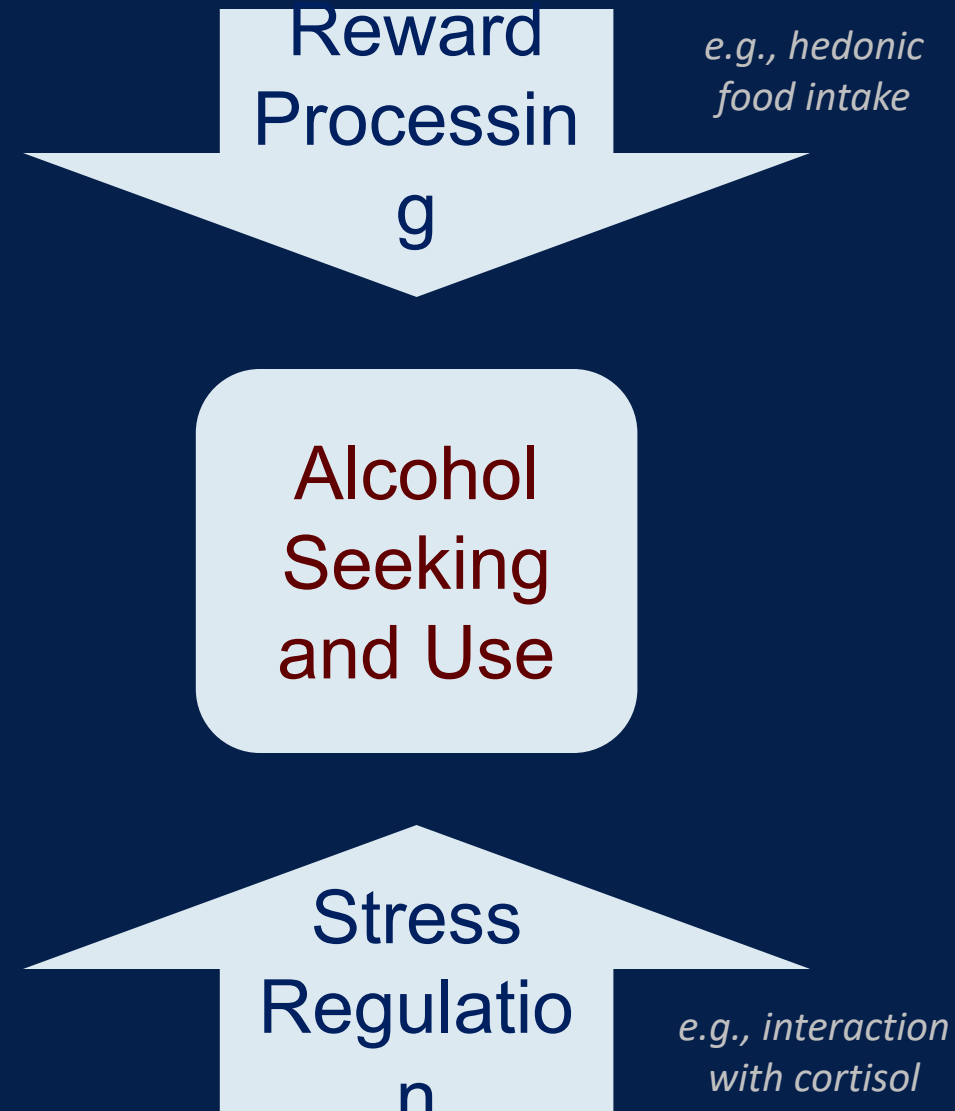


Cohen ND, et. al. The rationale for combining GLP-1 receptor agonists with basal insulin. Med J Aust 2013

GLP-1 is also a Neuropeptide



Baggio LL, et. al. Glucagon-like peptide-1 receptors in the brain: controlling food intake and body weight. J Clin Invest 2014



GLP-1 and Alcohol

- ◆ **GLP-1 receptor gene variants impact AUD risk and brain activity/connectivity**
- ◆ **Acute alcohol intake reduces circulating GLP-1 levels**
- ◆ **Chronic alcohol intake increases GLP-1 receptor gene expression in AUD-related brain regions**

► *Transl Psychiatry*. 2015 Jun 16;5(6):e583. doi: 10.1038/tp.2015.68.

The glucagon-like peptide-1 receptor as a potential treatment target in alcohol use disorder: evidence from human genetic association studies and a mouse model of alcohol dependence

P Suchankova ¹, J Yan ², M L Schwandt ³, B L Stangl ⁴, E C Caparelli ⁵, R Momenan ⁶, E Jerlhag ⁷, J A Engel ⁷, C A Hodgkinson ⁸, M Egli ⁹, M F Lopez ¹⁰, H C Becker ¹¹, D Goldman ⁸, M Heilig ³, V A Ramchandani ⁴, L Leggio ¹²

► *Sci Rep*. 2022 Jul 29;12(1):13027. doi: 10.1038/s41598-022-17190-3.

Differential association between the GLP1R gene variants and brain functional connectivity according to the severity of alcohol use

Mehdi Farokhnia ^{# 1 2 3}, Samantha J Fede ^{# 4}, Erica N Grodin ⁵, Brittney D Browning ⁶, Madeline E Crozier ⁶, Melanie L Schwandt ⁷, Colin A Hodgkinson ⁸, Reza Momenan ⁴, Lorenzo Leggio ^{9 10 11 12 13 14}

► *Addict Biol*. 2022 Sep;27(5):e13211. doi: 10.1111/adb.13211.

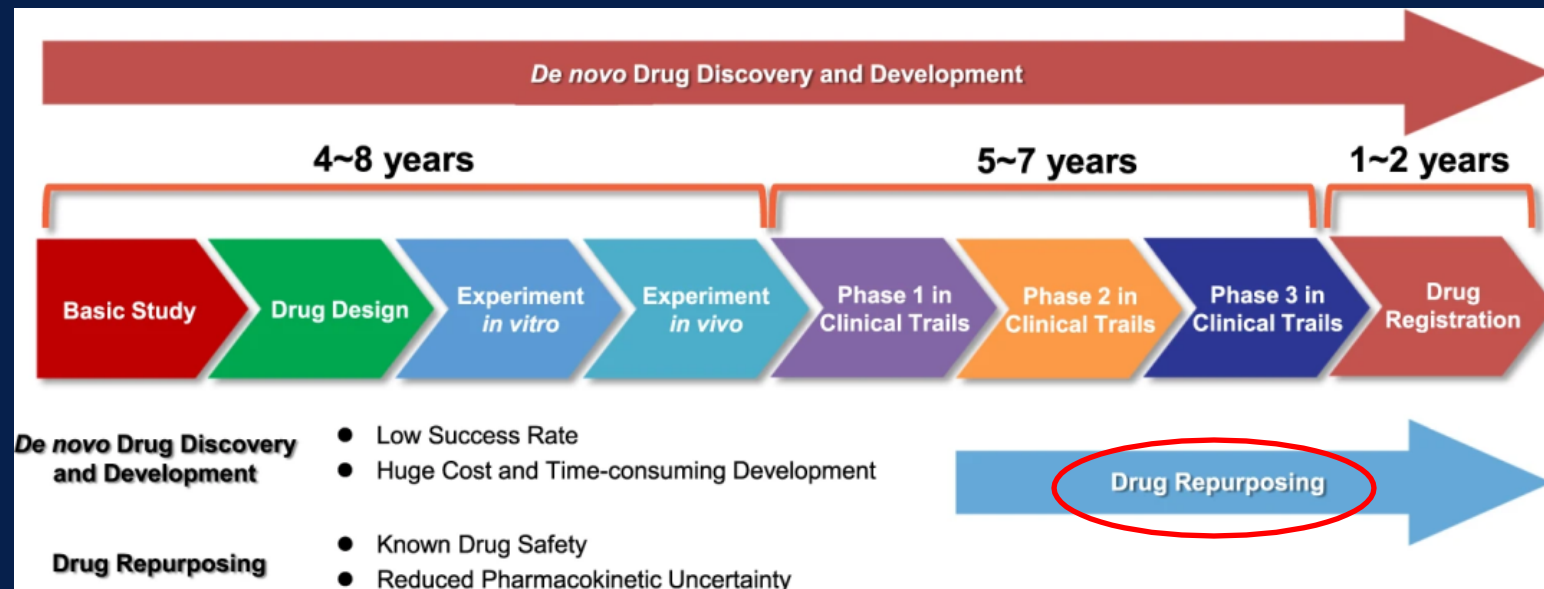
The glucagon-like peptide-1 system is modulated by acute and chronic alcohol exposure: Findings from human laboratory experiments and a post-mortem brain study

Mehdi Farokhnia ^{1 2 3}, Brittney D Browning ¹, Madeline E Crozier ¹, Hui Sun ⁴, Fatemeh Akhlaghi ⁵, Lorenzo Leggio ^{1 2 6 7 8 9}

Alcohol intake/dependence modulates the GLP-1 system

GLP-1 system as a pharmacotherapeutic target for AUD?

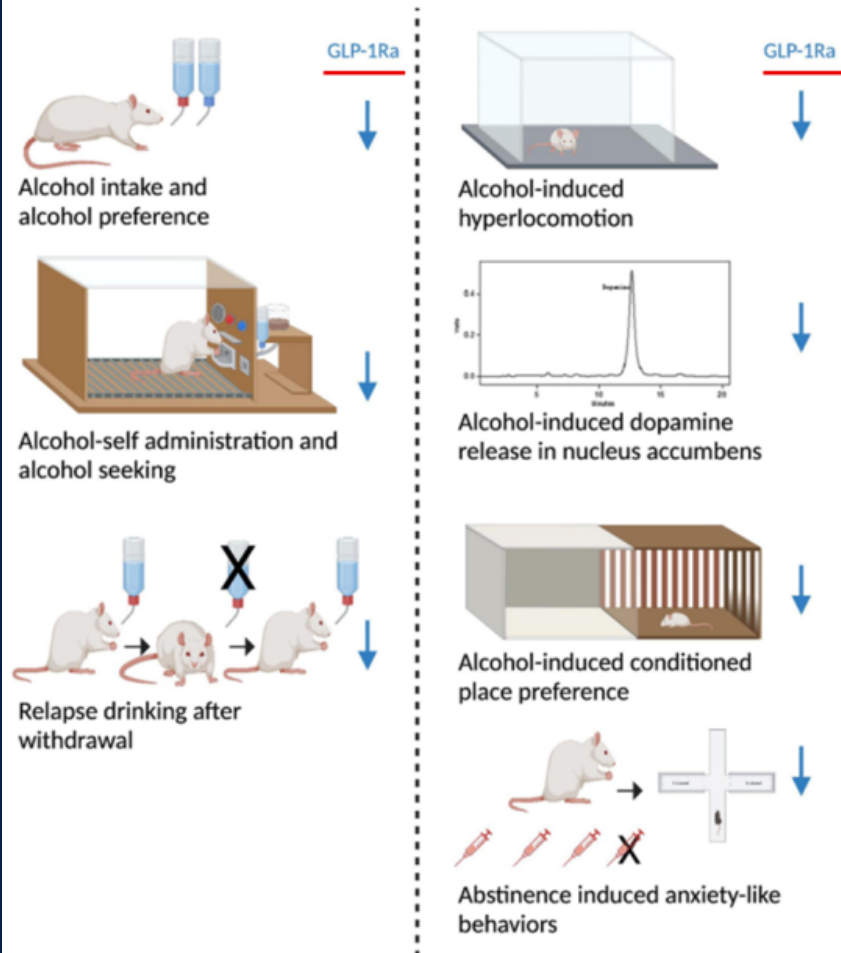
- ◆ FDA-approved GLP-1 analogues:
Exenatide, Lixisenatide, Albiglutide, Dulaglutide, Liraglutide, Semaglutide



Zhang Z, et. al. Overcoming cancer therapeutic bottleneck by drug repurposing. Signal Transduct Target Ther 2020

GLP-1 Analogues and Alcohol Use

Alcohol-related outcomes

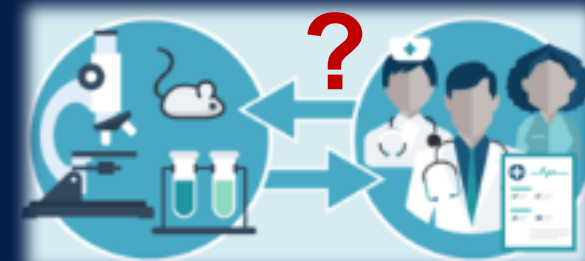


Tufvesson-Alm M, et. al. Insight into the role of the gut-brain axis in alcohol-related responses: Emphasis on GLP-1, amylin, and ghrelin. *Front Psychiatry* 2023

Summary of the effects of different GLP-1 receptor agonists on alcohol-mediated behaviours.

GLP-1 receptor agonist	Impact on consumption-related behaviours	Reference
Ex 4/exenatide	Prevents alcohol seeking in the progressive ratio test in the operant self-administration model in rats Decreases alcohol intake in the intermittent access model in rats	Egecioglu et al, 2013c
	Reduces the initial five days of alcohol intake in vervet monkeys Reduces the intravenous self-administration of alcohol in mice Declines the number of drinking bouts and protracts the latency to the first drink as well as prevents relapse drinking in socially housed mice	Egecioglu et al, 2013c Shirazi et al 2013 Thomsen et al, 2019 Sorensen et al, 2016 Thomsen et al, 2017
	Reduces alcohol intake in peripheral, but not central, GLP-1 receptor knockout mice	Sirohi et al, 2016
	Infusion into the NAc shell decreases alcohol intake in rats consuming high, but not low, amounts of alcohol	Vallof et al, 2019a
	Ex4 into the NAc shell, but not core, reduces alcohol intake in female rats	Abtahi et al, 2018
	Infusion into the LDTg decreases alcohol intake in rats consuming high, but not low, amounts of alcohol	Vallof et al, 2019a
	Infusion into the anterior VTA does not modulate alcohol intake in rats	Vallof et al, 2019a
	Infusion into the NTS decreases alcohol intake in alcohol-consuming rats	Vallof et al, 2019b

Jerlhag E. Alcohol-mediated behaviours and the gut-brain axis; with focus on glucagon-like peptide-1. *Brain Res* 2020



Novel GLP-1 Analogues

- ◆ Longer half-life → less frequent injections
- ◆ Higher affinity for the GLP-1R
- ◆ More effective in ↓ HbA1c and weight

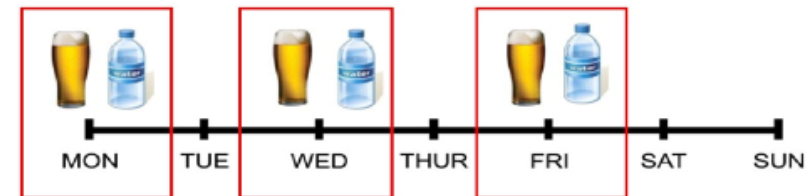
The Discovery and Development of Liraglutide and Semaglutide

Lotte Bjerre Knudsen^{1*} and Jesper Lau²

Front Endocrinol 2019

Long-Acting Glucagon-Like Peptide-1 Receptor Agonists Suppress Voluntary Alcohol Intake in Male Wistar Rats

Vincent N. Marty^{1†}, Mehdi Farokhnia^{2,3,4†}, Joseph J. Munier¹, Yatendra Mulpuri¹, Lorenzo Leggio^{2,3,5,6,7,8} and Igor Spigelman^{1*}

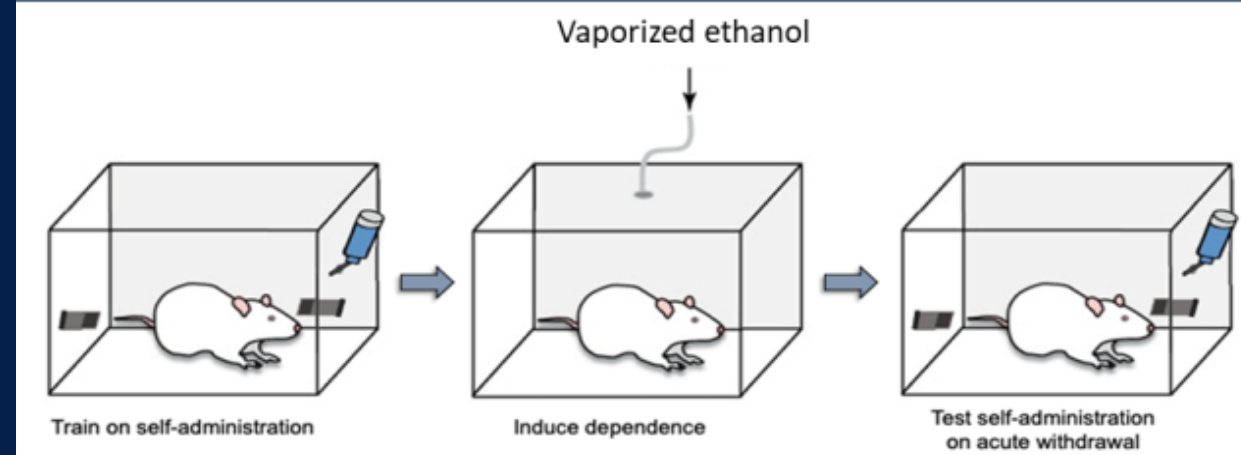
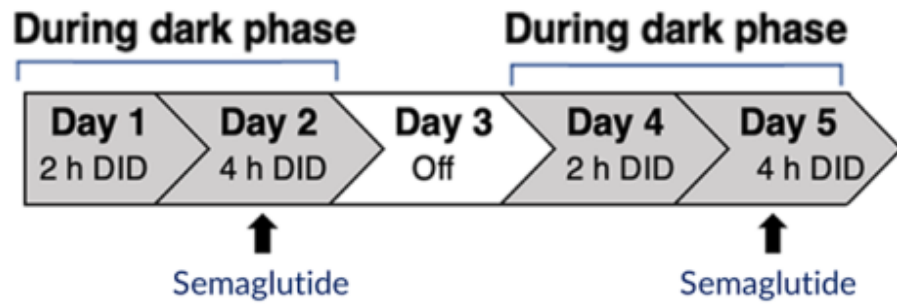


Front Neurosci 2020

- ◆ Semaglutide and liraglutide reduced alcohol intake
- ◆ Semaglutide's effects were more specific for alcohol

Semaglutide for Alcohol Use in Rodents

(unpublished)



Semaglutide dose-dependently reduced binge-like alcohol drinking in male and female mice

Semaglutide dose-dependently reduced alcohol self-administration in male and female rats

Semaglutide for Clinical Use



Ozempic

FDA-approved for diabetes



Wegovy

FDA-approved for obesity

Semaglutide for Alcohol Use in Humans

(RCT under development)

- ◆ **Outpatient, between-subject, double-blind, placebo-controlled**
- ◆ **Heavy-drinking individuals with AUD (N = 52 completers)**
- ◆ **Weekly injection of semaglutide/placebo up to 2.4 mg/week**
- ◆ **Outcomes:**
 - ◆ **Safety and tolerability**
 - ◆ **Alcohol use (drinks/week, etc.)**
 - ◆ **Three experimental procedures**

Visit / Week # →	1	2	3	4	5	6	7	8	9	10	11	12	13	14	15	16	17	18	19	20	Follow up	
Study Drug or Placebo (mg)	x	x	x	x	x	x	x	x	x	x	x	x	x	x	x	x	x	x	x	x	x	
Take Control	x			x			x			x			x			x			x			



Assessment S:

- ◆ Alcohol intake, side effects (co-primary outcome)
- ◆ Clinical assessments and labs
- ◆ Measures of craving, mood, anxiety, sleep, anhedonia, etc.



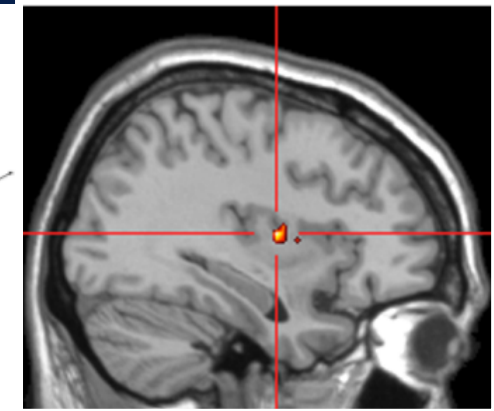
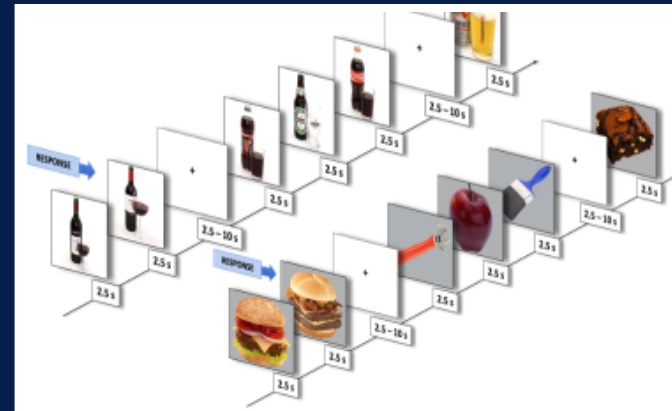
Virtual reality



Brain fMRI



Cue-reactivity



Take-Home Points

- ◆ **Current AUD medications are helpful but not sufficient**
- ◆ **Endocrine pathways (ghrelin, mineralocorticoids, GLP-1) are promising targets for AUD drug discovery**
- ◆ **Multiple drug discovery techniques are being pursued by the TAMB at the NIDA IRP**
 - ◆ **Testing of Investigational New Drugs (e.g., GLWL-01)**
 - ◆ **Repurposing of existing drugs (e.g., spironolactone, semaglutide)**