Ondansetron in alcohol use disorder: a systematic review

**Auteurs**

**Key words** *alcohol use disorder, ondansetron, 5-HT*3 *receptors, serotonin, craving, pharmacogenetics, personnalized medicine*

**Abstract**

**Context** Alcohol use disorder (AUD) is a frequent disorder. Few treatments have shown a great efficacy in clinical studies. It is hypothetized that this is due to the fact that AUD suffering people constitute a heterogeneous group composed of various endophenotypes. Ondansetron, a selective 5-HT3 receptor antagonist, has been evaluated in AUD and partciularly in certain subgroups of patients. This review presents clinical studies evaluating its impact.

**Methods** A systematic review was conducted on Pubmed and Wiley Online Library. All 19 clinical trials involving the use of ondansetron in AUD where included.

**Results** Ondansetron was found to be effective mostly in certain subgroups of AUD-suffering patients. The identified subgroups were based on clinical evaluation (age of onset, personality type) and genotype. A number of limitations remains and further studies are needed.

# Introduction

Alcohol use disorder (AUD) is a heterogeneous and chronic relapsing disorder resulting in a complex interaction between neurobiological, genetic and environmental factors. Despite the demonstrated efficacy of some approved medications (acamprosate, naltrexone, disulfiram), a key barrier is the fact that these medications are not effective in every patient pointing out the need for more personalized therapy approaches to overcome this heterogeneity. In this perspective, major advances in pharmacogenetics have highlighted distinct clinical subgroups of AUD according to genetic variation, that could be associated with differential treatment responses. Thus, the identification of patient subtypes that are most likely to respond favorably to different medications is crucial with a need for a better targeting of medication to specific patients.

Amongst the emerging pharmacotherapies for AUD, Ondansetron (IUPAC name: (RS)-9-Methyl-3- [(2-methyl-1H-imidazol1-yl)methyl]-2,3-dihydro-1Hcarbazol-4(9H)-one), a selective antagonist of the 5-HT3 receptor, has shown some promising results. Ondansetron is approved by ANSM in France and FDA in the USA as an antiemetic for cancer treatment-induced and anesthesia related nausea and vomiting. In the late 1980s, Haganetal et al. [1] showed that the injection of ondansetron in the ventral tegmental area of the rat brain lessened induced hyperactivation in the nucleus accumbens, pointing out the tight relationship between serotonin function and the mesolimbic dopaminergic reward system. Based on these findings and the role of dopaminergic activity on the rewarding effects of alcohol, ondansetron was thought to attenuate the pleasurable subjective effects of alcohol and thereby to reduce alcohol consumption in AUD suffering patients. Some phase 1 clinical studies ([2], [3]) found promising results in healthy male volunteers, and provided preliminary evidence on the role of ondansetron in reducing the reinforcing properties of alcohol and the desire to use by 5-HT3 receptor blockade. Later phase 2 clinical studies (([4], [5]) suggested differential effects among AUD patients depending on the age of onset (Early-onset alcoholism EOA/Late-onset alcoholism LOA) [6], <<<<which is hypothesized to be linked to individual genetic variations. Interestingly, compared to placebo, ondansetron was associated with reduced drinking and significant reduction in overall craving in randomized placebo-controlled studies ([4] +, Johnson 2002), but only among patients who developed AUD before age 25 only (EOA), and not among late-onset patients (LOA), presumably by ameliorating serotonergic abnormality>>>>.

Furthermore, as the functional state of the serotonin transporter protein (5-HTT) is an important factor of the serotonergic function control, more recent pharmacogenetic studies have investigated the potential role of 5-HTT genotype on drinking behaviors and alcohol craving ([7]). 5-HTT gene polymorphisms, involving two variants, a short form (S) and a long form (L), have been shown to be associated with differential 5-HT neurotransmission, which could moderate the rewarding effects and the craving for alcohol, and thereby ondansetron treatment response among AUD suffering patients. In line with this hypothesis, ondansetron was administered in a large study among AUD patients according to the 5-HTT polymorphism <<<<(ref Johnson 2011)>>>>. Participants with the LL genotype significantly reduced their drinking compared to the LS or SS genotype, suggesting that ondansetron could represent an interesting approach for the personalized treatment of AUD according to specific polymorphism of the 5-HTT gene.

Direct inhibition of the 5-HT3 receptor is thus hypothesized to reduce alcohol use and alleviate alcohol craving, that is currently considered as a key determinant of relapse vulnerability as well as a major treatment target. A better knowledge of the potential impact of 5-HT antagonist medication on alcohol consumption and craving in AUD, as well as individual clinical subtypes and genotype associated with treatment response is therefore a critical issue to improve treatment approaches and develop personalized medicine in the pharmacotherapy of alcohol use disorder. The aim of this systematic review is to address this issue by assessing scientific evidence of the efficacy of ondansetron on alcohol consumption and craving as well as clinical or genetic predictors of treatment response.

# Methods

**2.1 Research design**

## The study involved a systematic review of the literature based on the Preferred Reporting Items for Systematic reviews and Meta-Analyses (PRISMA) guidelines (35).

**2.2. Databases and search strategy**

This review was based on the following databases: PUBMED/MEDLINE, Psychinfo, Cochrane, Wiley Online Library. The search was performed for all years <<<<up to December, 2021>>>>.

<<<<The following search terms were used:>>>>

For Medline search, the relevant articles were identified by combining the terms: ("ondansetron"[MeSH Terms] OR "ondansetron"[All Fields]) AND ("alcoholism"[MeSH Terms] OR "alcoholism"[All Fields] OR "alcohol use disorder"[All Fields] OR "alcohol abuse"[All Fields] OR "AUD"[All Fields])

For the Wiley Online Library search we used the keywords "ondansetron" AND "alcoholism".

For the PsycInfo search, the keywords were "ondansetron or zofran" AND "alcoholism or alcohol dependence or alcohol abuse or alcoholic or alcohol addiction".

Finally, the Cochrane Library was used by searching ("alcohol use disorder" OR "alcohol dependance") AND "ondansetron" in "Title Abstract Keyword".

**2.3. Eligibility criteria**

Studies were included if they met the following inclusion criteria :

* Reported as a peer reviewed journal
* Concerning individuals suffering from AUD, with no restrictive criteria regarding age, sex, ethnic origin, or place of living.
* Assessing the impact of ondansetron on AUD and/or predictors of ondansetron treatment response
* Papers published in English

Studies were excluded if they were :

* Preclinical studies, reviews, opinion papers, protocols, case reports,
* Animal studies
* Studies in healthy volunteers
* Not published in English

**2.4. Study selection**

## Two authors independently examined all titles and abstracts. Relevant articles were obtained in full-text and assessed for inclusion criteria separately by the two reviewers based on the inclusion and exclusion criteria previously mentioned. Disagreements were resolved via discussion of each article for which conformity to inclusion and exclusion criteria were uncertain and a consensus was reached. The reference lists of major papers were also manually screened in order to ensure comprehensiveness of the review. All selected studies were read in full to confirm inclusion criteria, study type and study population.

**2.5. Assessment of risk of bias in included studies**

Two review authors independently assessed the risk of bias of each included study using the revised Cochrane tool for assessing risk of bias in randomized trials (RoB 2 [27]), in accordance with methods recommended by Cochrane collaboration. The risk-of-bias plot in Figure 2 was generated using the Robvis online tool [28]. The following judgments were used : high risk, low risk or unclear (either lack of information or uncertainty over the potential for bias). Authors resolved disagreements by consensus, and a third author was consulted to resolve disagreements if necessary.

The Newcastle Ottawa Scale (NOS, [29]) was used for assessing single-arm non-randomized studies. However, it had to be adapted by removing the Comparability item for two of the studies that lacked a control group.

**2.6. Collecting data**

<<<<Sample characteristics (including socio-demographic data, comorbidity), and information on study design, methods of assessment of clinical outcomes, and treatment dosage were extracted.>>>>

# 3. Results

**3.1. Study selection**

A total of 90 results were found in the MEDLINE database, 68 on PsycInfo, 35 on Cochrane Library and 8 additional studies were found on Wiley Online Library. A total of 134 articles were identified through the search of the databases. <<<<After review of titles and abstracts, x articles were selected for further examination.>>>> After reading the full text, 21 met inclusion criteria for this review. This process is described in the PRISMA flowchart **(Figure 1).** Among the 21 included studies, 18 were randomized controlled trials (RCT) or analysis of previous RCT, and 3 were prospective open-label studies ([5], [15], [16]). The study duration ranged from 2 to 12 weeks. The selected articles were published between 1994 and 2015.

**3.2. Quality and risk of bias assessment**

Randomized controlled trials where analyzed using the Cochrane Rob2 tool (see Figure 2 for the traffic-light plot). Two studies had a low concern of bias ([8], [17]), most of the studies showed some concern of bias and one had a high concern ([23]). The high dropout rate of the studies was the most concerning factor and affected the Domain 3 of the Risk of Bias tool which represent bias due to missing outcome result. For the 3 prospective study, the Newcastle-Ottawa scale was used (see Figure 6). The risk of bias was evaluated as acceptable as one study scored 7 out of 9 possible points and the two others scored 6 out of 7 (adapted score).

**3.3. Study results**

Results are presented according to their primary outcomes. Among the 21 included studies, x assessed odansetron efficacy through alcohol use (n=11), craving (n=) and mood effect (n=). X studies examined moderators of treatment outcomes. The first set of studies **(Tables 2, 3 and 4)** investigated alcohol use, craving and mood disturbances as clinical outcomes for odansetron efficacy, while the second set (**Table 5**) examined the moderators of treatment outcomes.

**3.3.1 Sample characteristics**

The included studies involved 11 distinct study populations whom characteristics are described in **Table 1.**

In total, 1088 subjects were enrolled, of which <<<<1071 (%)>>>> met criteria for alcohol use disorder. Patients were mostly males <<<<(n= ;75.3 %)>>>>, with a mean age of 41.0 years.

Patients enrolled were diagnosed as alcohol dependent according to the DSM-III-TR ([8], [4]), DSM-IV ([17], [9], [10], [5], [15], [19]), DSM-IV-TR ([18], [24]), DSM-5 ([26]) or ICD-10 ([23]). Some studies required additional criteria, such as more than 35 standard drinks per week for men or 28 for women ([18], [24]), more than 30 drinks per week for men or 21 for women ([21]), at least 15 standard drinks in the week before enrollement ([26]), more than 3 standard drinks per day and a Michigan Alcohol Screening Test greater than 5 ([9], [4], [10]), an AUDIT score greater than 8 ([19]) or a diagnosis before the age of 25 ([26]). Whereas most trials ([8], [9], [4], [10], [5], [15]) concerned treatment-seeking patients, some ([17], [18], [24]) did not. Drug use and disorder was considered as an exclusion criteria in most included studies. Thus, participants who reported drug use ([5], [17], [18], [19], [24]) and/or had a positive drug screening test ([8], [9], [4], [10], [17], [23]) were often excluded, except for cannabis use in a few trials ([15], [17]). Receipt of alcohol use disorder treatment prior to enrollment was an exclusion criteria; one study ([8]) considered treatment over the previous 12 months, and 3 ([30], [10], [15]) over the previous 30 days.

Some studies excluded participants who had been treated with stimulants, sedatives, hypnotics or with treatment that could have an effect on alcohol consumption or mood ([23], [4], [9], [10], [15]). In one study ([26]), having been treated with naltrexone, acamprosate, disulfiram or topiramate 2 weeks prior inclusion, or current treatment with phenytoin, carbamazepine, rifampicine, apomorphine or tramodol (due to potential interactions with ondansetron) were exclusion criteria.

Psychiatric disorders were exclusion criteria in most studies ([4], [9], [10] [5], [18], [19], [25]). Two studies considered only major diagnosis ([17]) or clinically significant disorders ([23] [15]) as exclusion criteria. In the study of Sellers *et al.* ([8]), a Montgomery/asberg Depression scale score below 15 and a Spielberger State-Trait anxiety inventory score below 55 were required. In contrast, one study ([26]) enrolled only people with a concurrent psychiatric diagnosis.

Many studies also required good health at enrollment and notably excluded frequent AUD comorbidities such as elevated bilirubin ([4], [18]), liver enzymes ([15], [18]), [26]), liver cirrhosis ([23], [26]) or severe alcohol withdrawal ([9], [4], [15]) ([26]).

Considering participants subtypes, 9 studies ([4], [9], [10], [11], [12], [13], [5], [14]), [26]) focused on clinical characteristics based on age of onset, before or after the age of 25 (EOA and LOA, while 6 studies examined patient subtypes according to their genotypes ([19], [21], [20], [25], [24], [22]). The most frequent investigated gene was SLC6A4, coding for the serotonin transporter that contains a polymorphism in the promoter region, the 5-HTT-linked polymorphic region, with a "short" (S) and "long" (L). <<<<Individuals possessing two long alleles (L/L) have been found to respond differently to serotoninergic treatments than those having either one (L/S) or two (S/S) short alleles. Another allele, rs1042173-TT, also on the serotonine transporter gene predicted a better response to ondansetron on alcohol use.>>>> Other genes of interest were HTR3A and HTR3B, which regulate the 5HT3 receptor had polymorphisms which influenced response to ondansetron (rs1150226-AG and rs1176713-GG in HTR3A and rs17614942-AC in HTR3B).

Ondansetron dosage ranged from 1 µg/kg bid (twice a day) to 16 mg per day with the most frequent dosage being 4 µg/kg bid. One study ([26]) used a flexible dosage according to treatment response that could range from 0.5 to 4 mg bid (with a mean dose of 3.24 ± 2.64 mg/day).

Four studies ([9], [10], [11], [17]), involving 127 patients, used ondansetron in combination with naltrexone (50 mg/d). Three studies (n=97 patients) compared ondansetron (0.5 mg/d) to sertraline (200 mg/d) ([18], [25], [24]). Treatment duration ranged from 8 days to 11 weeks. Pill count or riboflavin dosage were used for assessing treatment compliance.

**3.3.2 Treatment outcomes**

Efficacy was most often assessed by evaluating the number of standard drinks and derived variables such as defined by the Alcohol Timeline Followback (TLFB) method [31]. Drinking outcomes were drinks per day (DD), drinks per drinking day (DDD), percentage of day abstinent (PDA), heavy drinking days (days with more than 5 drinks per day), percentage of heavy drinking day (PHDD). Standard drink definition varied across the studies, either 12 g ([4][19]), 13 g ([8] or 14 g [23]) of pure ethanol. Furthermore, objective measures of alcohol use were used in 8 studies with either carbohydrate deficient transferrin (CDT) ([4],[11],[5],[26]), or the volume of alcohol consumed during self-administration ([18], [25], [24]), and 𝛾glutamyltransferase in one study ([26]). <<<<X studies examined the effects of ondasetron on alcohol craving. Some studies ([10], [12][17]) evaluated alcohol craving, either with a visual analogical scale or with the obsessive compulsive drinking scale (OCDS [32]) and the Penn Alcohol Craving Scale (PACS). ([26]) One study ([16]) used the Adolescent Obsessive–Compulsive Drinking Scale (A-OCDS). One study ([17]) used functional magnetic resonance imaging to determine ventral striatum activation during cue-exposure, in addition to cue-induced craving assessment>>>>. Finally, two studies examined the effcts of ondasetron on mood. One study used the Profile of Mood States [33] to evaluate attenuation of mood disturbances, and another study ([26]) the Hamilton Rating Scale for Depression (HRSD), the Young Mania Rating Scale (YMRS), and the Inventory of Depressive Symptomatology–Self-report (IDS-SR).

**3.3.3** **Effects of odonsetron on alcohol use reduction**

***{{{{d’abord RCT puis observationnelles}}}}***

Eleven studies, summarized in **Table 2**, evaluated the impact of ondansetron, alone or in combination with naltrexone, on alcohol use. The main outcomes were mostly self-reported changes in alcohol consumption according to the TLFB method, plasma CDT ([4], [11], [5], [26]), GGT ([26]) or the volume of alcohol consumed during self-administration ([18], [25], [24]).

In a sample of 71 males suffering from alcohol dependence (DSM-III-TR), Sellers *et al.* [8] showed a significant impact of ondansetron on alcohol reduction, when patients drinking more than 10 drinks per day at baseline were excluded from the analysis, with the lower ondansetron dose (0,25 mg/d) producing the greatest reduction from baseline.

One randomized control trial [9] (n = 20) examined the impact of a combination of ondansetron and naltrexone only among EOA participants. Results found significant effect on reduction of drinks per day, drinks per drinking day, as well as a trend in reducing the percentage of abstinent days, compared to placebo.

A subsequent analysis of the sample by Ait-Daoud *et al.* showed that the combination of ondansetron and naltrexone was associated with significantly lower CDT levels [11].

<<<<In the only trial taking place outside of North America, Corrêa Filho *et al.* [23] showed a significant reduction of heavy drinking days (7,8 % vs 11,7%, p=0.02) but not of other alcohol-related clinical outcomes.>>>>

**3.3.4. Effects of odonsetron on alcohol use reduction according patient subtypes**

***{{{{RCT puis observationnelles}}}}***

*EOA vs LOA*

In a later trial (Johnson *et al.* 2000b [4]), Johnson et al. examined odansetron efficacy according to patient subtypes (EOA vs LOA) among a sub-sample of 271 patients. Ondansetron was found to be significantly more effective than placebo in reducing alcohol consumption among EOAs but not LOAs. Ondansetron at 4 µg/kg *b.i.d.* was more effective than placebo on drinks per day (1.56 vs 3.30, p = 0.01), drinks per drinking day (4.28 vs 6.90, p = 0.004), percentage of day abstinent (70.10 vs 50.20, p = 0.02) and mean log CDT ratio (-0.19 vs 0.12, p = 0.01). Among EOAs, all other dosages were superior to placebo on the two first criteria. These results were subsequently replicated by Kranzler *et al.* in 2003 [5], who showed a significant reduction (compared to baseline) in most alcohol-related measures (drinks per day, drinks per drinking days, DrinC total score) among EOAs and LOAs who received ondansetron (4 µg/kg *b.i.d.*). A significant difference was also found according to patient subtypes, with significant greater decrease of drinks per day, drinks per drinking day and DrinC total score among EOAs. Changes in the level of carbohydrate-deficient transferrin were consistent with changes in self-reported drinking behavior.

*Patient genotypes*

In a randomized controlled trial, Johnson *et al.* [19] randomized 283 patients by genotype in the 5'-regulatory region of the 5-HTT gene (LL/LS/SS). Individuals with the L/L genotype receiving ondansetron significantly reduced their alcohol consumption, measured by drinks per drinking day and percentage of days abstinent as compared to placebo (respectively -1.62, p = 0.007 and 11.27%, p = 0.023).

Two other studies examined treatment response according to the L/L genotype using a self-administration experiment. The study of Kenna *et al.* [18], among 15 non-treatment seeking individuals, found that participants with the L/L genotype who were administered ondansetron (4 µg/kg *b.i.d.*) during 3 weeks drank significantly less alcohol during an alcohol self-administration procedure, than their counterparts being treated with sertaline (200 mg per day). In another clinical trial involving 77 participants, the same authors provided limited support that ondansetron may reduce drinking in non treatment-seeking individuals with the LL genotype ([24]). Further analysis ([25]) pinpointed gender differences as L/L women treated with ondansetron and S/L or S/S women treated with sertraline had significantly less drinks per drinking days and drank during alcohol self-administration experiment.

*Alcohol and bipolar disorder participants*

Recently, Sherwood *et al.* [26] evaluated the efficacy of ondansetron in 70 patients suffering from both alcohol use disorder and bipolar disorder. Results showed a trend in greater reduction of drinking as measured by the TLFB method that failed to reach significance.

**3.3.5. Effects on craving and cue-induced craving**

Four studies (**Table 3**) examined the impact of ondansetron on craving.

An analysis of the study population of Johnson *et al.* 2000a ([9]) found that the combination of ondansetron and naltrexone was significantly better than placebo in reducing craving among EOAs [10]. Johnson *et al.* 2002 [12] showed that ondansetron at 4 µg/kg *b.i.d.* was associated with a significant reduction in craving (measured by visual analog scale) compared to the placebo group, but only in EOAs. In contrast, craving was significantly increased by ondansetron at 1 µg/kg *b.i.d.* among LOAs.

In a BOLD-MRI laboratory study, Myrick *et al.* [17] examined ventral striatum activation of AUD patients during cue-exposure, according to treatment group (naltrexone, ondansetron, a combination of both or placebo), and compared to social-drinkers. Ventral striatum activation was significantly reduced in the naltrexone, combination and social-drinker groups, and this reduction was correlated with reduction in craving scores. Ondansetron alone was not significantly effective in the reduction of ventral striatum activation nor craving. There was however a trend in the reduction of both.

The study of Sherwood *et al.* [26] found no differences between ondansetron and placebo groups, among patients with both bipolar disorder and AUD.

**3.3.6. Effects on mood disturbances**

Two studies investigated the impact of ondansetron medication on mood (**Table 4**). One study ([33]) showed that ondansetron significantly reduced mood disturbances as measured by the POMS scale among EAOs only. Sherwood *et al.* [26] showed that ondansetron <<<<was significantly more efficacious than placebo in the reduction of HRSD scores, but not of YMRS or IDS-SR scores>>>>, among patients suffering from both bipolar disorder and AUD.

**3.3.7. Predictors of treatment outcomes**

Finally, six studies, presented in **Table 5,** provided data on predictors of treatment outcomes.

As ondansetron had previously been found useful in early onset alcoholism, Dawes *et al.* [15] evaluated ondansetron among adolescents in a prospective, open-label trial, which showed that ondansetron was safe and well tolerated in adolescents with alcohol dependence. In a subsequent study [16], they found that reduction of drinking correlated with reduction in craving. Roache *et al.* 2008 [14] compared the prediction capabilities of the EOA/LOA typology to the type A/type B typology described by Babor *et al.* 1992 [34][35](derived from Type I/II description by Cloninger *et al.* [36]). The A/B typology better described baseline severity of alcohol dependence but treatment response to ondansetron was significantly better predicted by the EOA/LOA typology.

Seneviratne *et al.* [21] produced some evidence that 5’HTTPLR mRNA levels could be used as biomarker to evaluate treatment effectiveness in L/L-subjects treated with ondansetron.

Two studies analysed the population of Johnson *et al.* 2011 [19] and identified genotypes predicting treatment success. Johnson *et al.* 2013 [20] found 5 genotypes that are highly prevalent in the general population and that predicted efficacy of ondansetron. Finally, Hou *et al.* [22] further worked on identifying ways of predicting effectiviness of ondansetron and found that data mining approaches, such as interaction trees and virtual twins could simplify subgroup identification while limiting statistical errors.

# 4. Discussion

{{{{Résumé des principaux résultats: nbre études, type d’étude, qualité méthodologique, traitements à l’étude

Résumé sur l’efficacité de l’ondasetron

Discuter des sous types et des classification EOA/LOA vs Cloninger}}}}

To our knowledge, the present review is the only systematic review assessing efficacy of including ondansetron for the treatment of alcohol use disorder. A subsequent research identified 5 systematic reviews in the last 10 years that included the keywords ’ondansetron’ and ’alcohol use disorder’ or ’alcoholism’. Bauer *et al.* 2015 [37] focused on the influence of serotonergic gene variation in substance use pharmacotherapy and included four out of the 21 studies presented here. Naglich *et al.* 2018 [38] focused on combined pharmacotherapy for the treament of alcohol use and thus included 2 studies involving ondansetron and naltrexone. Cservenka *et al.* 2017 [39] focused on pharmacogenetics and the implication of ethnic diversity in the treatment of AUD and included 2 papers. Castrén *et al.* 2019 [40] focused on the recent findings in AUD pharmacoterapy and mentioned ondansetron without including any of the clinical trials. Finally, Bharadwaj *et al.* 2018 [41] focused on the pharmacotherapy for relapse prevention in AUD in the Indian setting and also mentioned ondansetron but didn’t include any trial on this topic. One review [42] focuses on the role of the serotonin transporter gene in AUD and thus cites 6 of the latest pharmacogenetics studies.

Most of the clinical trials described in this systematic review have stringent inclusion criteria which greatly limits their external validity. Particularly, patients suffering from dual diagnosis or addicted to several substance (with the exception of nicotine) were often excluded. The high dropout rate (mean dropout rate 35.4%) could impact the validity of the findings, but evidence to identify whether or not dropout rate favors medication is lacking.

Finally, out of the seven registered trials that have no published papers yet, three haven’t had any updates for more than ten years whereas one reported non significant results. This may pose a publication bias that is to be taken in consideration.

**Conclusion** Whereas growing evidence tends to suggest efficacy of ondansetron as a treatment of alcohol use disorder in particular genetic subgroups, further studies are needed to fully conclude. In particular, there is a need for bigger studies evaluating long term changes in alcohol consumption. These studies should also have less exclusion criteria to maximize their external validity.

# References

1. RussellM.Hagan,ArthurButler,JuliaM.Hill,Christopher C. Jordan, Simon J. Ireland, and Michael B. Tyers. Effect of the 5-HT3 receptor antagonist, GR38032f, on responses to injection of a neurokinin agonist into the ventral tegmental area of the rat brain. *European Journal of Pharmacology*, 138(2):303–305, 1987.
2. Kathleen A. Grant and James E. Barrett. Blockade of the discriminative stimulus effects of ethanol with 5-HT3 receptor antagonists. *Psychopharmacology*, 104(4):451–456, 1991.
3. B. A. Johnson, G. M. Campling, P. Griffiths, and P. J. Cowen. Attenuation of some alcohol-induced mood changes and the desire to drink by 5-HT3 receptor blockade: a preliminary study in healthy male volunteers. *Psychopharmacology*, 112(1):142–144, 1993.
4. B. A. Johnson, J. D. Roache, M. A. Javors, C. C. DiClemente, C. R. Cloninger, T. J. Prihoda, P. S. Bordnick, N. Ait-Daoud, and J. Hensler. Ondansetron for reduction of drinking among biologically predisposed alcoholic patients: A randomized controlled trial. *JAMA*, 284(8):963–971, 2000.
5. Henry R. Kranzler, Amira Pierucci-Lagha, Richard Feinn, and Carlos Hernandez-Avila. Effects of ondansetron in early- versus late-onset alcoholics: A prospective, open-label study:. *Alcoholism: Clinical & Experimental Research*, 27(7):1150–1155, 2003.
6. Vijoy K Varma, Debasish Basu, Anil Malhotra, Avneet Sharma, and Surendra K Mattoo. Correlates of earlyand late-onset alcohol dependence. *Addictive behaviors*, 19(6):609–619, 1994.
7. Nassima Ait-Daoud, John D Roache, Michael A Dawes, Lei Liu, Xin-Qun Wang, Martin A Javors, Chamindi Seneviratne, and Bankole A Johnson. Can serotonin transporter genotype predict craving in alcoholism? *Alcoholism: Clinical and Experimental Research*, 33(8):1329–1335, 2009.
8. Edward M. Sellers, Tony Toneatto, Myroslava K. Romach, Gail R. Somer, Linda C. Sobell, and Mark B. Sobell. Clinical efficacy of the 5-HT3 antagonist ondansetron in alcohol abuse and dependence. *Alcoholism: Clinical and Experimental Research*, 18(4):879–885, 1994.
9. Bankole A. Johnson, Nassima Ait-Daoud, and Thomas J. Prihoda. Combining ondansetron and naltrexone effectively treats biologically predisposed alcoholics: From hypotheses to preliminary clinical evidence. *Alcoholism: Clinical and Experimental Research*, 24(5):737–742, 2000.
10. N. Ait-Daoud, B. A. Johnson, T. J. Prihoda, and I. D. Hargita. Combining ondansetron and naltrexone reduces craving among biologically predisposed alcoholics: preliminary clinical evidence. *Psychopharmacology*, 154(1):23–27, 2001.
11. Nassima Ait-Daoud, Bankole A. Johnson, Martin Javors, John D. Roache, and Nursen A. Zanca. Combining ondansetron and naltrexone treats biological alcoholics: Corroboration of self-reported drinking by serum carbohydrate deficient transferrin, a biomarker. *Alcoholism: Clinical and Experimental Research*, 25(6):847–849, 2001.
12. Bankole A. Johnson, John D. Roache, Nassima AitDaoud, Nursen A. Zanca, and Madeline Velazquez. Ondansetronreducesthecravingofbiologicallypredisposed alcoholics. *Psychopharmacology*, 160(4):408– 413, 2002.
13. Bankole A. Johnson, Nassima Ait-Daoud, Jennie Z. Ma, and Yanmei Wang. Ondansetron reduces mood disturbance among biologically predisposed, alcoholdependent individuals:. *Alcoholism: Clinical & Experimental Research*, 27(11):1773–1779, 2003.
14. John D. Roache, Yanmei Wang, Nassima Ait-Daoud, and Bankole A. Johnson. Prediction of serotonergic treatment efficacy using age of onset and type a/b typologies of alcoholism. *Alcoholism, Clinical and Experimental Research*, 32(8):1502–1512, 2008.
15. Michael A. Dawes, Bankole A. Johnson, Nassima AitDaoud, Jennie Z. Ma, and Jack R. Cornelius. A prospective, open-label trial of ondansetron in adolescents with alcohol dependence. *Addictive Behaviors*, 30(6):1077–1085, 2005.
16. Michael A. Dawes, Bankole A. Johnson, Jennie Z. Ma, Nassima Ait-Daoud, Suzanne E. Thomas, and Jack R. Cornelius. Reductions in and relations between "craving" and drinking in a prospective, open-label trial of ondansetron in adolescents with alcohol dependence. *Addictive Behaviors*, 30(9):1630–1637, 2005.
17. Hugh Myrick, Raymond F Anton, Xingbao Li, Scott Henderson, Patrick K Randall, and Konstantin Voronin. Effect of naltrexone and ondansetron on alcohol cue–induced activation of the ventral striatum in alcohol-dependent people. *Archives of general psychiatry*, 65(4):466–475, 2008.
18. George A. Kenna, William H. Zywiak, John E. McGeary, Lorenzo Leggio, Chinatsu McGeary, Shirley Wang, Andrea Grenga, and Robert M. Swift. A within-group design of nontreatment seeking 5HTTLPR genotyped alcohol-dependent subjects receiving ondansetron and sertraline. *Alcoholism, Clinical and Experimental Research*, 33(2):315–323, 2009.
19. Bankole A. Johnson, Nassima Ait-Daoud, Chamindi Seneviratne, John D. Roache, Martin A. Javors, Xin-

Qun Wang, Lei Liu, J. Kim Penberthy, Carlo C. DiClemente, and Ming D. Li. Pharmacogenetic approach at the serotonin transporter gene as a method of reducing the severity of alcohol drinking. *The American Journal of Psychiatry*, 168(3):265–275, 2011.

1. Bankole A. Johnson, Chamindi Seneviratne, Xin-Qun Wang, Nassima Ait-Daoud, and Ming D. Li. Determination of genotype combinations that can predict the outcome of the treatment of alcohol dependence using the 5-HT(3) antagonist ondansetron. *The American Journal of Psychiatry*, 170(9):1020–1031, 2013.
2. Chamindi Seneviratne and Bankole A. Johnson. Serotonin transporter genomic biomarker for quantitative assessment of ondansetron treatment response in alcoholics. *Frontiers in Psychiatry*, 3:23, 2012.
3. Jue Hou, Chamindi Seneviratne, Xiaogang Su, Jeremy Taylor, Bankole Johnson, Xin-Qun Wang, Heping Zhang, Henry R. Kranzler, Joseph Kang, and Lei Liu. Subgroup identification in personalized treatment of alcohol dependence. *Alcoholism, Clinical and Experimental Research*, 39(7):1253–1259, 2015.
4. João Maria Corrêa Filho and Danilo Antonio Baltieri. A pilot study of full-dose ondansetron to treat heavydrinking men withdrawing from alcohol in brazil. *Addictive Behaviors*, 38(4):2044–2051, 2013.
5. George A. Kenna, William H. Zywiak, Robert M. Swift, John E. McGeary, James S. Clifford, Jessica R. Shoaff, CynthiaVuittonet, SamuelFricchione, Michael Brickley, Kayla Beaucage, Carolina L. Haass-Koffler, and Lorenzo Leggio. Ondansetron reduces naturalistic drinking in nontreatment-seeking alcohol-dependent individuals with the LL 5’-HTTLPR genotype: a laboratory study. *Alcoholism, Clinical and Experimental Research*, 38(6):1567–1574, 2014.
6. George A. Kenna, William H. Zywiak, Robert M. Swift, John E. McGeary, James S. Clifford, Jessica R. Shoaff, Samuel Fricchione, Michael Brickley, Kayla Beaucage, Carolina L. Haass-Koffler, and Lorenzo Leggio. Ondansetron and sertraline may interact with 5-HTTLPR and DRD4 polymorphisms to reduce drinking in non-treatment seeking alcohol-dependent women: Exploratory findings. *Alcohol*, 48(6):515– 522, 2014.
7. E. Sherwood Brown, Meagan McArdle, Jayme Palka, Collette Bice, Elena Ivleva, Alyson Nakamura, Markey McNutt, Zena Patel, Traci Holmes, and Shane Tipton. A randomized, double-blind, placebo-controlled proof-of-concept study of ondansetron for bipolar and related disorders and alcohol use disorder. *European Neuropsychopharmacology: The Journal of the European College of Neuropsychopharmacology*, 43:92– 101, 2021.
8. Jonathan AC Sterne, Jelena Savović, Matthew J Page, Roy G Elbers, Natalie S Blencowe, Isabelle Boutron, Christopher J Cates, Hung-Yuan Cheng, Mark S Corbett, Sandra M Eldridge, et al. Rob 2: a revised tool for assessing risk of bias in randomised trials. *bmj*, 366:l4898, 2019.
9. Luke A. McGuinness and Julian P. T. Higgins. Risk-ofbias VISualization (robvis): An R package and Shiny web app for visualizing risk-of-bias assessments. *Research Synthesis Methods*, 12(1):55–61, 2021.
10. J Peterson, V Welch, M Losos, PJOOHRI Tugwell, et al. The newcastle-ottawa scale (nos) for assessing the quality of nonrandomised studies in meta-analyses. *Ottawa: Ottawa Hospital Research Institute*, pages 1– 12, 2011.
11. B. A. Johnson, C. R. Cloninger, J. D. Roache, P. S. Bordnick, and P. Ruiz. Age of onset as a discriminator between alcoholic subtypes in a treatment-seeking outpatient population. *The American Journal on Addictions*, 9(1):17–27, 2000.
12. Linda C Sobell and Mark B Sobell. *Timeline followback: user’s guide*. Addiction Research Foundation, 1996.
13. Raymond F Anton, Darlene H Moak, and Patricia K Latham. The obsessive compulsive drinking scale: A new method of assessing outcome in alcoholism treatment studies. *Archives of general psychiatry*, 53(3):225–231, 1996.
14. Douglas McNair, Maurice Lorr, and Leo Droppleman. Profile of mood states (poms). 1989.
15. Thomas F. Babor, Michael Hofmann, Frances K. DelBoca, Victor Hesselbrock, Roger E. Meyer, Zelig S. Dolinsky, and Bruce Rounsaville. Types of Alcoholics, I:EvidenceforanEmpiricallyDerivedTypologyBased on Indicators of Vulnerability and Severity. *Archives of General Psychiatry*, 49(8):599–608, 1992.
16. Mark D. Litt, Thomas F. Babor, Frances K. DelBoca, Ronald M. Kadden, and Ned L. Cooney. Types of

Alcoholics, II: Application of an Empirically Derived Typology to Treatment Matching. *Archives of General Psychiatry*, 49(8):609–614, 1992.

1. C Robert Cloninger. A systematic method for clinical descriptionandclassificationofpersonalityvariants: A proposal. *Archives of general psychiatry*, 44(6):573– 588, 1987.
2. Isabelle E. Bauer, David P. Graham, Jair C. Soares, and David A. Nielsen. Serotonergic gene variation in substance use pharmacotherapy: a systematic review. *Pharmacogenomics*, 16(11):1307–1314, 2015.
3. Andrew C. Naglich, Austin Lin, Sidarth Wakhlu, and Bryon H. Adinoff. Systematic Review of Combined Pharmacotherapy for the Treatment of Alcohol Use Disorder in Patients Without Comorbid Conditions. *CNS drugs*, 32(1):13–31, 2018.
4. Anita Cservenka, Megan M. Yardley, and Lara A. Ray. Review: Pharmacogenetics of alcoholism treatment: Implications of ethnic diversity. *The American Journal on Addictions*, 26(5):516–525, 2017.
5. Sari Castrén, Niklas Mäkelä, and Hannu Alho. Selecting an appropriate alcohol pharmacotherapy: review of recent findings. *Current Opinion in Psychiatry*, 32(4):266–274, 2019.
6. Balaji Bharadwaj, Nivedhitha Selvakumar, and Pooja Patnaik Kuppili. Pharmacotherapy for relapse prevention of alcohol use disorder in the Indian setting: A systematic review. *Industrial Psychiatry Journal*, 27(2):163–171, 2018.
7. Miles D. Thompson and George A. Kenna. Variation in the Serotonin Transporter Gene and Alcoholism: Risk and Response to Pharmacotherapy. *Alcohol and Alcoholism (Oxford, Oxfordshire)*, 51(2):164–171, 2016.