

Supplementary Online Content

Jonas DE, Amick HR, Feltner C, et al. Pharmacotherapy for adults with alcohol use disorders in outpatient settings: a systematic review and meta-analysis. JAMA. doi:10.1001/jama.2014.3628.

- eTable 1. Definitions of Alcohol Use Disorders: Evolution of Terminology
- eTable 2. Medications that are FDA-approved for treating adults with alcohol dependence
- eTable 3. Questions for the full technical report (all six) and for this manuscript (1 through 3)
- eTable 4. Characteristics of included studies
- eFigure 1. Return to any drinking for selected medications compared with placebo, sensitivity analysis including studies rated as high or unclear risk of bias
- eFigure 2. Return to heavy drinking for selected medications compared with placebo, sensitivity analysis including studies rated as high or unclear risk of bias
- eFigure 3. Return to any drinking for acamprosate compared with placebo, subgroup analysis by risk of bias
- eFigure 4. Return to any drinking for oral naltrexone (50mg/day) compared with placebo, subgroup analysis by risk of bias
- eFigure 5. Return to heavy drinking for acamprosate compared with placebo, subgroup analysis by risk of bias
- eFigure 6. Return to heavy drinking for oral naltrexone (50mg/day) compared with placebo, subgroup analysis by risk of bias
- eFigure 7. Percent drinking days for acamprosate compared with placebo, subgroup analysis by risk of bias
- eFigure 8. Percent drinking days for oral naltrexone (50mg/day) compared with placebo, subgroup analysis by risk of bias
- eFigure 9. Percent heavy drinking days for oral naltrexone (50mg/day) compared with placebo, subgroup analysis by risk of bias
- eFigure 10. Drinks per drinking day for oral naltrexone (50mg/day) compared with placebo, subgroup analysis by risk of bias
- eTable 5. Summary of findings and strength of evidence for alcohol consumption outcomes and health outcomes for medications used off-label or those under investigation
- eTable 6. Summary of meta-analyses of adverse events from head-to-head trials comparing acamprosate with naltrexone
- eTable 7. Summary of meta-analyses of adverse events from placebo-controlled trials of acamprosate, naltrexone, nalmefene, topiramate, and valproic acid
- eDiscussion Applicability to Primary Care Settings

This supplementary material has been provided by the authors to give readers additional information about their work.

eTable 1. Definitions of Alcohol Use Disorders: Evolution of Terminology^a

| Term | Definition |
|---|---|
| <p>Alcohol use disorder (DSM-5, 2013)¹</p> <p>Levels of severity</p> <p>Mild: 2-3</p> <p>Moderate: 4-5</p> <p>Severe: ≥6</p> | <ol style="list-style-type: none"> 1. Alcohol is taken in larger amounts or over a longer period than intended 2. Persistent desire or unsuccessful efforts to cut down or control alcohol use 3. A great deal of time is spent in activities necessary to obtain alcohol, use alcohol, or recover from its effects 4. Craving, or a strong desire or urge to use alcohol 5. Recurrent alcohol use resulting in a failure to fulfill major role obligations at work, school, or home 6. Continued alcohol use despite having persistent or recurrent social or interpersonal problems caused or exacerbated by the effects of alcohol 7. Important social, occupational, or recreational activities are given up or reduced because of alcohol use 8. Recurrent alcohol use in situations in which it is physically hazardous 9. Alcohol use is continued despite knowledge of having a persistent or recurrent physical or psychological problem that is likely to have been caused or exacerbated by alcohol 10. Tolerance, as defined by either of the following: <ol style="list-style-type: none"> a) A need for markedly increased amounts of alcohol to achieve intoxication or desired effect b) A markedly diminished effect with continued use of the same amount of alcohol 11. Withdrawal, as manifested by either of the following: <ol style="list-style-type: none"> a) The characteristic withdrawal syndrome for alcohol b) Alcohol (or a closely related substance, such as a benzodiazepine) is taken to relieve or avoid withdrawal symptoms |

| | |
|--|--|
| <p>Alcohol abuse (DSM-IV, 2000)⁴</p> | <p>A. A maladaptive pattern of alcohol use leading to clinically significant impairment or distress, as manifested by at least 1 of the following occurring within a 12-month period:</p> <ul style="list-style-type: none"> (1) recurrent alcohol use resulting in a failure to fulfill major role obligations at work, school, or home (e.g., repeated absences or poor work performance related to alcohol use; alcohol-related absences, suspensions, or expulsions from school; neglect of children or household); (2) recurrent alcohol use in situations in which it is physically hazardous (e.g., driving an automobile or operating a machine when impaired); (3) recurrent alcohol-related legal problems (e.g., arrests for alcohol-related disorderly conduct); or (4) continued alcohol use despite having persistent or recurrent social or interpersonal problems caused or exacerbated by the effects of alcohol (e.g., arguments with spouse about consequences of intoxication, physical fights). <p>B. The symptoms have never met the criteria for alcohol dependence.</p> |
| <p>Alcohol dependence (DSM-IV, 2000)⁴ (alcoholism, alcohol addiction)</p> | <p>A maladaptive pattern of alcohol use, leading to clinically significant impairment or distress, as manifested by at least 3 of the following occurring at any time in the same 12-month period:</p> <ul style="list-style-type: none"> (1) tolerance, as defined by either of the following: <ul style="list-style-type: none"> (a) a need for markedly increased amounts of alcohol to achieve intoxication or desired effect; or (b) markedly diminished effect with continued use of the same amount of alcohol; (2) withdrawal, as manifested by either of the following: <ul style="list-style-type: none"> (a) the characteristic withdrawal syndrome for alcohol; or (b) alcohol (or a closely related drug) is taken to relieve or avoid withdrawal symptoms; (3) alcohol is often taken in larger amounts or over a longer period than was intended; (4) there is a persistent desire or unsuccessful efforts to cut down or control alcohol use; (5) a great deal of time is spent in activities necessary to obtain alcohol, use alcohol, or recover from its effects; (6) important social, occupational, or recreational activities are given up or reduced because of alcohol use; or |

| | |
|---|---|
| | (7) alcohol use is continued despite knowledge of having a persistent or recurrent physical or psychological problem that is likely to have been caused or exacerbated by alcohol (e.g., continued drinking despite recognition that an ulcer was made worse by alcohol consumption). |
| Harmful use (ICD-10, 1994) ^{2,3} | A pattern of drinking that is already causing damage to health. The damage may be either physical (e.g., liver damage from chronic drinking) or mental (e.g., depressive episodes secondary to drinking). |

³The included literature used definitions from DSM-III or DSM-IV. DSM-5 (2013) describes a single alcohol use disorder category measured on a continuum from mild to severe, and no longer has separate categories for alcohol abuse and dependence.¹ Clinicians can also add “in early remission,” “in sustained remission,” “on maintenance therapy,” and “in a controlled environment” to the diagnosis.

eTable 2. Medications that are FDA-approved for treating adults with alcohol dependence

| Generic Drug Name | Mechanism | Dosing |
|-------------------|--|---|
| Acamprosate | Thought to modulate hyperactive glutamatergic NMDA receptors | Oral: 666 mg 3 times per day |
| Disulfiram | Inhibits ALDH2, causing accumulation of acetaldehyde during alcohol consumption, which produces a variety of adverse effects such as nausea, dizziness, flushing, and changes in heart rate and blood pressure | Oral: 250 to 500 mg per day |
| Naltrexone | Opioid antagonist; competitively binds to opioid receptors and blocks the effects of endogenous opioids such as β -endorphin | Oral: 50 to 100 mg per day Intramuscular injection: 380 mg per month |

Abbreviations: ALDH2 = aldehyde dehydrogenase; FDA = U.S. Food and Drug Administration; mg = milligram; NMDA = N-methyl-D-aspartate.

eTable 3. Questions for the full technical report (all six) and for this manuscript (1 through 3)

| | |
|----------|--|
| 1 | <p>a: Which medications are efficacious for improving consumption outcomes for adults with AUDs in outpatient settings?</p> <p>b: How do medications for adults with AUDs compare for improving consumption outcomes in outpatient settings?</p> |
| 2 | <p>a: Which medications are efficacious for improving health outcomes for adults with AUDs in outpatient settings?</p> <p>b: How do medications for adults with AUDs compare for improving health outcomes in outpatient settings?</p> |
| 3 | <p>a: What adverse effects are associated with medications for adults with AUDs in outpatient settings?</p> <p>b: How do medications for adults with AUDs compare for adverse effects in outpatient settings?</p> |
| 4 | <p>Are medications for treating adults with AUDs effective in primary care settings?</p> |
| 5 | <p>Are any of the medications more or less effective than other medications for men or women, older adults, young adults, racial or ethnic minorities, smokers, or those with co-occurring disorders?</p> |
| 6 | <p>Are any of the medications more or less effective for adults with specific genotypes (e.g., related to polymorphisms of the mu-opioid receptor gene [OPRM1])?</p> |

Abbreviation: AUD = alcohol use disorder

eTable 4. Characteristics of included studies

| Author, Year Trial Name | Arm Dose, mg/day (N) | Rx Dur, Wks (f-u) | Setting | Recruit- ment Method | Age, Yrs by arm ^a | % Non- white | % Fe m | % With Co- occurring Condition | Co-intervention | Risk of Bias |
|--|---|----------------------------|---|---|---|--------------------|----------------|---|--|--------------------|
| Addolorato, 2007 ¹⁴ | BAC 30 (42) Placebo (42) | 12 | University treatment and research center; Italy | People contacting alcohol treatment unit | 49 (range 43-61) 50 (range 44-60) | NR | 24- 31 | Liver cirrhosis 100 Hepatitis B 15 Hepatitis C 29 | Routine psychological support 100% | Med |
| Ahmadi, 2002 ¹⁵ ; Ahmadi, 2004 ¹⁶ | NTX 50 (58) Placebo (58) | 12 | Outpatient treatment; Iran | Self-referral | 43 (10) 43 (9) | NR | 0 | NR | Individual counseling 100% | Un- clear |
| ALK21-014, 2011 ¹⁷ | NTX inj 380 every 4 wks (152) Placebo (148) | 12 | Outpatient; Germany, Austria | NR | 46 (9) 46 (8) | NR | 20 | NR | NR | Med |
| Anton, 1999 ¹⁸ ; Anton, 2001 ¹⁹ | NTX 50 (68) Placebo (63) | 12 | Outpatient academic research center; U.S. | Ads, referrals for treatment- seekers | 41 (10) 44 (10) | 11- 18 | 27- 31 | 0 | CBT 100% | Med |
| Anton, 2003 ²⁰ | ACA 3,000 + CBI + MM (9) ACA 3,000 + MM (9) NTX 100 + CBI + MM (9) NTX 100 + MM (9) Placebo + CBI + MM (9) Placebo + MM (8) ^b | 16 | 11 U.S. academic sites | Ads, community resources, clinical referrals at 11 academic sites | ACA (any): 42 (11) NTX (any): 41 (7) PBO (any) 38 (7) | 17 to 22 | 22 to 33 | NR | As randomized | Med |

| | | | | | | | | | | |
|--|--|------------|---------------------------------|--|--|-----------|-----------|----|--|-----|
| Anton, 2004 ²¹ | NALM 5 (68) NALM 20 (66) NALM 40 (68) Placebo (68) | 12 | Outpatient; U.S. | Ads, recruited as outpatients | 45 (11) 46 (11) 44 (9) 45 (11) | 6-15 | 22- 33 | NR | MET 100% | Med |
| Anton, 2005 ²² | NTX 50+CBT (39) NTX 50+MET (41) Placebo+CBT (41) Placebo+MET (39) | 12 | Outpatient; U.S. | Ads, referred to clinical service | 44 (8) 43 (10) 45 (11) 43 (9) | 8-23 | 21- 27 | NR | CBT and MET as randomized | Med |
| Anton, 2006 ²³ Donovan, 2008 ²⁴ LoCastro, 2009 ²⁵ COMBINE | ACA 3,000+CBI + MM (151) ACA 3,000+MM (152) NTX 100+CBI+MM (155) NTX 100+MM (154) Placebo+CBI+MM (156) Placebo+MM (153) ^b | 16 (68) | 11 academic sites; U.S. | Ads, community resources, clinical referrals | 45 (10) 44 (11) 45 (1) 44 (10) 43 (10) 44 (9) | 23 | 31 | NR | As randomized; Community support group participation (like AA) encouraged | Low |
| Anton, 2008 ²⁶ | ARI 2 to 30 (149) Placebo (146) | 12 | 16 academic centers; U.S. | NR | 47 (9) 47 (9) | 15- 16 | 25- 38 | NR | Enhanced CBT 100% | Med |
| Anton, 2011 ²⁷ | NTX 50 (50) Placebo (50) NTX 50+6 wks gabapentin, with 1,200 maximum dose (50) | 16 | Outpatient; U.S. | NR | 44 (10) 43 (10) 47 (9) | 13 | 18 | NR | Used COMBINE's manual (CBT+MM+12- step techniques) 100% | Med |

| | | | | | | | | | | |
|--|--|-----------------|--|--|--------------------------------------|------|-----------|----------------------------------|--|------|
| Balldin, 2003 ²⁸ | NTX 50+CBT (25) NTX 50+ST (31) Placebo+CBT (30) Placebo+ST (32) | 26 | 10 sites outpatient; Sweden | Newspaper, outpatient treatment | 50 (7) 48 (8) 50 (8) 51 (8) | NR | 9- 23 | 0 | None | Low |
| Baltieri, 2004 ²⁹ | ACA 1,998 (40) Placebo (35) | 12 (24) | Outpatient; Brazil | Patients seeking treatment at outpatient SA clinic | Full sample ⁴ 4 (8) | NR | 0 | 0 | AA encouraged | Med |
| Baltieri, 2008 ³⁰ ; Baltieri, 2009 ³¹ | NTX 50 (49) TOP target 200, maximum 400 (52) Placebo (54) | 12 | Outpatient; Brazil | NR | 44 (7) 46 (9) 43 (9) | 29 | 0 | NR | Psychosocial 100% | High |
| Berger, 2013 ³² | ACA 1,998 (51) Placebo (49) | 12 | U.S.; 2 outpatient primary care clinics | Provider referral and ads | 47 (8) 48 (9) | 9 | 38 | NR | Brief structured behavioral intervention from primary care physician | Med |
| Besson, 1998 ³³ | ACA 1,300 to 1,998 (55) Placebo (55) | 52 (108) | Outpatient; 3 psychiatric treatment centers; Switzerland | From inpatient treatment unit | 42 43 (NR) | NR | 20 | 0 | Routine counseling 100% Voluntary DIS 22-24% | Med |
| Book, 2008 ³⁴ ; Thomas, 2008 ³⁵ | PAR 10-60 (20) Placebo (22) | 16 | NR; U.S. | Ads | 28 (7) 30 (8) | 0-18 | 45- 50 | SAD 100 MDD approx. 10 | None | Med |
| Brady, 2002 ³⁶ | VALP 1,500 (14) Placebo (15) | 12 | Outpatient; U.S. | Newspaper; several treatment settings | 40 (8) 41 (10) | 54 | 62 | 0 | CBT | Med |

| | | | | | | | | | | |
|---------------------------|--|----|---|--|-----------------------|----|-----------|---|---|------|
| Brady, 2005 ³⁷ | SERT 150 (49) Placebo (45) | 12 | Outpatient; U.S. | Ads, outpatient SA treatment programs | 37 (8) 37 (9) | NR | 43- 49 | PTSD 100 Depressive disorder 51 Anxiety disorder 38 | CBT 100% | Med |
| Brown, 2008 ³⁸ | QUET titrated from 25 to 600 over 6 weeks (52) Placebo (50) | 12 | NR; U.S | From community | 39 (10) 38 (9) | 39 | 37 | Bipolar 100 | NR | High |
| Brown, 2009 ³⁹ | NTX 50 (20) Placebo (23) | 12 | Outpatient; university health center; U.S. | Newspaper ads, physician referral, flyers and brochures at clinics | 40 (14) 42 (11) | 26 | 49 | Bipolar (current depressed or mixed mood) 100 Cannabis abuse 21 Cocaine abuse 12 Ampheta- mine abuse 7 | CBT 100% | High |
| Chick, 2000 ⁴⁰ | ACA 1,998 (289) Placebo (292) | 24 | Outpatient; U.K. | Recruited from treatment programs | 43 (NR) 44 (NR) | NR | 16 | 0 | Usual psychosocial outpatient program | Med |
| Chick, 2000 ⁴¹ | NTX 50 (90) Placebo (85) | 12 | Outpatient; U.K. | From patients starting outpatient alcohol rehabilitatio n program | 43 (8) 44 (10) | NR | 25 | 0 | “Usual psychosocial treatment program” | Med |

| | | | | | | | | | | |
|--|--|----|---|---|--------------------|-------|----|--|--|------|
| Chick, 2004 ⁴² | FLUV 100 to 300 (261) Placebo (260) | 52 | 10 outpatient sites; U.K., Eire, Austria, Switzerland | NR | 42 (10) 42 (9) | NR | 35 | NR | Psychosocial TAU at each site | Med |
| Cornelius, 1997 ⁴³ Cornelius, 1995 ⁴⁴ | FLUOX 20-40 (25) Placebo (26) | 12 | Inpatient psychiatric institute; U.S. | Recruited as inpatient | 36 (10) 34 (10) | 53 | 49 | MDD 100% | Usual care: psychotherapy 100% | Med |
| Corrêa Filho, 2013 ⁴⁵ | OND 16 (50) Placebo (52) | 12 | University-based outpatient SA treatment center; Brazil | Enrollees in the SA treatment program at study site | 44 (10) 42 (9) | 60-73 | 0 | NR | Standardized brief cognitive behavioral intervention | High |
| Coskunol, 2002 ⁴⁶ | SERT 100 (30) Placebo (29) | 26 | Inpatient (mean 1 month) followed by 6 months outpatient; SA treatment unit; Turkey | NR | 44 (8) 44 (9) | NR | 0 | For eligibility, required no concurrent Axis I disorders | Thiamine 500 mg/day 100% Pyridoxone 500 mg/day 100% AA during inpatient 100% | Med |
| De Sousa, 2004 ⁴⁷ | DIS 250 (50) NTX 50 (50) | 52 | India; outpatient | Recruited as inpatients | 46 (NR) 43 (NR) | NR | 0 | NR | Supportive group psychotherapy | High |

| | | | | | | | | | | |
|------------------------------|---|----|---|---|----------------------------|-----|----|--------------------------------|---|------|
| De Sousa, 2005 ⁴⁸ | ACA 1,998 (50) DIS 250 (50) | 35 | India; outpatient, private psychiatric hospital | Patients undergoing detoxificatio n | 42 (NR) 43 (NR) | 100 | 0 | NR | Weekly supportive group psychotherapy offered | High |
| Fawcett, 2000 ⁴⁹ | BUS 40 (48) Placebo (52) Lithium 1,200 (56) | 26 | Outpatient; U.S. | Ad, referral, inpatient/ outpatient programs | 39 (8) 40 (8) 41 (8) | 16 | 0 | Depression 48 | Supportive therapy | Med |
| Florez, 2008 ⁵⁰ | NTX 50 (51) TOP 50-400 (51) | 26 | Outpatient SA clinic, referrals; Spain | Recruited when presenting for treatment | 47 (9) 47 (9) | 0 | 15 | Personality disorders 27 | Therapy based on Relapse Prevention Model 100% | High |
| Florez, 2011 ⁵¹ | NTX 50 (91) TOP 200 (91) | 26 | Spain; Outpatient SA clinic, referrals | Recruited and screened when presenting for treatment | 47 (9) 48 (10) | NR | 15 | Personality disorders 23 | BRENDA 100% At least monthly meeting with psychiatrist 100% | High |
| Fogaca, 2011 ⁵² | NTX 50 (20) Placebo (20) NTX 50+PUFA (20) PUFA (20) | 12 | Outpatient; Brazil | Newspaper and radio ads | NR | NR | 0 | NR | None | High |
| Fuller, 1979 ⁵³ | DIS 250 (43) DIS 1 (43) RIB 50 (42) | 52 | Outpatient; VA hospital; U.S. | Patients presenting to VA hospital requesting | Full sample 43 (9) | 61 | 0 | NR | Counseling (unspecified) 100% | Med |

| | | | | | | | | | | |
|---|---|---------|--|---|-------------------------------|----|----|----|--|-----|
| | | | | treatment for alcoholism or patients admitted for alcohol-related illness | | | | | | |
| Fuller, 1986 ⁵⁴ | DIS 250 (202) DIS 1 (204) RIB 50 (199) | 52 | Outpatient; 9 VAMCs; U.S. | Screened as inpatients in 7 centers and outpatients at 2 | 41 (11) 42 (10) 42 (10) | 47 | 0 | NR | Counseling (loosely defined) % NR | Med |
| Garbutt, 2005 ⁵⁵ ; Pettinati, 2009 ⁵⁶ ; Lucey, 2008 ⁵⁷ | NTX inj 380 every 4 weeks (208) NTX inj 190 every 4 weeks (210) Placebo (209) | 26 | Inpatient and outpatient, private and VA; U.S. | NR | 45 (10) 45 (11) 45 (11) | 17 | 32 | NR | BRENDA standardized ST 100% | Med |
| Garbutt, 2010 ⁵⁸ | BAC 30 (40) Placebo (40) | 12 | Outpatient, details NR; U.S. | Newspaper and radio ads | 48 (8) 50 (7) | 4 | 45 | NR | BRENDA 100% | Med |
| Gastpar, 2002 ⁵⁹ | NTX 50 (84) Placebo (87) | 12 | 7 centers; Outpatient; Germany | Outpatient and inpatient recruitment | 43 (10) 42 (10) | 0 | 28 | 0 | Psychosocial treatment | Med |
| Geerlings, 1997 ⁶⁰ | ACA 1,332 to 1,998 (128) Placebo (134) | 26 (52) | Outpatient SA treatment centers; | Recruited from detox patients in same | 40 (9) 42 (8) | NR | 24 | NR | ACA: benzodiazepines 5% Placebo: | Med |

| | | | | | | | | | | |
|--------------------------------------|--|----|---|---|------------------------------|----|-------|--------------------------|--|------|
| | | | Belgium, the Netherlands, and Luxembourg | centers | | | | | benzodiazepines 6% | |
| George, 1999 ⁶¹ | BUS 60 (25) Placebo (24) | 52 | Outpatient; U.S. | Recruited from inpatient research unit at NIAAA | Full sample 42 (range 20-61) | NR | 0 | 0 | Care of psychiatrist and nurse at posthospital clinic 100% | High |
| Gual, 2001 ⁶² | ACA 1,998 (148) Placebo (148) | 26 | Outpatient; multicenter; hospitals; Spain | NR | 41 (9) 41 (9) | NR | 20-21 | NR | NR | Med |
| Gual, 2003 ⁶³ | SERT 50-150 (44) Placebo (39) | 24 | 1 center; Outpatient; Spain | Outpatient alcohol dependence treatment | 46 (9) 47 (10) | NR | 47 | Depression/dysthymia 100 | NR | Med |
| Gual, 2013 ⁶⁴ ESENSE 2 | NALM 20 as-needed (358) Placebo (360) | 24 | Belgium, Czech Republic, France, Italy, Poland, Portugal, Spain; 57 sites | Referrals and ads | 45 (11) 44 (11) | 1 | 26-29 | 0 | BRENDA | Med |
| Guardia, 2002 ⁶⁵ | NTX 50 (101) Placebo (101) | 12 | 7 centers, Outpatient; Spain | Recruited treatment-seeking patients | NR | NR | 25 | NR | Psychosocial | Med |

| | | | | | | | | | | |
|--|--|------------|--|---|-----------------------|-----|-------|----|---|------|
| Guardia, 2004 ⁶⁶ | OLA 5-15 (29) Placebo (31) | 12 (16) | Addictive behavior unit of a hospital psychiatry department; Spain | Treatment-seekers addictive behavior unit | 43 (10) 44 (14) | NR | 23-27 | NR | CBT 100% | Med |
| Heinala, 2001 ⁶⁷ | NTX 50 daily for 12 weeks then targeted+CS (34) Placebo+CS (33) NTX 50 daily for 12 weeks then targeted+ST (29) Placebo+ST (25) | 32 | Outpatient; Finland | Ads | Full sample 46 (8) | NR | 29 | 0 | None | High |
| Huang, 2005 ⁶⁸ | NTX 50 (20) Placebo (20) | 14 | Alcoholism treatment unit of an inpatient psychiatric hospital; 1 week inpatient, remainder outpatient; Taiwan | Recruited as inpatients after admission for detox | 38 (6) 43 (9) | 100 | 0 | NR | Weekly individual psychotherapy sessions 100% | High |
| Johnson, 2003 ⁶⁹ Ma, 2006 ⁷⁰ ; Johnson, 2004 ⁷¹ | TOP 25-300 (75) Placebo (75) | 12 | 1 site; outpatient; U.S. | Newspaper | 42 (9) 42 (9) | NR | 28-40 | 0 | None | Med |

| | | | | | | | | | | |
|--|---|---------|--|--|--------------------|----|-------|---|---------------------------|------|
| Johnson, 2004 ⁷² | NTX inj 400 every 28 days (25) Placebo inj (5) | 17 | 4 centers; outpatient; U.S., France, the Netherlands | NR | 42 (9) 45 (8) | 37 | 27 | NR | Psychosocial support 100% | High |
| Johnson, 2007 ⁷³ Johnson, 2008 ⁷⁴ | TOP 50-300, mean 171 (183) Placebo (188) | 14 | 17 academic sites; U.S. | From academic sites; by newspaper, radio, television ads | 47 (9) 48 (9) | 15 | 26-28 | NR | BBCET 100% | Low |
| Kabel, 1996 ⁷⁵ | FLUOX 20-60 (15) Placebo (13) | 15 | Inpatient SA treatment; U.S. | Inpatient recruitment | Full sample 47 (9) | 46 | 0 | Cocaine use 14% | NR | High |
| Kampman, 2007 ⁷⁶ | QUET 400 (29) Placebo (32) | 12 | Outpatient; U.S. | Community referrals, media ads | Full sample 47 (9) | 46 | 23 | MDD 15 Antisocial personality disorder 11 PTSD 8 Panic disorder 5 Social phobia 5 GAD 3 OCD 2 | BRENDA 100% | High |
| Kampman, 2013 ⁷⁷ | TOP 25-300 ^c (83) Placebo (87) | 12 (13) | Outpatient; U.S. | Treatment-seeking cocaine users | 45 (7) 43 (8) | 17 | 21 | Cocaine dependence 100 | Weekly individual CBCST | Med |

| | | | | | | | | | | |
|--|--|-----------------|--|--------------------------------|--|-------|-------|--|---|-----|
| Karhuvaara, 2007 ⁷⁸ | NALM 10 to 40 Targeted dose ^d (242) Placebo (161) | 28 ^e | 15 sites ^f ; Finland | Mainly by newspaper ads | 50 (9) 49 (8) | 0 | 19 | NR | Some elements of BRENDA | Med |
| Kiefer, 2003 ⁷⁹ Kiefer, 2004 ⁸⁰ Kiefer, 2005 ⁸¹ | ACA 1,998 (40) NTX 50 (40) Placebo (40) ACA 1,998+NTX 50 (40) | 12 | 1 site, Outpatient; Germany | Inpatient withdrawal treatment | 46 (8) 46 (8) 46 (11) 47 (10) | NR | 26 | 0 | Group therapy | Low |
| Killeen, 2004 ⁸² | NTX 50+TAU (54) Placebo+TAU(43) TAU alone (48) | 12 | Outpatient community SA treatment center; U.S. | Clinic treatment seekers | 38 (8) 36 (8) 38 (10) | 24 | 37 | Comorbid psychiatric disorder 51 Additional substance use disorder 35 | Several types and intensities | Med |
| Kranzler, 1994 ⁸³ | BUS 15-60, mean 52.5 (31) Placebo (30) | 12 (38) | Outpatient; university health center; U.S. | Ads | 39 (9) 40 (10) | 0-10 | 20-26 | GAD 37 to 46 Anxiety disorder 50 to 52 MDD 25 to 27 | CBT 100% | Med |
| Kranzler, 1995 ⁸⁴ | FLUOX 20-60, mean 47 (51) Placebo (50) | 12 (38) | Outpatient clinic; U.S. | Ads | Full sample 40 (9) | 5 | 20 | Major depression 14% | Group psychotherapy 79% Individual psychotherapy 21% | Med |
| Kranzler, 2004 ⁸⁵ | NTX inj 150 once a month (185) Placebo inj (157) | 12 | Outpatient; U.S. | Ads, recruited as outpatients | 44 (10) 44 (9) | 17-18 | 33-37 | NR | MET 100% | Med |
| Kranzler, 2009 ⁸⁶ | NTX 50 targeted (38) | 12 | Outpatient; | Media ads, | Full sample | 3 | 42 | Drug use disorder <1 | Brief coping | Med |

| | | | | | | | | | | |
|--|---|----------------|--|---|------------------------------|----|----|--|---|--|
| | NTX 50 once daily (45) Placebo targeted (39) Placebo once daily (41) | | U.S. | local provider referral | 49 (10) | | | Social phobia 3 Antisocial personality disorder 3 Dysthymic disorder <1 Agoraphobia without panic disorder <1 OCD <1 GAD <1 | skills training 100% | |
| Kranzler, 2011 ⁸⁷ ; Kranzler, 2012 ⁸⁸ | SERT 50-200 (63) Placebo (71) | 12 (26) | Outpatient; university health center; U.S. | Primarily ads, some clinician referrals | Full sample 48 (10) | 8 | 19 | Cannabis use disorder 17 Cocaine use disorder 19 Past MDD 21 | Coping skills training 100% | Med |
| Krystal, 2001 ⁸⁹ VACS 425 | NTX 50 for 12 months (209) NTX 50 for 3 months then placebo (209) Placebo (209) | 12 or 52 | Multicenter, outpatient; U.S. | VA clinics | 49 (10) 49 (1) 50 (10) | 37 | 3 | 0 | 12-step facilitation | Med |
| Laaksonen, 2008 ⁹⁰ Laaksonen, 2013 ⁹¹ | ACA 1,998 or 1,333 (81) DIS 100 to 200 (81) NTX 50 (81) | Up to 52 | 6 sites in 5 cities; Finland | Volunteers seeking outpatient treatment | 45 (8) 43 (9) 42 (9) | 0 | 29 | NR | Manual-based CBT targeted to match medication goals | Med for 12-wk outcomes; high for 52-wk |

| | | | | | | | | | | out-comes |
|--------------------------------------|--|------------|---|---|--|-----|----|-------|--|-----------|
| Latt, 2002 ⁹² | NTX 50 (56) Placebo (51) | 12 (26) | 4 hospitals; Outpatient; Australia | NR | Full sample 58 (range 23-70) | NR | 30 | 0 | No extensive psychosocial interventions | Med |
| Lee, 2001 ⁹³ | NTX 50 (35) Placebo (18) | 12 | Mixed: initially inpatient, discharged after 1 month from SA treatment center; Singapore | Direct recruitment from inpatient facility | 46 (9) 44 (10) | ≥88 | 0 | NR | Intensive inpatient rehabilitation program; postdischarge therapy encouraged 100% | High |
| Lhuintre, 1985 ⁹⁴ | ACA 1,000 to 2,250 (42) Placebo (43) | 13 | Outpatient; methadone mainten- ance clinics; France | Recruited as inpatients within 48 hours of admission | 43 (7) 40 (7) | NR | 11 | NR | Meprobamate 100% for first month | High |
| Lhuintre, 1990 ⁹⁵ | ACA 1,332 (279) Placebo (290) | 12 | Outpatient; multicenter; France | Recruited within 48 hours of hospitaliza- tion for alcohol withdrawal | 43 (9) 42 (10) | NR | 18 | NR | Psychotherapy allowed | Uncl |
| Likhitsathian, 2013 ⁹⁶ | TOP 100-300 (53) Placebo (53) | 12 | Outpatient; Thailand | Recruited during inpatient | Full sample 42 (9) | 100 | 0 | MDD 4 | Individual MET sessions plus | High |

| | | | | treatment at 3 sites | | | | | monthly MM | |
|-----------------------------------|--|-------------------|---|----------------------------------|--|-------------|-----------|--|--------------------------------------|------|
| Ling, 1983 ⁹⁷ | DIS 250 (41) Placebo (41) | 37 | Outpatient; VA; U.S. | Unclear | 39 (NR) | NR | NR | Heroin use 80 Marijuana use 36 Other drug use 67 Depression 83 Moderate to high depression 50 | Methadone 100% | High |
| Litten, 2013 ⁹⁸ | VAR 2 (99) Placebo (101) | 13 | 5 outpatient academic sites; U.S. | Ads and at treatment sites | 46 (11) 45 (12) | 30 to 38 | 27- 32 | Marijuana use 12-14 | Computerized self-help program | Low |
| Longabaugh, 2009 ⁹⁹ | NTX 50 for 24 weeks+BST (36) NTX 50 for 12 weeks then placebo for 12 weeks+BST (35) NTX 50 for 24 weeks+MET (33) NTX 50 for 12 weeks then placebo for 12 weeks+MET | 12- 24 (72) | Outpatient; U.S. | Newspaper ads | 45 (7) 46 (8) 44 (7) 44 (8) | 6-14 | 33- 43 | NR | None ^g | Med |

| | | | | | | | | | | |
|---------------------------------------|---|----|--|--|----------------------------|-------|-------|------------------|---|------|
| | (38) | | | | | | | | | |
| Malcolm, 1992 ¹⁰⁰ | BUS target 60, mean 52 (33) Placebo (34) | 26 | 1-2 weeks inpatient, then outpatient; VAMC alcohol dependence treatment unit; U.S. | Screened during inpatient stay for alcohol dependence treatment | 44 (SE 2) 42 (SE 1) | 15-18 | 0 | GAD 100 | None | Med |
| Malec, 1996 ¹⁰¹ | BUS 40 (28) Placebo (29) | 12 | Hospital research center; Canada | Media ad | 42 (8) 41 (8) | NR | 18 | NR | None prescribed but 37% received additional treatment: AA 7% Individual psychotherapy 3% | High |
| Mann, 2013 ¹⁰² PREDICT | ACA 1,998 (172) NTX 50 (169) Placebo (86) | 12 | Germany; NR | Recruited from inpatient facilities of 5 academic medical centers plus 2 state-run psychiatric hospitals | 45 (9) 45 (9) 47 (9) | NR | 23 | NR | MM | Med |
| Mann, 2013 ¹⁰³ ESENSE 1 | NALM 20 as-needed (306) Placebo (298) | 24 | Austria, Finland, Germany, Sweden; 39 sites | Referrals and ads | 51 (10) 52 (9) | <1 | 32-33 | 0 | BRENDA | Med |
| Martinotti, | ARI 5-15 (29) NTX 50 (28) | 16 | Italy; outpatient; | Direct | Full sample | NR | NR | Mood disorder 19 | None required | Med |

| | | | | | | | | | | |
|------------------------------|--|---------|--|--|--------------------------------|----------|----------|----------------------------------|---|------|
| 2009 ¹⁰⁴ | | | university hospital day clinic | recruitment from local facility | 40 (12) | | | Anxiety disorder 11 ^h | | |
| Mason, 1994 ¹⁰⁵ | NALM 10 (7) NALM 40 (7) Placebo (7) | 12 | NR; U.S. | Ads | Full sample 42 (9) | 10 | 29 | 0 | Group therapy 0 to 14% AA 0 to 29% | High |
| Mason, 1996 ¹⁰⁶ | DESIP 200 (37) Placebo (34) | 26 | Psychiatry outpatient departments at 2 urban medical centers; U.S. | Inpatient/outpatient referral and public service announcements | Full sample median 40 (IQR 16) | 38 | 17 | Depression 39 | AA attendance encouraged | High |
| Mason, 1999 ¹⁰⁷ | NALM 20 or 80 (70) Placebo (35) | 12 | Outpatient SA treatment; academic research center; U.S. | Ads, press releases, other non-specified sources | 42 (8) 42 (10) | 17-19 | 31-37 | 0 | CBT 100% | Med |
| Mason, 2006 ¹⁰⁸ | ACA 2,000 (258) ACA 3,000 (83) Placebo (260) | 24 (32) | 21 outpatient clinics ⁱ ; U.S. | Primarily by newspaper ads | 45 (11) 44 (9) 45 (10) | 14-15 | 29-36 | NR | Brief abstinence-oriented protocol-specific counseling and self-help materials 100% | Low |
| Mason, 2014 ¹⁰⁹ | GAB 900 (54) GAB 1,800 (47) Placebo (49) | 12 (36) | Outpatient clinical research site; U.S. | Print and internet ads | 42 (10) 45 (11) 47 (11) | 14 to 26 | 34 to 57 | NR | Weekly manual-guided counseling | Low |
| McGrath, 1996 ¹¹⁰ | IMI 50 to 300 (36) Placebo (33) | 12 | University-based depression | Ads and referrals | 37 (7) 11 (9) ^j | 17-22 | 49-53 | MDD 71-72 Bipolar 11-12 | Individual relapse prevention | Med |

| | | | | | | | | | | |
|--|--|------------|--|--|--|----|----|--|---|-----|
| | | | research clinic; U.S. | | | | | Atypical depression 70-72 Other SA 16 | counseling | |
| Moak, 2003 ¹¹¹ | SERT 50-200 (38) Placebo (44) | 12 | 1 site; outpatient; U.S. | Newspaper, outpatient treatment | 41 (11) 42 (10) | 1 | 39 | Depression/ dysthymia 100 | CBT | Med |
| Monterosso, 2001 ¹¹² | NTX 100 (121) Placebo (62) | 12 | Outpatient; U.S. | Ads | Full sample 46 (12) | 27 | 27 | NR | BRENDA | Med |
| Monti, 2001 ¹¹³ ; Rohsenow, 2007 ¹¹⁴ ; Rohsenow, 2000 ¹¹⁵ | NTX 50 (64) Placebo (64) | 12 (52) | 2 weeks partial hospital (pre-medication); 52 weeks outpatient; U.S. | Recruited from partial hospital program in an urban private psychiatric hospital | Full sample 39 (9) | 3 | 24 | Cocaine use 23 Sedative use 8 Opiate use 4 | Brief physician outpatient contacts (intensive therapy occurred prior to medication portion of trial) | Med |
| Morgenstern, 2012 ¹¹⁶ | NTX 100+MBSCT (51) NTX 100 (51) Placebo+MBSCT (50) Placebo (48) | 12 | NR; U.S. | Ads, community outreach | 38 (10) 39 (11) 42 (11) 40 (11) | 26 | 0 | HIV 15 Any drug use 67 | BBCET 100% | Med |
| Morley, 2006 ¹¹⁷ Morley, 2010 ¹¹⁸ | ACA 1,998 (55) NTX 50 (53) Placebo (61) | 12 | 3 treatment centers with "medical care typically | Patients who had attended an inpatient detox program, | 45 (9) 48 (9) 42 (9) | NR | 30 | Severe concurrent illness (psychiatric or other) – NOS 3 | All offered 4-6 sessions of manualized compliance therapy; up-take / attendance NR | Low |

| | | | | | | | | | | |
|-------------------------------|---|---------|---|--|--|-----|------|---|--|------|
| | | | available at hospital based drug and alcohol treatment services"; Australia | outpatient treatment or followup or who responded to ads | | | | | | |
| Morris, 2001 ¹¹⁹ | NTX 50 (55) Placebo (56) | 12 | Outpatient; Australia | Outpatient, self-referral | 47 (8) 48 (8) | NR | 0 | PTSD 23 GAD 32 Panic disorder 4 MDD 6 BPD 1 | Group psychoeducation and social support | Med |
| Naranjo, 1995 ¹²⁰ | CIT 40 (53) Placebo (46) | 12 (20) | Outpatient research center; Canada | Newspaper ad | 44 (10) 47 (10) | NR | 44 | NR | Brief psychosocial intervention 100% | High |
| Narayana, 2008 ¹²¹ | ACA 1,332 to 1,998 (28) NTX 50 (26) TOP 100 to 125 (38) | 52 | India; military, outpatient | Members of the Armed Forces | 38 (range 30-54) 37 (range 27-51) 40 (range 29-55) | 100 | 0 | NR | Various psychotherapies were offered | High |
| Nava, 2006 ¹²² | DIS 200 (28) NTX 50 (24) GHB 150 mg/kg/day (28) | 52 | Italy; outpatient | Advertisements, word of mouth, press release | 43 (5) 41 (7) 39 (8) | NR | 15 % | 0 | CBT | High |

| | | | | | | | | | | |
|---|---|------------|---|---|--|----|-----|--------------------|--|-----|
| O'Malley, 1992 ¹²³ , O'Malley, 1996 ¹²⁴ | NTX 50+CS (29) NTX 50+ST (23) Placebo+CS (25) Placebo+ST (27) | 12 (38) | Outpatient; university alcohol treatment unit; U.S. | Ads and those seeking treatment at unit | NTX (any) 43 (10) PBO (any) 39 (9) | 7 | 26 | NR | See arms | Med |
| O'Malley, 2007 ¹²⁵ | NTX 50 (57) Placebo (50) Randomization stratified by presence of eating disorder | 12 | University mental health center; U.S. | Newspaper ads and patients seeking SA treatment | Full sample 40 (8) | 11 | 100 | Eating disorder 28 | CBCST 100%, based on manualized approach used in Project MATCH | Med |
| O'Malley, 2008 ¹²⁶ | NTX 50 (34) Placebo (34) NTX 50+SERT 100 (33) | 16 | Outpatient; U.S. | Direct community recruitment, health clinic referral, local ads | 42 (11) 39 (10) 39 (8) | 70 | 34 | NR | MM 100% | Med |
| Oslin, 1997 ¹²⁷ | NTX 100 on Monday and Wednesday, 150 on Friday (21) Placebo (23) | 12 | Outpatient SA clinic and VAMC; U.S. | From a VA hospital | 57 (7) 59 (7) | 64 | NR | 0 | Group therapy and case manager 100% | Med |
| Oslin, 2008 ¹²⁸ | NTX 100+CBT (40) NTX 100+BRENDA (39) NTX 100+doctor only (41) Placebo+CBT (40) | 24 | Outpatient psychiatry clinic; U.S. | Ads in local media | 45 (11) 44 (10) 42 (11) 44 (13) 44 (10) 44 (11) | 27 | 27 | NR | None | Med |

| | | | | | | | | | | |
|--|---|------------|--|---|----------------------------|----|----|---|---|------|
| | Placebo+BREND A (40) Placebo+doctor only (40) | | | | | | | | | |
| Paille, 1995 ¹²⁹ | ACA 1.3 g (188) ACA 2 g (173) Placebo (177) | 52 (78) | NR ^K ; France | Referral from alcohol specialist centers | 44 (9) 43 (8) 43 (9) | NR | 20 | NR | Supportive psychotherapy 100% Hypnotics 6 to 7% Anxiolytics 8 to 12% Antidepressants 8 to 9% | Med |
| Pelc, 1996 ¹³⁰ , Pelc, 1992 ¹³¹ | ACA 1,332 to 1,998 (55) Placebo (47) | 26 | Outpatient; multicenter; Belgium | Post-inpatient detox | 48 (8) 43 (9) | NR | 31 | NR | Supportive psychotherapy 100% | High |
| Pelc, 1997 ¹³² | ACA 1,332 (63) ACA 1,998 (63) Placebo (62) | 13 | Outpatient; after inpatient detox; Belgium, France | Inpatient referral | NR | NR | NR | NR | Counseling, social support when needed 100% | Med |
| Petrakis, 2004 ¹³³ , Ralevski, 2006 ¹³⁴ | NTX 50 (16) Placebo (15) | 12 | At least 3 outpatient centers—MIRECC clinics; U.S. | Direct recruitment from participating centers | 47 (5) 46 (6) | 19 | 0 | Schizophrenia or schizoaffective disorder 100 | CBT+psychiatric TAU Neuroleptics 52% Benzodiazepines 16% Thymoleptics 39% | Med |

| | | | | | | | | | | |
|---|--|----|--|---|--|----|----|--------------------------------|--|---------------------------------------|
| Petrakis, 2005 ¹³⁵ Ralevski, 2007 ¹³⁶ Petrakis, 2007 ¹³⁷ Petrakis, 2006 ¹³⁸ VA MIRECC | DIS 250 (66) NTX 50 (59) Placebo (64) NTX 50+DIS 250 (65) | 12 | Outpatient; VA; U.S. | Recruited as outpatients or ad | 46 (9) 48 (7) 46 (7) 48 (9) | 26 | 3 | Axis I disorder 100 | Psychiatric TAU 100% | Med for NTX; high for DIS |
| Petrakis, 2012 ¹³⁹ | DESIP 200+placebo (24) ^l PAR 40+placebo (20) DESIP 200+NTX 50 (22) PAR 40+NTX 50 (22) | 12 | U.S.; out- patient; multiple mental illness centers, most from VAs | Ads (non- veterans); mental illness centers (veterans) | 47 (10) 49 (9) 47 (10) 45 (7) | 25 | 9 | PTSD 100 | Clinical management/ compliance enhancement therapy 100% | High |
| Pettinati, 2001 ¹⁴⁰ | SERT 200 (50) Placebo (50) | 14 | Outpatient; U.S. | Ads and referral | Full sample 45 (10) | 80 | 48 | Depress-ion 47 | 12-step facilitation | Uncl |
| Pettinati, 2008 ¹⁴¹ | NTX 150 (82) Placebo (82) Participants also randomized to either CBT or BRENDA (2x2 design) ^m | 12 | University- affiliated outpatient SA treatment research facility; U.S. | Those seeking treatment at the facility | Full sample 39 (7) | 76 | 29 | Cocaine depend- ence 100 | NR | Med |
| Pettinati, 2010 ¹⁴² | NTX 100 (49) SERT 200 (40) Placebo (39) SERT 200+NTX | 14 | Outpatient; U.S. | Newspaper ads, referrals from local | 43 (8) 44 (12) 43 (9) 43 (10) | 35 | 38 | Depress-ion 100 | CBT 100% | Med |

| | | | | | | | | | | |
|--|--|------------|---|---|--------------------|-----------|-----------|--|---|------|
| | 100 (42) | | | professional or friends/ family | | | | | | |
| Plebani, 2013 ¹⁴³ | VAR 2 (19) Placebo (21) | 12 (13) | Outpatient; U.S. | Ads, treatment seekers | 45 (12) 48 (11) | 29- 58 | 9- 21 | NR | Individual weekly MM | High |
| Poldrugo, 1997 ¹⁴⁴ | ACA 1,332 to 1,998 (122) Placebo (124) | 26 (52) | Inpatient for 1-2 weeks then outpatient; multicenter community based alcohol rehabilita- tion program; Italy | From acute inpatient withdrawal treatment | 43 (10) 45 (9) | NR | 23- 31 | 0 | Community- based rehabilitation program with group sessions, alcohol education, community meetings 100% | Med |
| Ralevski, 2011 ¹⁴⁵ , Ralevski, 2011 ¹⁴⁶ | ACA 1,998 (12) Placebo (11) | 12 | Outpatient; university and VA health centers; U.S. | From community and through referrals from treatment facilities at a university and a VA facility | 52 (9) 49 (5) | 65 | 17 | Schizo- phrenia spectrum disorders 100 | Weekly skills training that incorporated CB drug relapse prevention strategies 100% | High |
| Rubio, 2001 ¹⁴⁷ | ACA 1,665-1,998 (80) | 52 | Spain; | Patients presenting to hospital | 44 (12) 43 (10) | NR | 0 | 0 | Supportive group therapy weekly; weekly | High |

| | | | | | | | | | | |
|------------------------------|---|------------|---|------------------------|--------------------------|----|----|---|---|------|
| | NTX 50 (77) | | outpatient | for detox | | | | | visits with a psychiatrist for 3 months, then biweekly until end of study | |
| Rubio, 2009 ¹⁴⁸ | TOP 250 (31) Placebo (32) ⁿ | 12 | Outpatient; Spain | NR | 43 (9) 42 (9) | NR | 0 | NR | Supportive group therapy offered | High |
| Salloum, 2005 ¹⁴⁹ | VALP 750+ (29) Placebo (30) | 24 | Outpatient SA service at university clinic; U.S. | Treatment seekers | 37 (9) 38 (9) | 25 | 29 | Bipolar I disorder 100 Mixed bipolar subtype 58 Manic 21 Depress-ed 21 Cannabis abuse or depen- dence 29 Cocaine abuse 29 | Lithium and weekly individual dual diagnosis recovery counseling 100% | Med |
| Sass, 1996 ¹⁵⁰ | ACA 1,332 to 1,998 (136) Placebo (136) | 48 (96) | Psychiatric outpatient; Germany | Outpatient referral | 42 (8) 41 (9) | NR | 22 | NR | Counseling / psychotherapy 100% | Med |
| Schmitz, 2004 ¹⁵¹ | NTX 50+RPT (20) NTX 50+DC (20) Placebo+RPT (20) Placebo+DC (20) | 12 | Outpatient; U.S. | Ads | Full sample 36 (6) | 71 | 16 | Cocaine depend- ence 100 | RPT or DC as randomized | High |

| | | | | | | | | | | |
|-------------------------------|--|---------|--|---|--|-------|----|-----------------------------|-----------------------------------|------|
| Schmitz, 2009 ¹⁵² | NTX 100+CBT (20) NTX 100+CBT and CM (25) Placebo+CBT (27) Placebo+CBT and CM (14) | 12 | Outpatient SA clinic; U.S. | Media ads | 33 (1) 33 (1) 33 (1) 42 (7) | 84-93 | 13 | Cocaine use disorder 100 | CBT 100% | High |
| SENSE, 2013 ¹⁵³ | NALM 20 as-needed (509) Placebo (166) | 52 (56) | Czech Republic, Estonia, Hungary, Latvia, Lithuania, Poland, Russian Federation, Slovakia, Ukraine, U.K. | NR | 44 (11) 44 (12) | NR | 23 | NR | NR | High |
| Stedman, 2010 ¹⁵⁴ | QUET 300-800 (175) Placebo (186) | 12 | Outpatient; multicenter; U.S. | NR | 39 (9) 38 (10) | 12 | 37 | Bipolar 100 | None | High |
| Tempesta, 2000 ¹⁵⁵ | ACA 1,998 (164) Placebo (166) | 26 (39) | Outpatient; Italy | Recruited from outpatient internal medicine, neurology and addiction treatment programs | 46 (11) 46 (11) | NR | 17 | 0 | Medical and behavioral counseling | Med |

| | | | | | | | | | | |
|--|--|-----------------|---|---|----------------------------------|-----------|-----------|---|---|------|
| Tiihonen, 1996 ¹⁵⁶ | CIT 40 (31) Placebo (31) | 13 (17) | Outpatient; community- based alcohol rehabilita- tion center; Finland | Inpatient / outpatient referral | 45 (7) 47 (9) | NR | 0 | 0 | Supportive psychotherapy intervention 100% | High |
| Tollefson, 1991 ¹⁵⁷ ; Tollefson, 1992 ¹⁵⁸ | BUS 15-60 (26) Placebo (25) | 24 | Outpatient; U.S. | Referred based on recent inpatient SA treatment discharges | Full sample 38 (SE 0.4) | NR | 44 | GAD with depress-ive features 100% | Controlled AA participation | High |
| Volpicelli, 1995 ¹⁵⁹ Volpicelli, 1992 ¹⁶⁰ | NTX 50 (54) Placebo (45) ^o | 12 | SA treatment unit of a VAMC; U.S. | Patients in the SA treatment program of a VAMC | 44 (9) 43 (9) | ≥78 | 0 | NR | Outpatient treatment program and group therapy 100% | Uncl |
| Volpicelli, 1997 ¹⁶¹ | NTX 50 (48) Placebo (49) | 12 | Outpatient SA treatment, university/V A treatment research center; U.S. | Receiving outpatient treatment | 39 (9) 38 (9) | 60- 65 | 18- 26 | NR | Counseling 100% | Med |
| Whitworth, 1996 ¹⁶² | ACA 1,332 or 1,998 (224) Placebo (224) | 52 (104) | Outpatient specialty; Austria | Inpatient recruitment | 42 (8) 42 (9) | NR | 21 | NR | NR | Med |
| Wilens, 2008 ¹⁶³ | ATO 25 to 100 (72) | 12 | Multi- institution; | NR | 34 (10) 35 (10) | 12 | 15 | ADHD 100 | 12-step allowed; all other co- | High |

| | Placebo (75) | | U.S. and Canada | | | | | | interventions prohibited | |
|-----------------------------|--|---------|--|---------------------------------|----------------------------|----|----|----|--------------------------|-----|
| Wolwer, 2011 ¹⁶⁴ | ACA 1,998+IBT (124) ACA 1,998+TAU (122) ^d Placebo+IBT (125) | 24 (52) | Outpatient; 4 university hospitals 1 non-academic clinic; Germany | Recruited after inpatient detox | 45 (8) 46 (8) 46 (8) | NR | 29 | NR | NR | Med |

^a Age is reported as mean (SD) by treatment group unless otherwise noted.

^b Three additional treatment arms were included in COMBINE but were not relevant to our Key Questions: ACA + NTX + CBI + MM, ACA + NTX + MM, and CBI only (no pills).

^c Dose was titrated up from 25 to 300mg/day over 8 weeks

^d Targeted dosing; medication was taken when participants believed drinking to be imminent, rather than as a daily scheduled medication.

^e 52 weeks total (28 weeks of initial nalmefene vs. placebo, then another randomization for nalmefene responders).

^f Sites included 5 specialist treatment clinics, 6 private general practices, 2 occupational health care offices, and 2 outpatient clinical research facilities

^g This study is not focused on NTX versus placebo comparison; it is a different design and has 4 arms, aiming to compare 12 versus 24 weeks of NTX and to compare MET versus BST (to determine whether the type of psychosocial treatment delivered in combination with duration of NTX may partially explain inconsistent findings regarding efficacy of NTX).

^h Study also reported the following percentages of participants with co-occurring disorders: impulse control disorder 5%, eating disorder 1%, somatoform disorder 1%. Personality disorders: borderline 8%, antisocial 4%, avoidant 4%, histrionic 1%, paranoid 1%, dependent 1%, passive-aggressive 1%, schizoid 1%, cannabis abuse 12%, cocaine abuse 8%, benzodiazepine abuse 1%, MDMA abuse 1%.

ⁱ Clinics were affiliated with academic medical centers and had investigators experienced in alcoholism treatment.

^j The study reported 11 years, but it was clearly a reporting error; likely 31 or 41 years.

^k The article was not explicit about the setting, but patients received psychotherapy and psychiatric medication management suggesting a psychiatric outpatient setting.

^l Because 2 of the 4 arms are combinations, they are not eligible/not comparisons of interest; only the head-to-head comparison of paroxetine+placebo and desipramine+placebo is eligible.

^m Study stratified randomization by sex and reports the results overall and separately by sex.

ⁿ Numbers entered are those analyzed; 76 total were randomized, but dropouts were not reported by arm.

^o Data are from Volpicelli 1995,¹⁵⁹ which reported pooled results of 99 participants. Data from a smaller subset (N=70) of this sample was reported in Volpicelli 1992.¹⁶⁰ For our data analyses, we used data from Volpicelli 1995 to use the larger, more complete sample and did not use data from Volpicelli 1992 to avoid double counting.

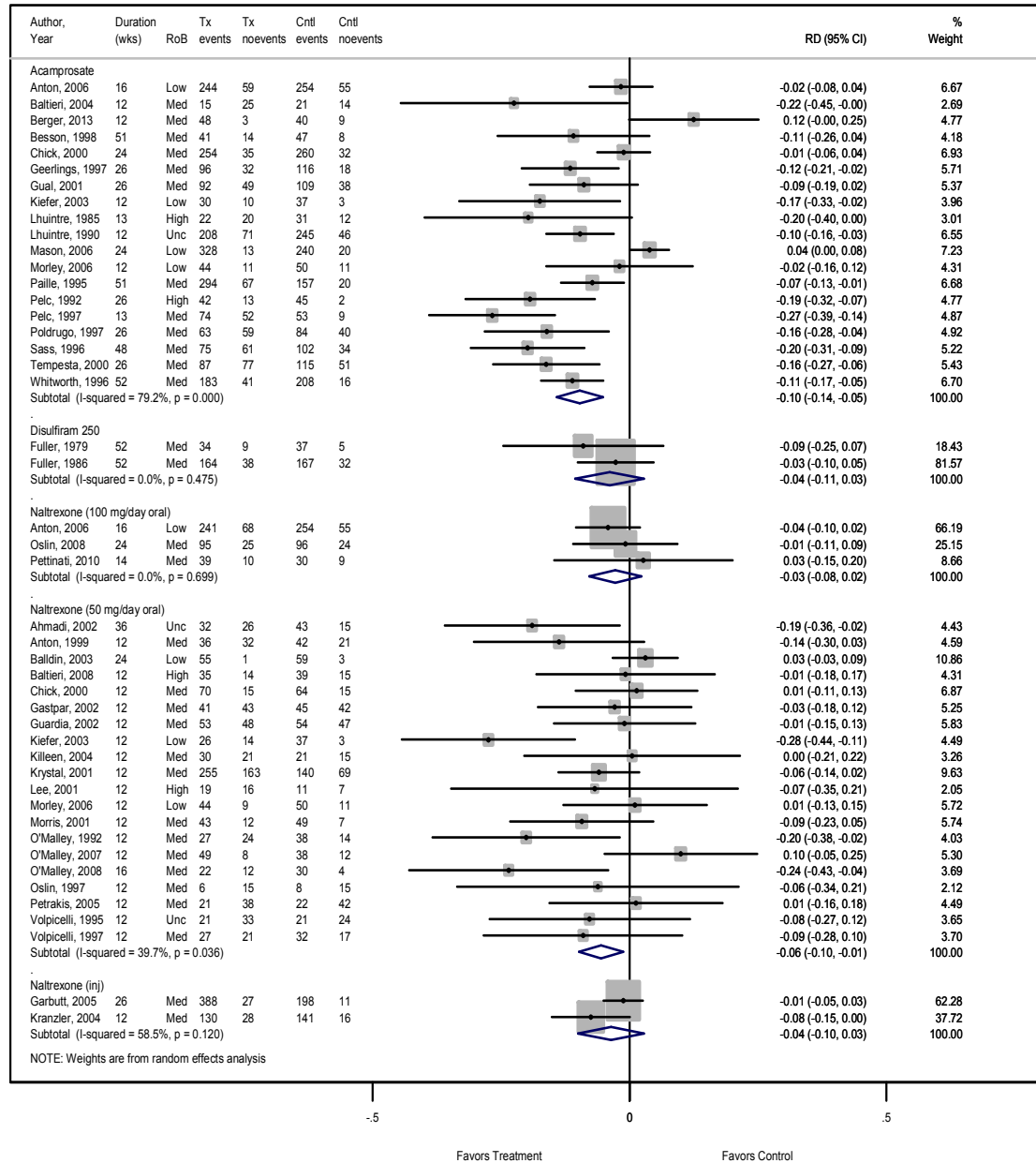
^p Treatment as usual, seen once per week in an individual setting; MI techniques allowed.

Abbreviations: AA = Alcoholics Anonymous; ACA = acamprosate; ADHD = attention deficit hyperactivity disorder; ARI = aripiprazole; ATO = atomoxetine; BAC = baclofen; BBCET = brief behavioral compliance enhancement treatment; BPD = bipolar disorder; BRENDA = BRENDA is an acronym based on the components of the intervention: (B)rief psychosocial evaluation, (R)eport to the patient on assessment, (E)mpathic understanding of the patient's situation, (N)eeds collaboratively identified by the patient and treatment provider, (D)irect advice to the patient on how to meet those needs, (A)ssess reaction of the patient to advice and adjust as necessary for best care; BST = broad spectrum treatment; BUS = buspirone; CB = cognitive behavioral; CBCST = cognitive behavioral coping skills therapy; CBI = combined behavioral intervention; CBT = cognitive behavioral therapy; CIT = citalopram; CS = coping skills; DC = drug counseling; DESIP = desipramine; detox = detoxification; DIS = disulfiram; Dur = duration; f-u = followup; Fem = female; FLUOX = fluoxetine; FLUV = fluvoxamine; g = grams; GAB = gabapentin; GAD = generalized anxiety disorder; GHB = γ -Hydroxybutyric acid; HIV = human immunodeficiency virus; IBT = integrative behavior therapy; IMI = imipramine; inj = injectable; IQR = interquartile range; kg = kilogram; MBSCT = modified behavioral self-control therapy; MDD = major depressive disorder; Med = medium; MET = motivational enhancement therapy; mg = milligram; MIRECC = Mental Illness Research, Education and Clinical Center; MM = medical management; N = number; NALM = nalmefene; NIAAA = National Institute on Alcohol Abuse and Alcoholism; NOS = not otherwise specified; NR = not reported; NTX = naltrexone; OCD = obsessive-compulsive disorder; OLA = olanzapine; OND = ondansetron; PAR = paroxetine; PTSD = post-traumatic stress disorder; PUFA = polyunsaturated fatty acid; QUET = quetiapine; RIB = riboflavin;

RPT = relapse prevention therapy; Rx = medication; SA = substance abuse; SE = standard error; SAD = social anxiety disorder; SERT = sertraline; ST = supportive therapy; TAU = treatment as usual; TOP = topiramate; U.K. = United Kingdom; U.S. = United States; Uncl = unclear; VA = Veterans Affairs; VACS = Veterans Affairs Cooperative Study; VALP = valproic acid; VAMC, Veterans Administration Medical Center; VAR = varenicline; Wks = weeks; Yrs = years.

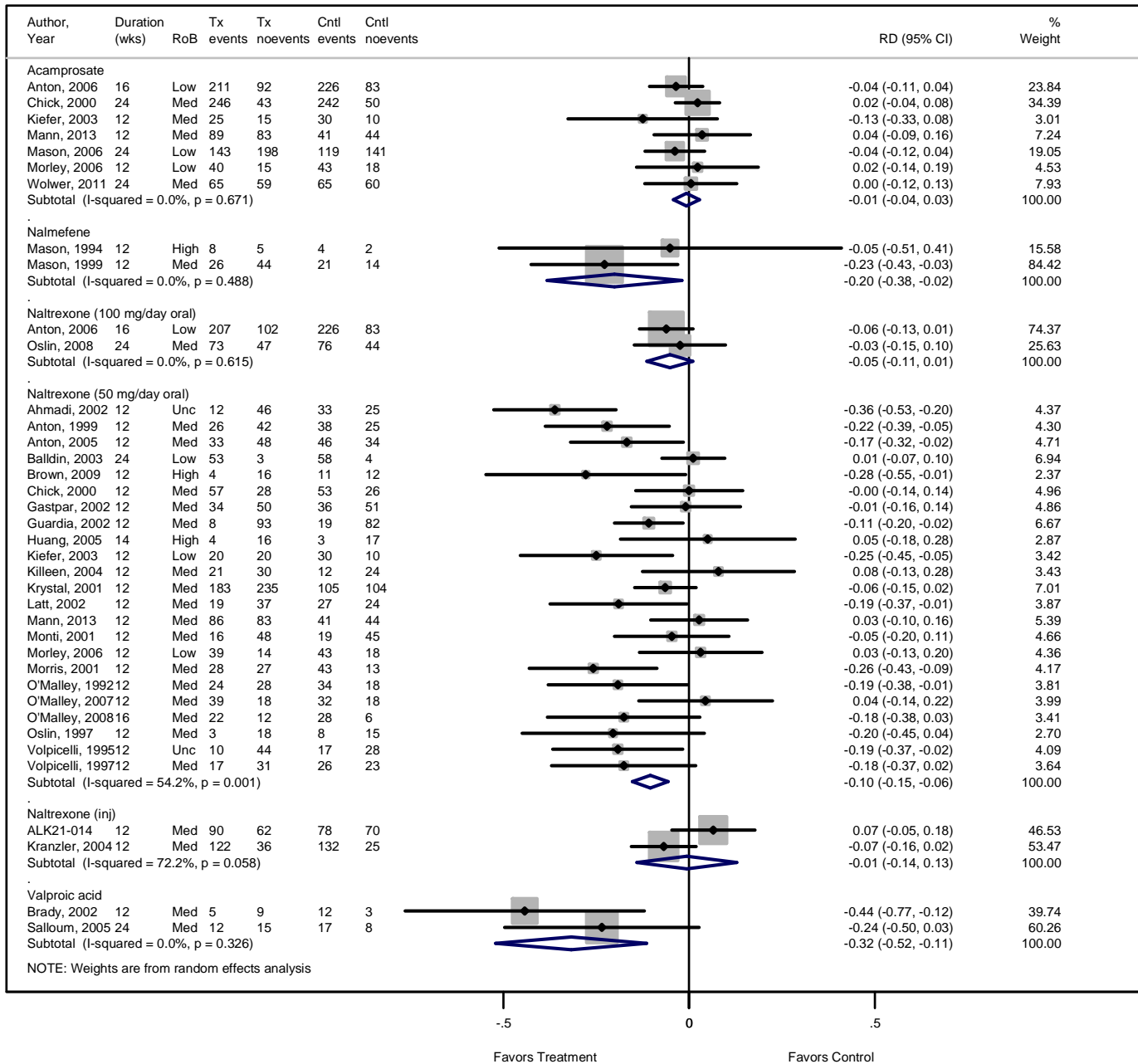
eFigure 1. Return to any drinking for selected medications compared with placebo, sensitivity analysis including studies rated as high or unclear risk of bias

Sensitivity Analysis - Return to Any Drinking Selected Medications Compared with Placebo



Abbreviations: CI = confidence interval; Cntl events = number of events in the control group; Cntl noevents = number of no events in the control group; Med = medium; RD = risk difference; RoB = risk of bias; Tx events = number of events in treatment group; Tx noevents = number of no events in the treatment group; wks =weeks.

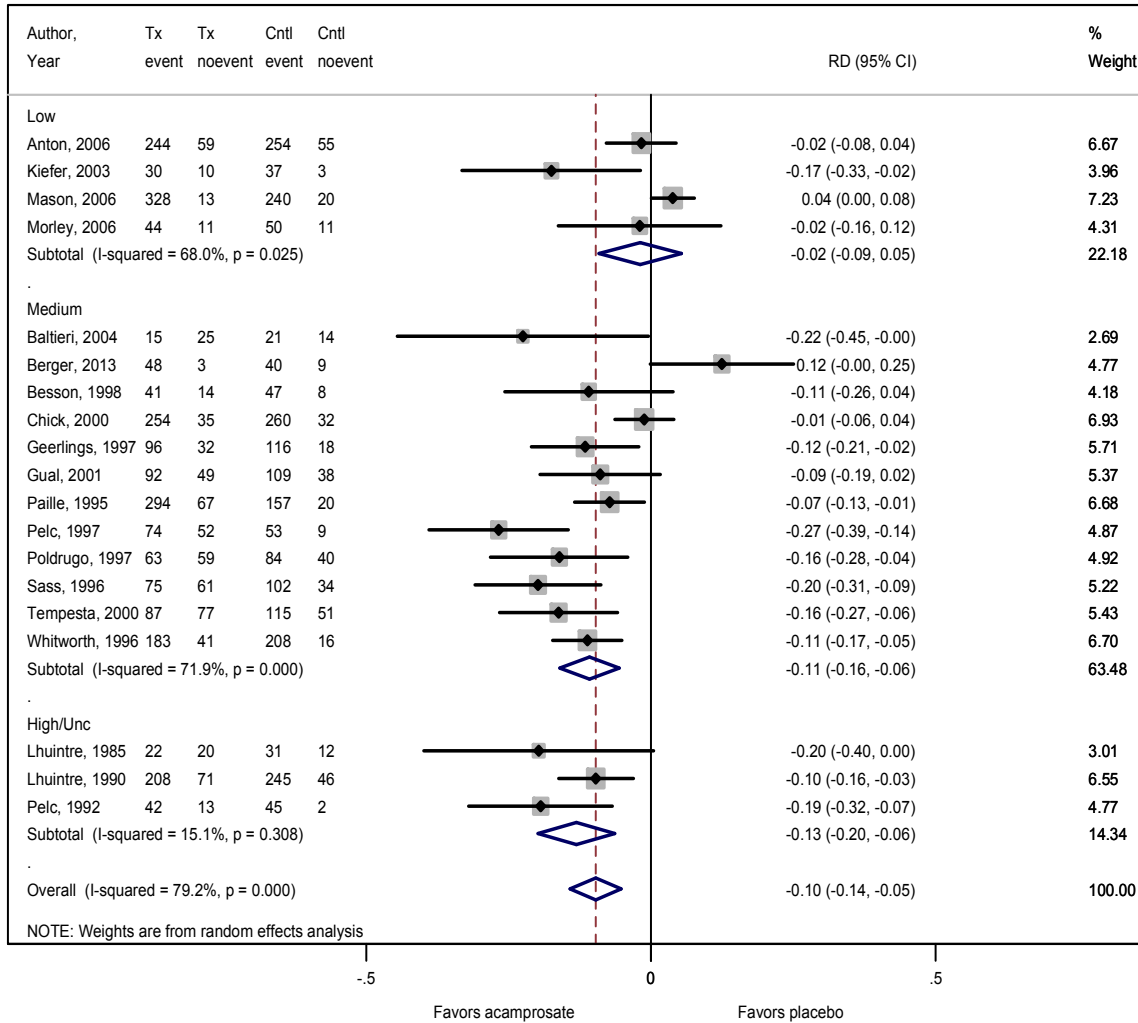
eFigure 2. Return to heavy drinking for selected medications compared with placebo, sensitivity analysis including studies rated as high or unclear risk of bias



Abbreviations: CI = confidence interval; Cntl events = number of events in the control group; Cntl noevents = number of no events in the control group; Med = medium; RD = risk difference; RoB = risk of bias; Tx events = number of events in treatment group; Tx noevents = number of no events in the treatment group; wks =weeks.

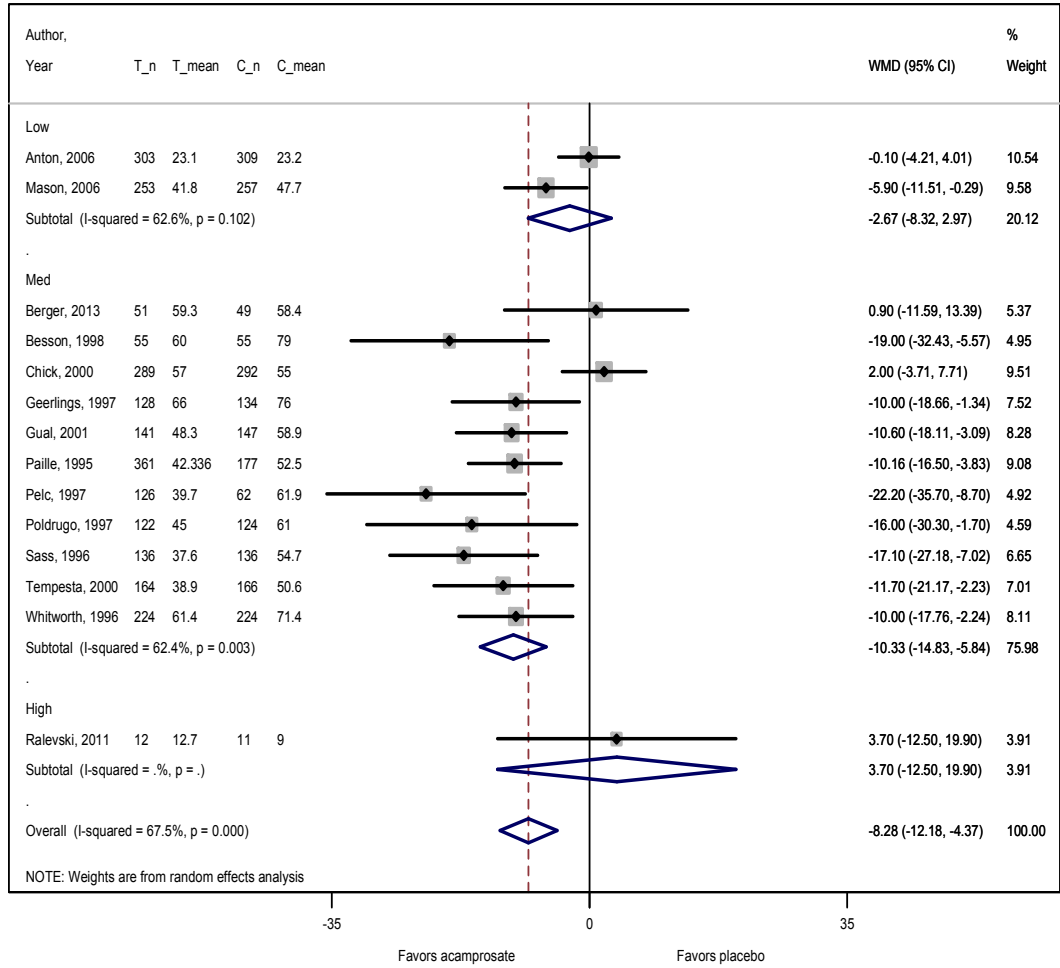
eFigure 3. Return to any drinking for acamprosate compared with placebo, subgroup analysis by risk of bias

Acamprosate versus Placebo Return to Any Drinking by Risk of Bias

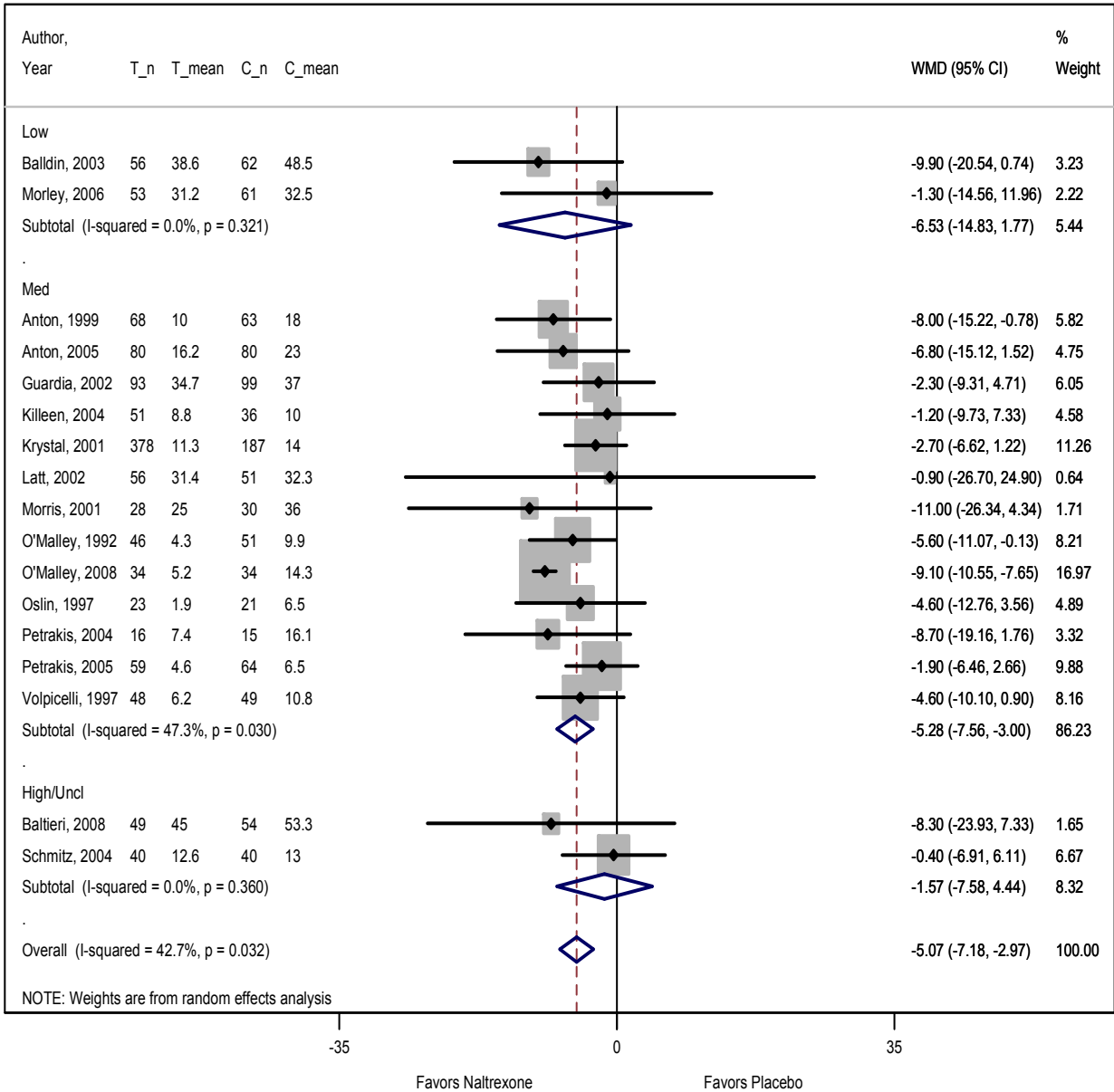


eFigure 7. Percent drinking days for acamprosate compared with placebo, subgroup analysis by risk of bias

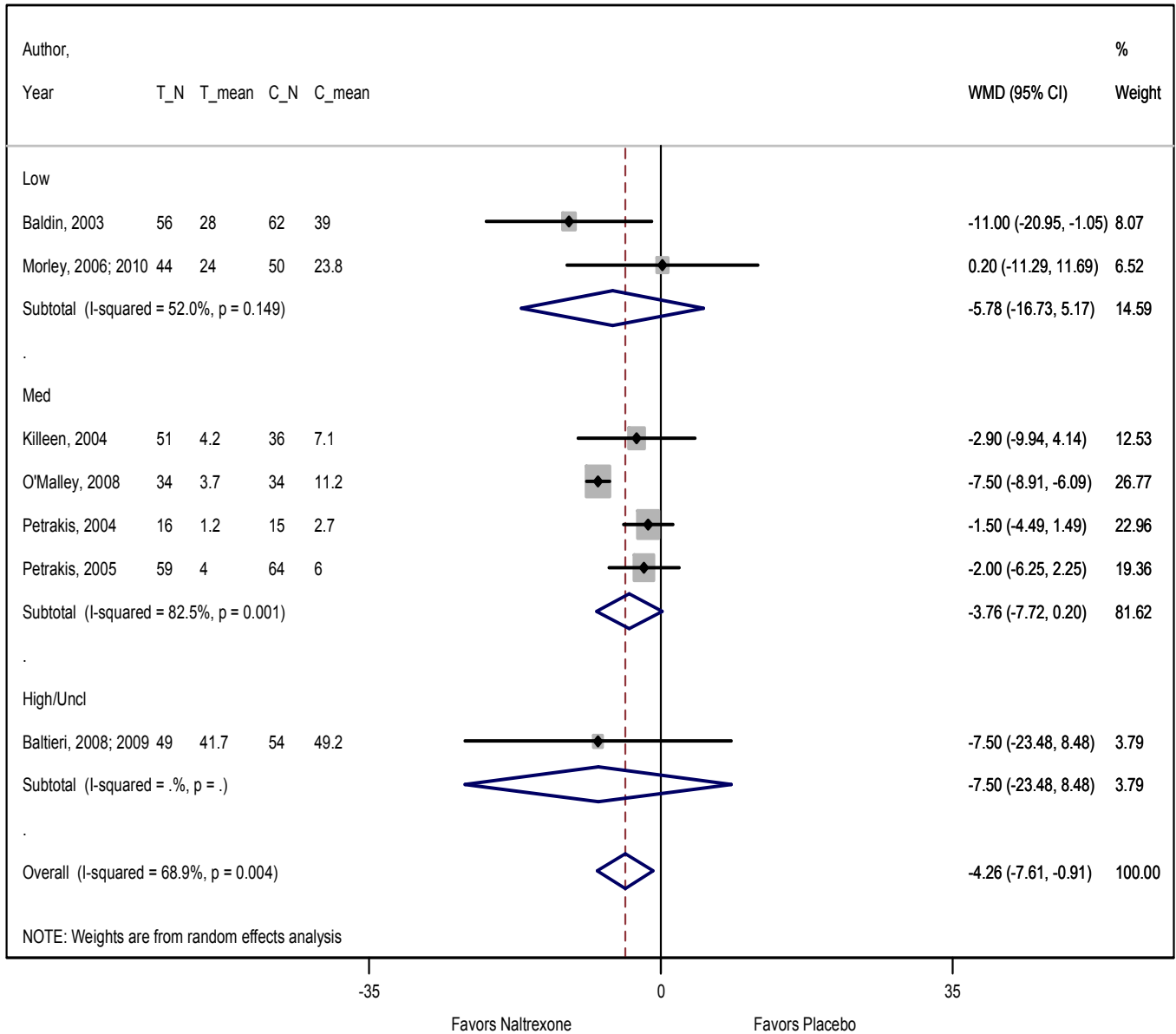
Acamprosate versus Placebo - Percent Drinking Days by Risk of Bias



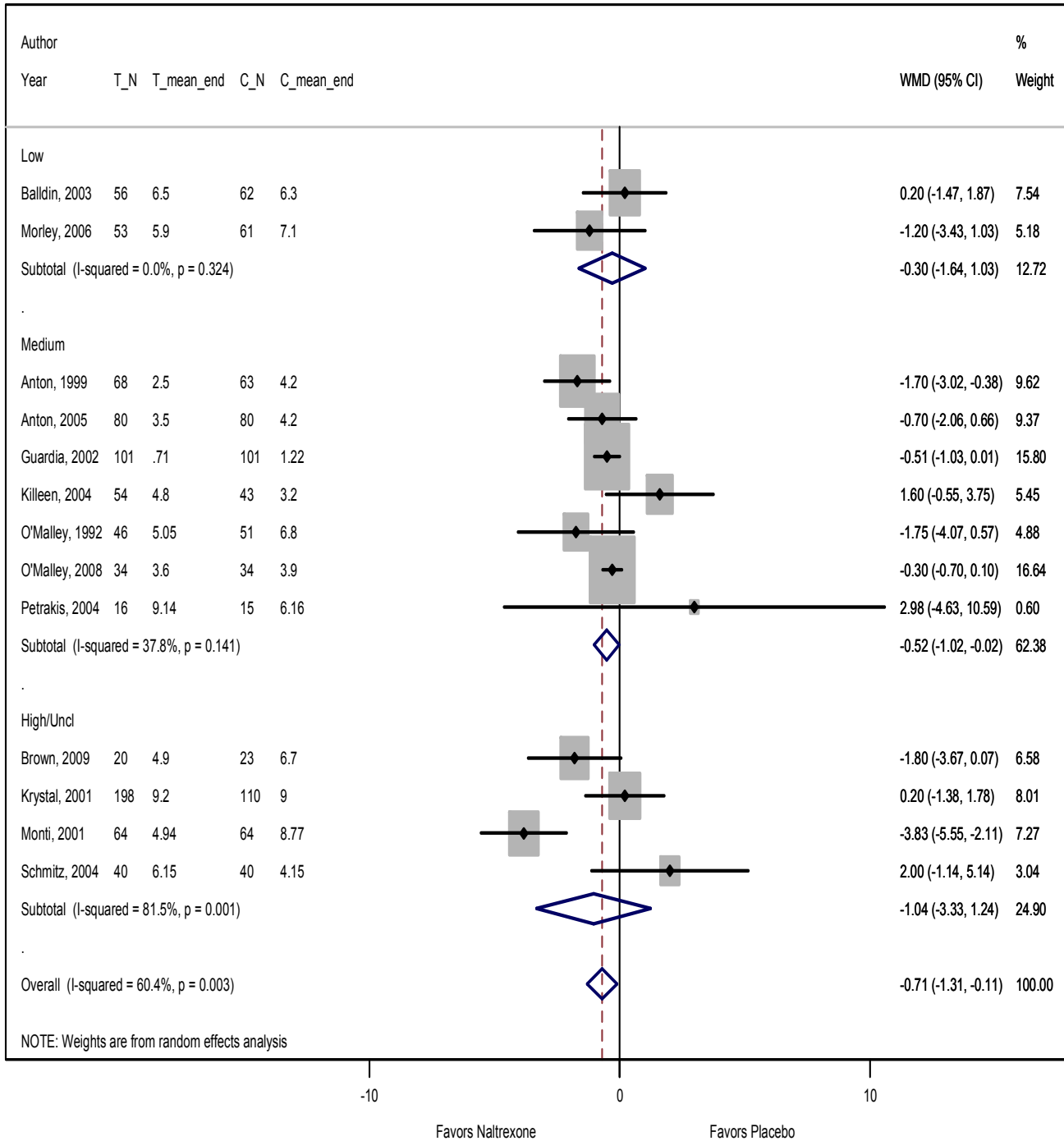
eFigure 8. Percent drinking days for oral naltrexone (50mg/day) compared with placebo, subgroup analysis by risk of bias



eFigure 9. Percent heavy drinking days for oral naltrexone (50mg/day) compared with placebo, subgroup analysis by risk of bias



eFigure 10. Drinks per drinking day for oral naltrexone (50mg/day) compared with placebo, subgroup analysis by risk of bias



eTable 5. Summary of findings and strength of evidence for alcohol consumption outcomes and health outcomes for medications used off-label or those under investigation

| Intervention | Outcome | N studies ^a | N participants | Results Effect Size (95% CI) ^b | NNT (95% CI) ^c | Strength of Evidence |
|---------------|--------------------------|------------------------|----------------|--|---------------------------|----------------------|
| Amitriptyline | Return to any drinking | 0 | 0 | NA | NA | Insufficient |
| | Return to heavy drinking | 0 | 0 | NA | NA | Insufficient |
| | % DD | 0 | 0 | NA | NA | Insufficient |
| | % HDD | 0 | 0 | NA | NA | Insufficient |
| | Drinks per DD | 0 | 0 | NA | NA | Insufficient |
| | Accidents or injuries | 0 | 0 | NA | NA | Insufficient |
| | QoL or function | 0 | 0 | NA | NA | Insufficient |
| | Mortality | 0 | 0 | NA | NA | Insufficient |
| Aripiprazole | Return to any drinking | 1 ²⁶ | 288 | 89% (ARI) vs. 78% (PBO); P=0.02 | NA | Insufficient |
| | Return to heavy drinking | 1 ²⁶ | 288 | 73% (ARI) vs. 73% (PBO); P=0.98 | NA | Insufficient |
| | % DD | 1 ²⁶ | 288 | 41% (ARI) vs. 37% (PBO); P=0.23 | NA | Insufficient |
| | % HDD | 0 | 0 | NA | NA | Insufficient |
| | Drinks per DD | 1 ²⁶ | 288 | 4.4 (ARI) vs. 5.5 (PBO); P<0.001 | NA | Insufficient |
| | Accidents or injuries | 0 | 0 | NA | NA | Insufficient |
| | QoL or function | 0 | 0 | NA | NA | Insufficient |
| | Mortality | 0 | 0 | NA | NA | Insufficient |
| Atomoxetine | Return to any drinking | 0 | 0 | NA | NA | Insufficient |

| | | | | | | |
|-----------|--------------------------|--------------------|-----|---|----|--------------|
| | Return to heavy drinking | 0 | 0 | NA | NA | Insufficient |
| | % DD | 0 | 0 | NA | NA | Insufficient |
| | % HDD | 0 | 0 | NA | NA | Insufficient |
| | Drinks per DD | 0 | 0 | NA | NA | Insufficient |
| | Accidents or injuries | 0 | 0 | NA | NA | Insufficient |
| | QoL or function | 0 | 0 | NA | NA | Insufficient |
| | Mortality | 0 | 0 | NA | NA | Insufficient |
| Baclofen | Return to any drinking | 2 ^{14,58} | 164 | Study 1: OR 6.3, 95% CI 2.4, 16.1 Study 2: NSD | NA | Insufficient |
| | Return to heavy drinking | 2 ^{14,58} | 164 | Study 1: BAC significantly lower than PBO (data in Figure, P=0.006) Study 2: HR 0.92, P=0.76 | NA | Insufficient |
| | % DD | 1 ⁵⁸ | 80 | 50.1% (BAC) vs. 49.4% (PBO); P=0.50 | NA | Insufficient |
| | % HDD | 1 ⁵⁸ | 80 | 25.9% (BAC) vs. 25.5% (PBO); P=0.73 | NA | Insufficient |
| | Drinks per DD | 0 | 0 | NA | NA | Insufficient |
| | Accidents or injuries | 0 | 0 | NA | NA | Insufficient |
| | QoL or function | 0 | 0 | NA | NA | Insufficient |
| | Mortality | 0 | 0 | NA | NA | Insufficient |
| Buspirone | Return to any drinking | 1 | 54 | RD: 0.07 (-0.19 to 0.34) | NA | Insufficient |
| | Return to heavy drinking | 0 | 0 | NA | NA | Insufficient |
| | % DD | 2 ^{49,83} | 161 | WMD: -3.4 (-9.2 to 2.4) | NA | Low |
| | % HDD | 0 | 0 | NA | NA | Insufficient |

| | | | | | | |
|-------------|--------------------------|--------------------|-----|---------------------------|----|--------------|
| | Drinks per DD | 1 ⁸³ | 61 | 0.7 vs. 2.1; P NSD | NA | Insufficient |
| | Accidents or injuries | 0 | 0 | NA | NA | Insufficient |
| | QoL or function | 0 | 0 | NA | NA | Insufficient |
| | Mortality | 0 | 0 | NA | NA | Insufficient |
| Citalopram | Return to any drinking | 0 | 0 | NA | NA | Insufficient |
| | Return to heavy drinking | 0 | 0 | NA | NA | Insufficient |
| | % DD | 0 | 0 | NA | NA | Insufficient |
| | % HDD | 0 | 0 | NA | NA | Insufficient |
| | Drinks per DD | 0 | 0 | NA | NA | Insufficient |
| | Accidents or injuries | 0 | 0 | NA | NA | Insufficient |
| | QoL or function | 0 | 0 | NA | NA | Insufficient |
| | Mortality | 0 | 0 | NA | NA | Insufficient |
| Desipramine | Return to any drinking | 0 | 0 | NA | NA | Insufficient |
| | Return to heavy drinking | 0 | 0 | NA | NA | Insufficient |
| | % DD | 0 | 0 | NA | NA | Insufficient |
| | % HDD | 0 | 0 | NA | NA | Insufficient |
| | Drinks per DD | 0 | 0 | NA | NA | Insufficient |
| | Accidents or injuries | 0 | 0 | NA | NA | Insufficient |
| | QoL or function | 0 | 0 | NA | NA | Insufficient |
| | Mortality | 0 | 0 | NA | NA | Insufficient |
| Fluoxetine | Return to any drinking | 1 ⁴³ | 51 | RD: -0.13 (-0.35 to 0.10) | NA | Insufficient |
| | Return to heavy drinking | 0 | 0 | NA | NA | Insufficient |
| | % DD | 2 ^{43,84} | 146 | WMD: -3.2 (-18.2 to 11.9) | NA | Low |
| | % HDD | 1 ⁴³ | 51 | 4.8 (FLUOX) vs. 16 | NA | Insufficient |

| | | | | | | |
|-------------|--------------------------|--------------------|-----|---|----|--------------|
| | | | | (PBO); P=0.04 | | |
| | Drinks per DD | 2 ^{43,84} | 146 | WMD: -1.2 (-4.6 to 2.2) | NA | Low |
| | Accidents or injuries | 0 | 0 | NA | NA | Insufficient |
| | QoL or function | 0 | 0 | NA | NA | Insufficient |
| | Mortality | 0 | 0 | NA | NA | Insufficient |
| Fluvoxamine | Return to any drinking | 1 ⁴² | 492 | 12 weeks: 58% (FLUV) vs. 54% (PBO); P=0.40 52 weeks: 71% (FLUV) vs. 71% (PBO); P=0.94 | NA | Insufficient |
| | Return to heavy drinking | 1 ⁴² | 492 | 12 weeks: 46% (FLUV) vs. 40% (PBO); P=0.18 52 weeks: 64% (FLUV) vs. 64% (PBO); P=0.47 | NA | Insufficient |
| | % DD | 1 ⁴² | 492 | 12 weeks: 31% (FLUV) vs. 23% (PBO); P=0.009 52 weeks: 44% (FLUV) vs. 38% (PBO); P=0.13 | NA | Insufficient |
| | % HDD | 0 | 0 | NA | NA | Insufficient |
| | Drinks per DD | 0 | 0 | NA | NA | Insufficient |
| | Accidents or injuries | 0 | 0 | NA | NA | Insufficient |
| | QoL or function | 0 | 0 | NA | NA | Insufficient |
| | Mortality | 1 ⁴² | 492 | 52 weeks: 1 (FLUV) vs. 1 (PBO) | NA | Insufficient |
| Gabapentin | Return to any drinking | 1 ¹⁰⁹ | 150 | 89% (78% to 95%) (GAB 900) vs. 83% (70% to 91%) (GAB 1800) vs. 96% (86% to 99%) (PBO) | NA | Insufficient |

| | | | | | | |
|------------|--------------------------|---------------------|-----|---|----|--------------|
| | Return to heavy drinking | 1 ¹⁰⁹ | 150 | 70% (57% to 81%) (GAB 900) vs. 55% (41% to 69%) (GAB 1800) vs. 78% (63% to 86%) (PBO) | NA | Insufficient |
| | % DD | 0 | 0 | NA | NA | Insufficient |
| | % HDD | 0 | 0 | NA | NA | Insufficient |
| | Drinks per DD | 0 | 0 | NA | NA | Insufficient |
| | Accidents or injuries | 0 | 0 | NA | NA | Insufficient |
| | QoL or function | 0 | 0 | NA | NA | Insufficient |
| | Mortality | 1 ¹⁰⁹ | 150 | No deaths in any group | NA | Insufficient |
| Imipramine | Return to any drinking | 1 ¹¹⁰ | 56 | 69% (IMI) vs. 79% (PBO); P NSD | NA | Insufficient |
| | Return to heavy drinking | 0 | 0 | NA | NA | Insufficient |
| | % DD | 1 ¹¹⁰ | 56 | 28.3% (IMI) vs. 30.8% (PBO); P NSD | NA | Insufficient |
| | % HDD | 1 ¹¹⁰ | 56 | 13.5% (IMI) vs. 9.0% (PBO); P NSD | NA | Insufficient |
| | Drinks per DD | 1 ¹¹⁰ | 56 | 3.7 (IMI) vs. 4.1 (PBO); P NS | NA | Insufficient |
| | Accidents or injuries | 0 | 0 | NA | NA | Insufficient |
| | QoL or function | 0 | 0 | NA | NA | Insufficient |
| | Mortality | 0 | 0 | NA | NA | Insufficient |
| Nalmefene | Return to any drinking | 0 | 0 | NA | NA | Insufficient |
| | Return to heavy drinking | 1 ¹⁰⁷ | 105 | RD: -0.23 (-0.43 to -0.03) | NA | Insufficient |
| | % DD | 2 ^{78,107} | 508 | WMD: -1.1 (-7.6 to 5.4) | NA | Low |
| | % HDD | 1 ⁷⁸ | 403 | 18.1% (NALM) vs. 29.7% (PBO); P=0.02 | NA | Insufficient |
| | Change in HDDs per | | | | | |

| | | | | | | |
|-------------|---------------------------------------|--|--------------|--|----|--------------|
| | month: OC analysis PMI analysis | 2 ^{64,103} 2 ^{64,103} | 806 1,234 | WMD: -2.0 (-3.0 to -1.0) WMD: -1.3 (-2.2 to -0.3) | | Moderate |
| | Drinks per DD | 3 ^{21,78,107} | 608 | WMD: -1.02 (-1.77 to -0.28) | NA | Moderate |
| | Accidents or injuries | 0 | 0 | NA | NA | Insufficient |
| | QoL or function | 0 | 0 | NA | NA | Insufficient |
| | Mortality | 2 ^{64,103} | 1,253 | Study 1: 0 (NALM) vs. 2 (PBO) Study 2: 1 (NALM) vs. 1 (PBO) | | |
| Olanzapine | Return to any drinking | 0 | 0 | NA | NA | Insufficient |
| | Return to heavy drinking | 1 ⁶⁶ | 60 | 37.9% (OLA) vs. 29.0% (PBO); P=0.50 | NA | Insufficient |
| | % DD | 1 ⁶⁶ | 60 | 13.1% (OLA) vs. 22.7% (PBO); P=0.18 | NA | Insufficient |
| | % HDD | 0 | 0 | NA | NA | Insufficient |
| | Drinks per DD | 1 ⁶⁶ | 60 | 1.8 (OLA) vs. 2.0 (PBO); P=0.71 | NA | Insufficient |
| | Accidents or injuries | 0 | 0 | NA | NA | Insufficient |
| | QoL or function | 0 | 0 | NA | NA | Insufficient |
| | Mortality | 0 | 0 | NA | NA | Insufficient |
| Ondansetron | Return to any drinking | 0 | 0 | NA | NA | Insufficient |
| | Return to heavy drinking | 0 | 0 | NA | NA | Insufficient |
| | % DD | 0 | 0 | NA | NA | Insufficient |
| | % HDD | 0 | 0 | NA | NA | Insufficient |
| | Drinks per DD | 0 | 0 | NA | NA | Insufficient |

| | | | | | | |
|------------|--------------------------|------------------------|-----|------------------------------------|----|--------------|
| | Accidents or injuries | 0 | 0 | NA | NA | Insufficient |
| | QoL or function | 0 | 0 | NA | NA | Insufficient |
| | Mortality | 0 | 0 | NA | NA | Insufficient |
| Paroxetine | Return to any drinking | 0 | 0 | NA | NA | Insufficient |
| | Return to heavy drinking | 0 | 0 | NA | NA | Insufficient |
| | % DD | 1 ³⁴ | 42 | 34% (PAR) vs. 35% (PBO); P=NSD | NA | Insufficient |
| | % HDD | 1 ³⁴ | 42 | 54% (PAR) vs. 55% (PBO); P=NSD | NA | Insufficient |
| | Drinks per DD | 1 ³⁴ | 42 | 5.9 (PAR) vs. 7.0 (PBO); P=NSD | NA | Insufficient |
| | Accidents or injuries | 0 | 0 | NA | NA | Insufficient |
| | QoL or function | 0 | 0 | NA | NA | Insufficient |
| | Mortality | 0 | 0 | NA | NA | Insufficient |
| Quetiapine | Return to any drinking | 0 | 0 | NA | NA | Insufficient |
| | Return to heavy drinking | 0 | 0 | NA | NA | Insufficient |
| | % DD | 0 | 0 | NA | NA | Insufficient |
| | % HDD | 0 | 0 | NA | NA | Insufficient |
| | Drinks per DD | 0 | 0 | NA | NA | Insufficient |
| | Accidents or injuries | 0 | 0 | NA | NA | Insufficient |
| | QoL or function | 0 | 0 | NA | NA | Insufficient |
| | Mortality | 0 | 0 | NA | NA | Insufficient |
| Sertraline | Return to any drinking | 1 ¹⁴² | 79 | 72.5% (SER) vs. 76.9% (PBO); P NSD | NA | Insufficient |
| | Return to heavy drinking | 2 ^{46,63} | 142 | RD: -0.04 (-0.31 to 0.23) | NA | Low |
| | % DD | 3 ^{63,87,111} | 299 | WMD: 1.8 (-6.25 to 9.86) | NA | Low |

| | | | | | | |
|---------------|--------------------------|-----------------------|-----|--|------------|----------------------|
| | % HDD | 2 ^{37,87} | 228 | WMD: 1.9 (0.70 to 3.0) | NA | Low (favors placebo) |
| | Drinks per DD | 2 ^{37,111} | 176 | WMD: -0.86 (-2.2 to 0.48) | NA | Low |
| | Accidents or injuries | 0 | 0 | NA | NA | Insufficient |
| | QoL or function | 1 ⁶³ | 83 | Graph only; data NR (P=0.03) | NA | Insufficient |
| | Mortality | 1 ¹⁴² | 79 | 0 (SER) vs. 0 (PBO) | NA | Insufficient |
| Topiramate | Return to any drinking | 0 | 0 | NA | NA | Insufficient |
| | Return to heavy drinking | 0 | 0 | NA | NA | Insufficient |
| | % DD | 2 ^{73,77} | 541 | WMD: -6.5 (-12.0 to -1.0) | NA | Moderate |
| | % HDD | 3 ^{69,73,77} | 691 | WMD: -9.0 (-15.3 to -2.7) | NA | Moderate |
| | Drinks per DD | 3 ^{69,73,77} | 691 | WMD: -1.0 (-1.6 to -0.5) | NA | Moderate |
| | Accidents or injuries | 1 ⁷³ | 371 | 4.4% (TOP) vs. 11.7% (PBO); P=0.01 | NA | Insufficient |
| | QoL or function | 1 ⁹¹ | 106 | SF-36 physical: 89.9 (TOP) vs. 89.4 (PBO); P=0.85 SF-36 mental: 84.0 (TOP) vs. 84.2 (PBO); P=0.92 | NA | Insufficient |
| | Mortality | 1 ⁷³ | 371 | 0 (TOP) vs. 1 (PBO) | NA | Insufficient |
| Valproic acid | Return to any drinking | 1 ³⁶ | 29 | 81% (VAL) vs. 83% (PBO); P NSD | NA | Insufficient |
| | Return to heavy drinking | 2 ^{36,149} | 81 | RD: -0.32 (-0.52 to -0.11) | 4 (2 to 9) | Low |
| | % DD | 1 ³⁶ | 29 | 15.9 (VAL) vs. 19.6 (PBO); P NSD | NA | Insufficient |
| | % HDD | 2 ^{36,149} | 81 | WMD: -8.5 (-15.9 to - | NA | Low |

| | | | | | | |
|-------------|--------------------------|---------------------|-----|--|----|--------------|
| | | | | 1.1) | | |
| | Drinks per DD | 2 ^{36,149} | 81 | WMD: -2.6 (-5.0 to -0.15) | NA | Low |
| | Accidents or injuries | 0 | 0 | NA | NA | Insufficient |
| | QoL or function | 0 | 0 | NA | NA | Insufficient |
| | Mortality | 0 | 0 | NA | NA | Insufficient |
| Varenicline | Return to any drinking | 1 ⁹⁸ | 200 | 98% (VAR) vs. 98% (PBO); P=0.81 | NA | Insufficient |
| | Return to heavy drinking | 1 ⁹⁸ | 200 | 93% (VAR) vs. 95% (PBO); P=0.50 | NA | Insufficient |
| | % DD | 1 ⁹⁸ | 200 | 60% (VAR) vs. 64% (PBO); P=0.29 | NA | Insufficient |
| | % HDD | 1 ⁹⁸ | 200 | 38% (VAR) vs. 48% (PBO); P=0.03 | NA | Insufficient |
| | Drinks per DD | 1 ⁹⁸ | 200 | 5.8 (VAR) vs. 6.8 (PBO); P=0.03 | NA | Insufficient |
| | Accidents or injuries | 0 | 0 | NA | NA | Insufficient |
| | QoL or function | 1 ⁹⁸ | 200 | SF-12 mental mean difference=0.7; P=0.55 SF-12 physical mean difference=0.4; P=0.38 | NA | Insufficient |
| | Mortality | 1 ⁹⁸ | 200 | 1 (VAR) vs. 0 (PBO) | NA | Insufficient |

^a Includes only studies rated as low or medium risk of bias included in the main analyses; these numbers do not include studies rated as high or unclear risk of bias that were included in sensitivity analyses.

^b Negative effect sizes favor intervention over placebo/control.

^c NA entry for numbers needed to treat (NNT) indicates that the risk difference (95% CI) was not statistically significant, so we did not calculate a NNT, or that the effect measure was not one that allows direct calculation of NNT (e.g., WMD).

Abbreviations: ARI = aripiprazole; BAC = baclofen; CI = confidence interval; DD, drinking day; FLUOX = fluoxetine; FLUV = fluvoxamine; HDD, heavy drinking day; HR = hazard ratio; IMI = imipramine; N = number; NA = not applicable; NALM = nalmefene; NNT = number needed to treat; NR = not reported; NSD = no statistically significant difference; NTX = naltrexone; OC = observed cases; OLA = olanzapine; OR = odds ratio; PAR = paroxetine; PBO = placebo; PMI = placebo bean imputation; QoL, quality of life; RD = risk difference; SER = sertraline; TOP = topiramate; VAL = valproate / valproic acid; VAR = varenicline; vs. = versus; WMD = weighted mean difference.

eTable 6. Summary of meta-analyses of adverse events from head-to-head trials comparing acamprosate with naltrexone

| Outcome | N trials ^a | N participants | RD ^b | 95% CI | Heterogeneity I ² |
|-------------------------------------|----------------------------------|----------------|-----------------|----------------|------------------------------|
| Withdrawal due to adverse events | 2 ^{23,102} | 953 | 0.0 | -0.04 to 0.07 | 73.4% |
| Withdrawal due to adverse events—SA | 3 ^{23,102,147} | 1,110 | 0.0 | -0.04 to 0.04 | 64.5% |
| Diarrhea | 4 ^{20,23,79,117} | 836 | 0.1 | -0.02 to 0.37 | 89.3% |
| Diarrhea—SA | 5 ^{20,23,79,117,147} | 993 | 0.1 | -0.07 to 0.35 | 96.0% |
| Dizziness | 2 ^{20,117} | 144 | 0.0 | -0.23 to 0.39 | 81.7% |
| Dizziness—SA | 3 ^{20,90,117} | 306 | 0.0 | -0.19 to 0.13 | 82.4% |
| Headache | 3 ^{20,117,147} | 301 | - | -0.12 to 0.01 | 0.0% |
| Headache—SA | 4 ^{20,90,117,147} | 463 | 0.0 | -0.16 to -0.01 | 36.3% |
| Insomnia | 2 ^{20,117} | 144 | 0.0 | -0.20 to 0.34 | 65.5% |
| Nausea | 4 ^{20,23,79,117} | 836 | - | -0.18 to 0.02 | 72.8% |
| Nausea—SA | 6 ^{20,23,79,90,117,147} | 1,155 | 0.0 | -0.18 to -0.02 | 71.6% |
| Vomiting | 2 ^{20,23} | 648 | - | -0.11 to -0.01 | 0.0% |

^a Our main analyses for each outcome (i.e., the first row for each outcome) included studies rated as low or medium risk of bias. We did not include studies rated as high or unclear risk of bias in our main analyses, but included them in sensitivity analyses (i.e., rows labeled SA).

^b Positive risk differences favor naltrexone. Table only includes rows for outcomes with sufficient data for meta-analyses.

Abbreviations: CI = confidence interval; N = number of trials or participants contributing data; NA, not applicable; RD = risk difference; SA = sensitivity analysis.

eTable 7. Summary of meta-analyses of adverse events from placebo-controlled trials of acamprosate, naltrexone, nalmefene, topiramate, and valproic acid^a

| Outcome | N trials ^b | N participants | RD ^c | 95% CI | NNH (95% CI) | Heterogeneity I ² |
|-------------------------------------|---|----------------|-----------------|---------------|-------------------|------------------------------|
| Acamprosate | | | | | | |
| Withdrawal due to adverse events | 13 ^{23,32,33,40,60,62,102,108,129,144,150,155,162} | 4,653 | 0.0 | -0.00 to 0.02 | NA | 8.5% |
| Withdrawal due to adverse events—SA | 16 ^{23,32,33,40,60,62,94,95,102,108,129,132,144,150,155,162} | 5,480 | 0.0 | 0.00 to 0.01 | 125 (67 to 1,000) | 0.0% |
| Anxiety | 1 ¹⁰⁸ | 601 | 0.1 | 0.10 to 0.23 | NA | NA |
| Anxiety—SA | 2 ^{108,145} | 624 | 0.1 | 0.09 to 0.22 | 7 (5 to 11) | 0.0% |
| Diarrhea | 12 ^{20,23,29,32,33,60,79,117,129,150,155,162} | 2,978 | 0.1 | 0.03 to 0.17 | 11 (6 to 34) | 88.9% |
| Diarrhea—SA | 14 ^{20,23,29,32,33,60,79,95,117,129,132,150,155,162} | 3,797 | 0.0 | 0.04 to 0.15 | 11 (7 to 27) | 85.8% |
| Dizziness | 2 ^{20,117} | 151 | 0.0 | -0.22 to 0.38 | NA | 80.3% |
| Headache | 6 ^{20,29,117,144,150,155} | 1,074 | 0.0 | -0.05 to 0.05 | NA | 67.0% |
| Headache—SA | 7 ^{20,29,95,117,144,150,155} | 1,643 | 0.0 | -0.04 to 0.04 | NA | 60.2% |
| Insomnia | 3 ^{20,32,117} | 251 | 0.0 | -0.10 to 0.14 | NA | 73.5% |
| Insomnia—SA | 4 ^{20,32,95,117} | 820 | 0.0 | -0.06 to 0.09 | NA | 63.8% |
| Nausea | 7 ^{20,23,32,79,95,117,144} | 1,758 | 0.0 | -0.01 to 0.02 | NA | 0.3% |
| Nausea—SA | 8 ^{20,23,32,79,94,95,117,144} | 1,828 | 0.0 | -0.01 to 0.03 | NA | 12.5% |

| | | | | | | |
|---|--|----------------|----------------------|--------------------------------------|--|----------------|
| Numbness Numbness—SA | 1 ⁶⁰ 2 ^{60,95} | 262 831 | 0.0 1 0.0 1 | -0.01 to 0.03 -0.01 to 0.03 | NA NA | NA 0.0% |
| Rash Rash--SA | 1 ²⁰ 2 ^{20,94} | 35 105 | 0.1 1 0.0 7 | -0.07 to 0.29 -0.01 to 0.16 | NA NA | NA 0.0% |
| Suicide attempts or suicidal ideation Suicide attempts or suicidal ideation—SA | 1 ⁴⁰ 3 ^{40,95,145} | 581 1,173 | 0.0 1 0.0 0 | -0.00 to 0.02 -0.01 to 0.01 | NA NA | NA 14.6% |
| Vomiting Vomiting – SA | 4 ^{20,23,95,108} 5 ^{20,23,95,108,145} | 1,817 1,840 | 0.0 2 0.0 2 | 0.01 to 0.04 0.01 to 0.04 | 42 (24 to 143) 42 (25 to 167) | 0.0% 0.0% |
| Naltrexone | | | | | | |
| Withdrawal due to adverse events Withdrawal due to adverse events - SA | 17 ^{18,23,41,65,82,86,92,102,116,123,125,126,133,142,161,165} 20 ^{18,23,41,52,65,72,82,86,92,102,116,123,125,126,133,142,152,161,165} | 2,743 2,899 | 0.0 2 0.0 2 | 0.01 to 0.03 0.01 to 0.03 | 48 (30 to 112) 50 (32 to 125) | 0.0% 0.0% |
| Anorexia | 1 ⁴¹ | 175 | 0.0 8 | 0.01 to 0.14 | NA | NA |
| Anxiety Anxiety—SA | 7 ^{41,55,119,125,128,166,167} 9 ^{15,41,55,119,125,128,159,166,167} | 1,461 1,676 | 0.0 1 0.0 1 | -0.02 to 0.04 -0.02 to 0.04 | NA NA | 0.0% 0.0% |
| Cognitive dysfunction | 1 ¹³⁵ | 123 | 0.1 9 | 0.04 to 0.34 | NA | NA |
| Diarrhea Diarrhea - SA | 11 ^{17,20,23,41,55,59,79,92,117,125,133} 12 ^{17,20,23,30,41,55,59,79,92,117,125,133} | 2,358 2,461 | 0.0 1 | -0.01 to 0.04 | NA NA | 21.9% 30.2% |

| | | | | | | |
|---------------------|---|-------|----------|------------------|---------|-------|
| | | | 0.0 1 | -0.02 to 0.03 | | |
| Dizziness | 13 ^{17,20,41,55,59,82,86,89,117,119,123,125,126} | 2,675 | 0.0 | 0.04 to | 16 (12 | 37.4% |
| Dizziness - SA | 17 ^{15,17,20,30,41,55,59,72,82,86,89,93,117,119,123,125,126} | 2,977 | 6 | 0.09 | to 28) | 27.1% |
| | | | 0.0 | 0.04 to | 17 (13 | |
| Headache | 17 ^{17,20,41,55,59,65,82,85,89,92,117,119,125,127,128,133,141} | 3,347 | 6 | 0.08 | to 27) | |
| Headache - SA | 22 ^{15,17,20,41,55,59,65,67,72,82,85,89,92,117,119,125,127,128,133,141,152,159} | 3,799 | 1 | -0.02 to | NA | 8.0% |
| | | | 0.0 | 0.03 | NA | 20.9% |
| | | | 0 | -0.02 to | | |
| Insomnia | 8 ^{20,41,55,82,117,119,128,141} | 1,637 | 0.0 | -0.00 to | NA | 0.0% |
| Insomnia - SA | 12 ^{15,20,30,41,55,67,82,93,117,119,128,141} | 2,030 | 3 | 0.06 | 39 (20 | 0.0% |
| | | | 0.0 | 0.00 to | to | |
| | | | 3 | 0.05 | 1,000) | |
| Nausea | 24 ^{17,18,20,23,41,55,59,79,82,85,86,89,92,113,117,119,123,125-128,133,135,141} | 4,655 | 0.1 | 0.07 to | 9 (7 to | 69.6% |
| Nausea - SA | 31 ^{15,17,18,20,23,30,41,55,59,67,72,79,82,85,86,89,92,93,113,117,119,123,125-128,133,135,141,152,159} | 5,263 | 1 | 0.15 | 14) | 65.9% |
| | | | 0.1 | 0.07 to | 10 (8 | |
| | | | 0 | 0.13 | to 15) | |
| Numbness | 1 ¹³⁵ | 123 | - | -0.19 to | NA | NA |
| Numbness - SA | 2 ^{30,135} | 226 | 0.0 | 0.17 | NA | 0.0% |
| | | | 1 | -0.08 to | | |
| | | | - | 0.04 | | |
| | | | 0.0 | | | |
| | | | 2 | | | |
| Rash | 4 ^{17,20,125,133} | 469 | - | -0.06 to | NA | 41.6% |
| Rash - SA | 5 ^{17,20,93,125,133} | 522 | 0.0 | 0.04 | NA | 21.5% |
| | | | 1 | -0.06 to | | |
| | | | - | 0.02 | | |
| | | | 0.0 | | | |
| | | | 2 | | | |
| Taste abnormalities | 1 ¹³⁵ | 123 | - | -0.18 to | NA | NA |
| | | | 0.0 | 0.17 | | |
| | | | 1 | | | |
| Vision changes | 2 ^{133,135} | 133 | 0.0 | -0.17 to | NA | 46.3% |

| | | | | | | |
|---------------------------------------|--|-------|-----|------------|---------|-------|
| | | | 8 | 0.33 | | |
| Vomiting | 9 ^{17,20,23,41,55,59,128,135,141} | 2,438 | 0.0 | 0.02 to | 24 (17 | 0.0% |
| Vomiting - SA | 11 ^{17,20,23,41,55,59,72,128,135,141,159} | 2,567 | 4 | 0.06 | to 44) | 1.1% |
| | | | 0.0 | 0.02 to | 27 (19 | |
| | | | 4 | 0.06 | to 56) | |
| Nalmefene | | | | | | |
| Withdrawal due to adverse events | 5 ^{21,64,78,103,107} | 2,054 | 0.0 | 0.02 to | 12 (7 | 86.4% |
| Withdrawal due to adverse events – SA | 7 ^{21,64,78,103,105,107,153} | 2,750 | 8 | 0.15 | to 50) | 79.5% |
| | | | 0.0 | 0.04 to | 13 (9 | |
| | | | 8 | 0.12 | to 28) | |
| Cognitive dysfunction | 1 ²¹ | 265 | 0.0 | 0.01 to | NA | NA |
| Cognitive dysfunction – SA | 2 ^{21,153} | 830 | 5 | 0.09 | NA | 81.9% |
| | | | 0.0 | -0.03 to | | |
| | | | 3 | 0.08 | | |
| Diarrhea | 2 ^{78,153} | 1,081 | - | -0.06 to - | NA | 0.0% |
| | | | 0.0 | 0.01 | (more | |
| | | | 3 | | with | |
| | | | | | placeb | |
| | | | | | o) | |
| Dizziness | 4 ^{21,64,78,103} | 1,944 | 0.1 | 0.11 to | 7 (5 to | 58.4% |
| Dizziness – SA | 6 ^{21,64,78,103,105,153} | 2,630 | 6 | 0.21 | 10) | 53.5% |
| | | | 0.1 | 0.10 to | 7 (6 to | |
| | | | 4 | 0.18 | 10) | |
| Headache | 3 ^{64,103,107} | 1,401 | 0.0 | 0.01 to | 26 (15 | 0.0% |
| Headache – SA | 4 ^{64,103,107,153} | 2,066 | 4 | 0.07 | to | 0.0% |
| | | | 0.0 | 0.01 to | 143) | |
| | | | 4 | 0.07 | 25 (15 | |
| | | | | | to 77) | |
| Insomnia | 5 ^{21,64,78,103,107} | 2,049 | 0.1 | 0.06 to | 10 (8 | 47.3% |
| Insomnia – SA | 6 ^{21,64,78,103,107,153} | 2,714 | 0 | 0.14 | to 17) | 34.9% |
| | | | 0.0 | 0.06 to | 11 (9 | |
| | | | 9 | 0.12 | to 16) | |
| Nausea | 5 ^{21,64,78,103,107} | 2,049 | 0.1 | 0.10 to | 7 (5 to | 75.8% |
| Nausea – SA | 6 ^{21,64,78,103,107,153} | 2,714 | 6 | 0.22 | 11) | 69.8% |
| | | | 0.1 | 0.11 to | 7 (5 to | |
| | | | 6 | 0.21 | 9) | |

| | | | | | | |
|--|----------------------------|-------|---------------------------|--------------------------|----------------------------|-------|
| Suicide attempts or suicidal ideation | 2 ^{64,103} | 1,253 | - | -0.02 to | NA | 0.0% |
| Suicide attempts or suicidal ideation – SA | 3 ^{64,103,153} | 1,918 | 0.0 1 | 0.00 -0.01 to 0.01 | NA | 39.2% |
| Taste abnormalities | 1 ⁷⁸ | 403 | 0.0 4 | 0.01 to 0.07 | NA | NA |
| Vomiting | 3 ^{64,78,103} | 1,679 | 0.0 | 0.02 to | 17 (11 | 68.1% |
| Vomiting – SA | 4 ^{64,78,103,153} | 2,344 | 6 0.0 7 | 0.10 0.03 to 0.11 | to 48) 15 (10 to 30) | 76.2% |
| Topiramate | | | | | | |
| Withdrawal due to adverse events | 3 ^{69,73,77} | 691 | 0.0 5 | -0.08 to 0.17 | NA | 93.4% |
| Withdrawal due to adverse events - SA | 4 ^{69,73,77,148} | 767 | 0.0 5 | -0.05 to 0.14 | NA | 86.9% |
| Anorexia | 1 ⁷³ | 371 | 0.1 3 | 0.06 to 0.20 | NA | NA |
| Anxiety | 1 ⁷³ | 371 | - | -0.17 to | NA | NA |
| Anxiety – SA | 2 ^{73,96} | 477 | 0.0 8 - 0.0 4 | 0.01 -0.14 to 0.05 | NA | 69.8% |
| Cognitive dysfunction | 2 ^{69,73} | 521 | 0.0 | 0.01 to | 12 (7 | 38.5% |
| Cognitive dysfunction –SA | 3 ^{69,73,96} | 627 | 8 0.0 7 | 0.16 0.03 to 0.12 | to 84) 14 (9 to 39) | 0.0% |
| Diarrhea | 1 ⁷³ | 371 | 0.0 | -0.03 to | NA | NA |
| Diarrhea – SA | 2 ^{30,73} | 477 | 4 0.0 0 | 0.10 -0.07 to 0.08 | NA | 61.1% |
| Dizziness | 2 ^{69,73} | 521 | 0.1 | -0.01 to | NA | 65.0% |
| Dizziness – SA | 4 ^{30,69,73,96} | 733 | 0 | 0.22 | NA | 80.2% |

| | | | | | | |
|---|--|------------|--------------------------------|--------------------------------------|--------------------------------|----------------|
| | | | 0.0 5 | -0.03 to 0.13 | | |
| Headache Headache –SA | 2 ^{73,77} 3 ^{73,77,96} | 541 647 | - 0.0 2 0.0 0 | -0.16 to 0.12 -0.09 to 0.09 | NA NA | 83.3% 78.7% |
| Insomnia Insomnia – SA | 1 ⁷³ 2 ^{30,73} | 371 477 | 0.0 3 0.0 3 | -0.05 to 0.11 -0.03 to 0.10 | NA NA | NA 0.0% |
| Nausea Nausea – SA | 1 ⁷³ 2 ^{30,73} | 371 477 | - 0.0 6 - 0.0 2 | -0.13 to 0.01 -0.11 to 0.06 | NA NA | NA 62.0% |
| Numbness Numbness – SA | 3 ^{69,73,77} 5 ^{30,69,73,77,96} | 691 903 | 0.3 2 0.2 6 | 0.15 to 0.48 0.12 to 0.41 | 4 (3 to 7) 4 (3 to 9) | 86.5% 88.0% |
| Taste abnormalities Taste abnormalities – SA | 1 ⁷³ 2 ^{73,96} | 371 477 | 0.1 8 0.1 5 | 0.11 to 0.25 0.07 to 0.23 | NA 7 (5 to 15) | NA 39.7% |
| Valproic acid | | | | | | |
| Withdrawal due to adverse events | 1 ¹⁴⁹ | 52 | 0.0 4 | -0.07 to 0.14 | NA | NA |
| Diarrhea | 1 ¹⁴⁹ | 52 | 0.1 0 | -0.12 to 0.32 | NA | NA |
| Headache | 1 ¹⁴⁹ | 52 | 0.0 5 | -0.20 to 0.30 | NA | NA |
| Nausea | 1 ¹⁴⁹ | 52 | 0.2 5 | 0.05 to 0.46 | NA | NA |
| Vision changes | 1 ¹⁴⁹ | 52 | - 0.0 2 | -0.26 to 0.22 | NA | NA |

^a Table includes rows for outcomes/comparisons for which there was at least one study rated low or medium risk of bias.

^b Our main analyses for each outcome (i.e., the first row for each outcome) included studies rated as low or medium risk of bias. We did not include studies rated as high or unclear risk of bias in our main analyses, but included them in sensitivity analyses (i.e., rows labeled SA).

^cPositive risk differences favor placebo. Sensitivity analyses include studies rated as high risk of bias.

Abbreviations: CI = confidence interval; N = number of trials or participants contributing data; NA = not applicable; NNH = number needed to harm; RD = risk difference; SA = sensitivity analysis.

eDiscussion. Applicability to Primary Care Settings

This supplement summarizes additional information about four studies that did not meet our inclusion criteria (due to study design or comparators) that have important implications for primary care settings.¹⁶⁸⁻¹⁷¹ While these studies found conflicting results, they demonstrate approaches to managing AUDs in primary care. In general, the interventions involve formal clinic structure, staffing, and protocols. They used variations of chronic care management, multidisciplinary team-based care, and care-coordination between primary care and mental health providers.

First, O'Malley et al.¹⁶⁹ found no difference in avoiding persistent heavy drinking between those who received naltrexone plus "primary care management" (PCM) and those who received naltrexone plus cognitive behavioral therapy. Among responders enrolled in a maintenance trial, those who received naltrexone and PCM had significantly better response (no more than 2 days of heavy drinking in final 28 days) than those who received placebo and PCM (80.8% vs. 51.9%, $p=0.03$). Primary care management was provided by nurse practitioners, physician assistants, and one internist in an initial 45-minute visit, followed by seven 15- to 20-minute sessions in the following 10 weeks.

Second, Kiritze-Topor et al.¹⁶⁸ conducted a trial with 149 general practitioners in France, randomizing patients ($N=422$) to acamprosate plus standard care or standard care alone. The trial reported better outcomes for the acamprosate group for alcohol-related health, personal, and social problems, and for quality of life.

Third, in a U.S.-based RCT ($N=163$), Oslin et al.¹⁷⁰ compared a primary-care based alcohol care management (ACM) program with a specialty outpatient addiction treatment program. The trial found that participants in the ACM program were more likely to receive naltrexone (65.9% vs. 11.5%), to be engaged in treatment (OR, 5.36; 95% CI, 2.99 to 9.59), and to have a lower percentage of heavy drinking days (OR, 2.16; 95% CI, 1.27 to 3.66) than participants in the specialty treatment program. Overall abstinence did not differ between groups.

Fourth, in the U.S.-based AHEAD trial ($N=563$), Saitz et al.¹⁷¹ compared chronic care management (CCM) with no CCM for people with alcohol or drug dependence who were not currently engaged in primary care. Of those enrolled, 12 percent had alcohol dependence without also meeting criteria for other drug dependence. CCM involved longitudinal care coordinated by a primary care clinician, and included motivational enhancement therapy; relapse prevention counseling; on-site medical, addiction, and psychiatric treatment; social work assistance; and referrals. The no-CCM group received a primary care appointment and a list of treatment resources including a telephone number to arrange counseling. The trial found no difference between groups for the primary outcome of abstinence over 12 months.

REFERENCES

1. American Psychiatric Association. *Diagnostic and statistical manual of mental disorders (5th ed.)*. Arlington, VA: American Psychiatric Publishing; 2013.
2. Isaac M, Janca A, Sartorius N. *ICD-10 symptom glossary for mental disorders*. Geneva: Division of Mental Health, World Health Organization; 1994.
3. Janca A, Ustun TB, van Drimmelen J, Dittmann V, Isaac M. *ICD-10 symptom checklist for mental disorders, version 1.1*. Geneva: Division of Mental Health, World Health Organization; 1994.
4. American Psychiatric Association. *Diagnostic and statistical manual of mental disorders, 4th ed. Text rev.* Washington, DC: American Psychiatric Publishing, Inc.; 2000.
5. Barrias JA, Chabac S, Ferreira L, Fonte A, Potgieter A, Teixeira de Sousa E. Acamprosate: multicenter Portuguese efficacy and tolerance evaluation study. *Psiquiatria Clinica*. 1997;18:149-160.
6. Huang X, Huang X, Peng H, Mai G. Placebo-controlled trial of naltrexone in outpatient treatment of alcohol dependence. *Chinese Mental Health Journal*. 2002;16(5):302-303, 295.
7. Krupitski EM, Burakov AM, Ivanov VB, et al. [The use of baclofen for treating affective disorders in alcoholism]. *Zhurnal nevrologii i psikiatrii imeni S.S. Korsakova / Ministerstvo zdravookhraneniia i meditsinsko? promyshlennosti Rossi?sko? Federatsii, Vserossi?skoe obshchestvo nevrologov [i] Vserossi?skoe obshchestvo psikiatrov*. 1994;94(1):57-61.
8. Ladewig D, Knecht T, Leher P, Fendl A. [Acamprosate--a stabilizing factor in long-term withdrawal of alcoholic patients]. *Therapeutische Umschau. Revue thérapeutique*. 1993;50(3):182-188.
9. Castro LA, Laranjeira R. [A double blind, randomized and placebo-controlled clinical trial with naltrexone and brief intervention in outpatient treatment of alcohol dependence]. *J. Bras. Psiquiatr*. 2009;58(2):79-85.
10. Roussaux JP, Hers D, Ferauge M. Does acamprosate influence alcohol consumption of weaned alcoholics? *J. Pharm. Belg*. 1996;51(2):65-68.
11. Geerlings P, Ansoms C, Van DBW. Acamprosate and relapse prevention in outpatient alcoholics; results from a randomized, placebo-controlled double-blind study in the Benelux. *Tijdschrift Voor Alcohol, Drugs En Andere Psychotrope Stoffen*. 1995;21(3):129-141.
12. Kiefer F, Jahn H, Holzbach R, Briken P, Stracke R, Wiedemann K. The NALCAM-study: Efficacy, tolerability, outcome. *Sucht*. 2003;49(6):342-351.
13. Sass H, Mann K, Soyka M. Drug support for prevention of relapse in alcoholic patients with acamprosate: Results of a double blind, randomized, placebo controlled study. *Sucht*. 1996;42(5):316-322.
14. Addolorato G, Leggio L, Ferrulli A, et al. Effectiveness and safety of baclofen for maintenance of alcohol abstinence in alcohol-dependent patients with liver cirrhosis: randomised, double-blind controlled study. *Lancet*. Dec 8 2007;370(9603):1915-1922.
15. Ahmadi J, Ahmadi N. A double blind, placebo-controlled study of naltrexone in the treatment of alcohol dependence. *German Journal of Psychiatry*. 2002;5(4):85-89.
16. Ahmadi J, Babaebeigi M, Maany I, et al. Naltrexone for alcohol-dependent patients. *Ir. J. Med. Sci*. Jan-Mar 2004;173(1):34-37.

17. ALK21-014: Efficacy and safety of Medisorb naltrexone (Vivitrol) after enforced abstinence; 2011. <http://www.clinicaltrials.gov/ct2/show/NCT00501631>. Accessed April 22, 2014
18. Anton RF, Moak DH, Waid LR, Latham PK, Malcolm RJ, Dias JK. Naltrexone and cognitive behavioral therapy for the treatment of outpatient alcoholics: results of a placebo-controlled trial. *A. J. Psychiatry*. Nov 1999;156(11):1758-1764.
19. Anton RF, Moak DH, Latham PK, et al. Posttreatment results of combining naltrexone with cognitive-behavior therapy for the treatment of alcoholism. *J. Clin. Psychopharmacol*. Feb 2001;21(1):72-77.
20. Anton RF. Testing combined pharmacotherapies and behavioral interventions for alcohol dependence (the COMBINE study): A pilot feasibility study. *Alcohol*. 2003;27(7):1123-1131.
21. Anton RF, Pettinati H, Zweben A, et al. A multi-site dose ranging study of nalmefene in the treatment of alcohol dependence. *J. Clin. Psychopharmacol*. Aug 2004;24(4):421-428.
22. Anton RF, Moak DH, Latham P, et al. Naltrexone combined with either cognitive behavioral or motivational enhancement therapy for alcohol dependence. *J. Clin. Psychopharmacol*. Aug 2005;25(4):349-357.
23. Anton RF, O'Malley SS, Ciraulo DA, et al. Combined pharmacotherapies and behavioral interventions for alcohol dependence: the COMBINE study: a randomized controlled trial. *JAMA*. May 3 2006;295(17):2003-2017.
24. Donovan DM, Anton RF, Miller WR, Longabaugh R, Hosking JD, Youngblood M. Combined pharmacotherapies and behavioral interventions for alcohol dependence (The COMBINE Study): examination of posttreatment drinking outcomes. *J Stud Alcohol Drugs*. Jan 2008;69(1):5-13.
25. LoCastro JS, Youngblood M, Cisler RA, et al. Alcohol treatment effects on secondary nondrinking outcomes and quality of life: the COMBINE study. *J Stud Alcohol Drugs*. Mar 2009;70(2):186-196.
26. Anton RF, Kranzler H, Breder C, Marcus RN, Carson WH, Han J. A randomized, multicenter, double-blind, placebo-controlled study of the efficacy and safety of aripiprazole for the treatment of alcohol dependence. *J. Clin. Psychopharmacol*. Feb 2008;28(1):5-12.
27. Anton RF, Myrick H, Wright TM, et al. Gabapentin combined with naltrexone for the treatment of alcohol dependence. *A. J. Psychiatry*. Jul 2011;168(7):709-717.
28. Balldin J, Berglund M, Borg S, et al. A 6-month controlled naltrexone study: combined effect with cognitive behavioral therapy in outpatient treatment of alcohol dependence. *Alcohol. Clin. Exp. Res*. Jul 2003;27(7):1142-1149.
29. Baltieri DA, De Andrade AG. Acamprosate in alcohol dependence: a randomized controlled efficacy study in a standard clinical setting. *J. Stud. Alcohol*. Jan 2004;65(1):136-139.
30. Baltieri DA, Daro FR, Ribeiro PL, de Andrade AG. Comparing topiramate with naltrexone in the treatment of alcohol dependence. *Addiction*. Dec 2008;103(12):2035-2044.
31. Baltieri DA, Daro FR, Ribeiro PL, Andrade AG. Effects of topiramate or naltrexone on tobacco use among male alcohol-dependent outpatients. *Drug Alcohol Depend*. Nov 1 2009;105(1-2):33-41.
32. Berger L, Fisher M, Brondino M, et al. Efficacy of acamprosate for alcohol dependence in a family medicine setting in the United States: A randomized, double-blind, placebo-controlled study. *Alcohol*. 2013;37(4):668-674.

33. Besson J, Aeby F, Kasas A, Lehert P, Potgieter A. Combined efficacy of acamprosate and disulfiram in the treatment of alcoholism: a controlled study. *Alcohol. Clin. Exp. Res.* May 1998;22(3):573-579.
34. Book SW, Thomas SE, Randall PK, Randall CL. Paroxetine reduces social anxiety in individuals with a co-occurring alcohol use disorder. *J. Anxiety Disord.* 2008;22(2):310-318.
35. Thomas SE, Randall PK, Book SW, Randall CL. A complex relationship between co-occurring social anxiety and alcohol use disorders: what effect does treating social anxiety have on drinking? *Alcohol. Clin. Exp. Res.* Jan 2008;32(1):77-84.
36. Brady KT, Myrick H, Henderson S, Coffey SF. The use of divalproex in alcohol relapse prevention: a pilot study. *Drug Alcohol Depend.* Aug 1 2002;67(3):323-330.
37. Brady KT, Sonne S, Anton RF, Randall CL, Back SE, Simpson K. Sertraline in the treatment of co-occurring alcohol dependence and posttraumatic stress disorder. *Alcohol. Clin. Exp. Res.* Mar 2005;29(3):395-401.
38. Brown ES, Garza M, Carmody TJ. A randomized, double-blind, placebo-controlled add-on trial of quetiapine in outpatients with bipolar disorder and alcohol use disorders. *J. Clin. Psychiatry.* May 2008;69(5):701-705.
39. Brown ES, Carmody TJ, Schmitz JM, et al. A randomized, double-blind, placebo-controlled pilot study of naltrexone in outpatients with bipolar disorder and alcohol dependence. *Alcohol. Clin. Exp. Res.* Nov 2009;33(11):1863-1869.
40. Chick J, Howlett H, Morgan MY, Ritson B. United Kingdom Multicentre Acamprosate Study (UKMAS): a 6-month prospective study of acamprosate versus placebo in preventing relapse after withdrawal from alcohol. *Alcohol Alcohol.* Mar-Apr 2000;35(2):176-187.
41. Chick J, Anton R, Chęcinski K, et al. A multicentre, randomized, double-blind, placebo-controlled trial of naltrexone in the treatment of alcohol dependence or abuse. *Alcohol Alcohol.* Nov-Dec 2000;35(6):587-593.
42. Chick J, Aschauer H, Hornik K. Efficacy of fluvoxamine in preventing relapse in alcohol dependence: a one-year, double-blind, placebo-controlled multicentre study with analysis by typology. *Drug Alcohol Depend.* Apr 9 2004;74(1):61-70.
43. Cornelius JR, Salloum IM, Ehler JG, et al. Fluoxetine in depressed alcoholics. A double-blind, placebo-controlled trial. *Arch. Gen. Psychiatry.* Aug 1997;54(8):700-705.
44. Cornelius JR, Salloum IM, Cornelius MD, et al. Preliminary report: double-blind, placebo-controlled study of fluoxetine in depressed alcoholics. *Psychopharmacol. Bull.* 1995;31(2):297-303.
45. Corrêa Filho JM, Baltieri DA. A pilot study of full-dose ondansetron to treat heavy-drinking men withdrawing from alcohol in Brazil. *Addict. Behav.* 2013;38(4):2044-2051.
46. Coskunol H, Gökden O, Ercan ES, Bayraktar E, Tuglular I, Saygili R. Long-term efficacy of sertraline in the prevention of alcoholic relapses in alcohol-dependent patients: a single-center, double-blind, randomized, placebo-controlled, parallel-group study. *Current Therapeutic Research.* 2002;63(11):759-771.
47. De Sousa A, De Sousa A. A one-year pragmatic trial of naltrexone vs disulfiram in the treatment of alcohol dependence. *Alcohol Alcohol.* Nov-Dec 2004;39(6):528-531.
48. De Sousa A, De Sousa A. An open randomized study comparing disulfiram and acamprosate in the treatment of alcohol dependence. *Alcohol Alcohol.* Nov-Dec 2005;40(6):545-548.

49. Fawcett J, Kravitz HM, McGuire M, et al. Pharmacological treatments for alcoholism: revisiting lithium and considering buspirone. *Alcohol. Clin. Exp. Res.* May 2000;24(5):666-674.
50. Florez G, Garcia-Portilla P, Alvarez S, Saiz PA, Nogueiras L, Bobes J. Using topiramate or naltrexone for the treatment of alcohol-dependent patients. *Alcohol. Clin. Exp. Res.* Jul 2008;32(7):1251-1259.
51. Florez G, Saiz PA, Garcia-Portilla P, Alvarez S, Nogueiras L, Bobes J. Topiramate for the treatment of alcohol dependence: comparison with naltrexone. *Eur. Addict. Res.* 2011;17(1):29-36.
52. Fogaca MN, Santos-Galduroz RF, Eserian JK, Galduroz JC. The effects of polyunsaturated fatty acids in alcohol dependence treatment--a double-blind, placebo-controlled pilot study. *BMC Clin Pharmacol.* 2011;11:10.
53. Fuller RK, Roth HP. Disulfiram for the treatment of alcoholism. An evaluation in 128 men. *Ann. Intern. Med.* Jun 1979;90(6):901-904.
54. Fuller RK, Branchey L, Brightwell DR, et al. Disulfiram treatment of alcoholism. A Veterans Administration cooperative study. *JAMA.* Sep 19 1986;256(11):1449-1455.
55. Garbutt JC, Kranzler HR, O'Malley SS, et al. Efficacy and tolerability of long-acting injectable naltrexone for alcohol dependence: a randomized controlled trial. *JAMA.* Apr 6 2005;293(13):1617-1625.
56. Pettinati HM, Gastfriend DR, Dong Q, Kranzler HR, O'Malley SS. Effect of extended-release naltrexone (XR-NTX) on quality of life in alcohol-dependent patients. *Alcohol. Clin. Exp. Res.* Feb 2009;33(2):350-356.
57. Lucey MR, Silverman BL, Illeperuma A, O'Brien CP. Hepatic safety of once-monthly injectable extended-release naltrexone administered to actively drinking alcoholics. *Alcohol. Clin. Exp. Res.* Mar 2008;32(3):498-504.
58. Garbutt JC, Kampov-Polevoy AB, Gallop R, Kalka-Juhl L, Flannery BA. Efficacy and safety of baclofen for alcohol dependence: a randomized, double-blind, placebo-controlled trial. *Alcohol. Clin. Exp. Res.* Nov 2010;34(11):1849-1857.
59. Gastpar M, Bonnet U, Boning J, et al. Lack of efficacy of naltrexone in the prevention of alcohol relapse: results from a German multicenter study. *J. Clin. Psychopharmacol.* Dec 2002;22(6):592-598.
60. Geerlings PJ, Ansoms C, Van Den Brink W. Acamprosate and prevention of relapse in alcoholics. Results of a randomized, placebo-controlled, double-blind study in out-patient alcoholics in the Netherlands, Belgium and Luxembourg. *Eur. Addict. Res.* 1997;3(3):129-137.
61. George DT, Rawlings R, Eckardt MJ, Phillips MJ, Shoaf SE, Linnoila M. Buspirone treatment of alcoholism: age of onset, and cerebrospinal fluid 5-hydroxyindolacetic acid and homovanillic acid concentrations, but not medication treatment, predict return to drinking. *Alcohol. Clin. Exp. Res.* Feb 1999;23(2):272-278.
62. Gual A, Leher P. Acamprosate during and after acute alcohol withdrawal: a double-blind placebo-controlled study in Spain. *Alcohol Alcohol.* 2001;36(5):413-418.
63. Gual A, Balcells M, Torres M, Madrigal M, Diez T, Serrano L. Sertraline for the prevention of relapse in detoxicated alcohol dependent patients with a comorbid depressive disorder: a randomized controlled trial. *Alcohol Alcohol.* Nov-Dec 2003;38(6):619-625.
64. Gual A, He Y, Torup L, van den Brink W, Mann K. A randomised, double-blind, placebo-controlled, efficacy study of nalmefene, as-needed use, in patients with alcohol dependence. *Eur. Neuropsychopharmacol.* Apr 3 2013.

65. Guardia J, Caso C, Arias F, et al. A double-blind, placebo-controlled study of naltrexone in the treatment of alcohol-dependence disorder: results from a multicenter clinical trial. *Alcohol. Clin. Exp. Res.* Sep 2002;26(9):1381-1387.
66. Guardia J, Segura L, Gonzalvo B, et al. A double-blind, placebo-controlled study of olanzapine in the treatment of alcohol-dependence disorder. *Alcohol. Clin. Exp. Res.* May 2004;28(5):736-745.
67. Heinala P, Alho H, Kiianmaa K, Lonnqvist J, Kuoppasalmi K, Sinclair JD. Targeted use of naltrexone without prior detoxification in the treatment of alcohol dependence: a factorial double-blind, placebo-controlled trial. *J. Clin. Psychopharmacol.* Jun 2001;21(3):287-292.
68. Huang MC, Chen CH, Yu JM, Chen CC. A double-blind, placebo-controlled study of naltrexone in the treatment of alcohol dependence in Taiwan. *Addict Biol.* Sep 2005;10(3):289-292.
69. Johnson BA, Ait-Daoud N, Bowden CL, et al. Oral topiramate for treatment of alcohol dependence: a randomised controlled trial. *Lancet.* May 17 2003;361(9370):1677-1685.
70. Ma JZ, Ait-Daoud N, Johnson BA. Topiramate reduces the harm of excessive drinking: implications for public health and primary care. *Addiction.* Nov 2006;101(11):1561-1568.
71. Johnson BA, Ait-Daoud N, Akhtar FZ, Ma JZ. Oral topiramate reduces the consequences of drinking and improves the quality of life of alcohol-dependent individuals: a randomized controlled trial. *Arch. Gen. Psychiatry.* Sep 2004;61(9):905-912.
72. Johnson BA, Ait-Daoud N, Aubin HJ, et al. A pilot evaluation of the safety and tolerability of repeat dose administration of long-acting injectable naltrexone (Vivitrex) in patients with alcohol dependence. *Alcohol. Clin. Exp. Res.* Sep 2004;28(9):1356-1361.
73. Johnson BA, Rosenthal N, Capece JA, et al. Topiramate for treating alcohol dependence: a randomized controlled trial. *JAMA.* Oct 10 2007;298(14):1641-1651.
74. Johnson BA, Rosenthal N, Capece JA, et al. Improvement of physical health and quality of life of alcohol-dependent individuals with topiramate treatment: US multisite randomized controlled trial. *Arch. Intern. Med.* Jun 9 2008;168(11):1188-1199.
75. Kabel DI, Petty F. A placebo-controlled, double-blind study of fluoxetine in severe alcohol dependence: adjunctive pharmacotherapy during and after inpatient treatment. *Alcohol. Clin. Exp. Res.* Jun 1996;20(4):780-784.
76. Kampman KM, Pettinati HM, Lynch KG, et al. A double-blind, placebo-controlled pilot trial of quetiapine for the treatment of Type A and Type B alcoholism. *J. Clin. Psychopharmacol.* Aug 2007;27(4):344-351.
77. Kampman KM, Pettinati HM, Lynch KG, Spratt K, Wierzbicki MR, O'Brien CP. A double-blind, placebo-controlled trial of topiramate for the treatment of comorbid cocaine and alcohol dependence. *Drug Alcohol Depend.* 1 2013;133(1):94-99.
78. Karhuvaara S, Simojoki K, Virta A, et al. Targeted nalmefene with simple medical management in the treatment of heavy drinkers: a randomized double-blind placebo-controlled multicenter study. *Alcohol. Clin. Exp. Res.* Jul 2007;31(7):1179-1187.
79. Kiefer F, Jahn H, Tarnaske T, et al. Comparing and combining naltrexone and acamprosate in relapse prevention of alcoholism: a double-blind, placebo-controlled study. *Arch. Gen. Psychiatry.* Jan 2003;60(1):92-99.
80. Kiefer F, Andersohn F, Otte C, Wolf K, Jahn H, Wiedemann K. Long-term effects of pharmacotherapy on relapse prevention in alcohol dependence. *Acta Neuropsychiatrica.* 2004;18:233-238.

81. Kiefer F, Helwig H, Tarnaske T, Otte C, Jahn H, Wiedemann K. Pharmacological relapse prevention of alcoholism: clinical predictors of outcome. *Eur. Addict. Res.* 2005;11(2):83-91.
82. Killeen TK, Brady KT, Gold PB, et al. Effectiveness of naltrexone in a community treatment program. *Alcohol. Clin. Exp. Res.* Nov 2004;28(11):1710-1717.
83. Kranzler HR, Burleson JA, Del Boca FK, et al. Buspirone treatment of anxious alcoholics. A placebo-controlled trial. *Arch. Gen. Psychiatry.* Sep 1994;51(9):720-731.
84. Kranzler HR, Burleson JA, Korner P, et al. Placebo-controlled trial of fluoxetine as an adjunct to relapse prevention in alcoholics. *A. J. Psychiatry.* Mar 1995;152(3):391-397.
85. Kranzler HR, Wesson DR, Billot L. Naltrexone depot for treatment of alcohol dependence: a multicenter, randomized, placebo-controlled clinical trial. *Alcohol. Clin. Exp. Res.* Jul 2004;28(7):1051-1059.
86. Kranzler HR, Tennen H, Armeli S, et al. Targeted naltrexone for problem drinkers. *J. Clin. Psychopharmacol.* Aug 2009;29(4):350-357.
87. Kranzler HR, Armeli S, Tennen H, et al. A double-blind, randomized trial of sertraline for alcohol dependence: moderation by age of onset [corrected] and 5-hydroxytryptamine transporter-linked promoter region genotype. *J. Clin. Psychopharmacol.* Feb 2011;31(1):22-30.
88. Kranzler HR, Armeli S, Tennen H. Post-treatment outcomes in a double-blind, randomized trial of sertraline for alcohol dependence. *Alcohol. Clin. Exp. Res.* Apr 2012;36(4):739-744.
89. Krystal JH, Cramer JA, Krol WF, Kirk GF, Rosenheck RA. Naltrexone in the treatment of alcohol dependence. *N. Engl. J. Med.* Dec 13 2001;345(24):1734-1739.
90. Laaksonen E, Koski-Jannes A, Salaspuro M, Ahtinen H, Alho H. A randomized, multicentre, open-label, comparative trial of disulfiram, naltrexone and acamprosate in the treatment of alcohol dependence. *Alcohol Alcohol.* Jan-Feb 2008;43(1):53-61.
91. Laaksonen E, Vuoristo-Myllys S, Koski-Jannes A, Alho H. Combining medical treatment and CBT in treating alcohol-dependent patients: Effects on life quality and general well-being. *Alcohol Alcohol.* 2013;48(6):687-693.
92. Latt NC, Jurd S, Houseman J, Wutzke SE. Naltrexone in alcohol dependence: a randomised controlled trial of effectiveness in a standard clinical setting. *Med. J. Aust.* Jun 3 2002;176(11):530-534.
93. Lee A, Tan S, Lim D, et al. Naltrexone in the treatment of male alcoholics—An effectiveness study In Singapore. *Drug and Alcohol Review.* 2001;20(2):193-199.
94. Lhuintre JP, Daoust M, Moore ND, et al. Ability of calcium bis acetyl homotaurine, a GABA agonist, to prevent relapse in weaned alcoholics. *Lancet.* May 4 1985;1(8436):1014-1016.
95. Lhuintre JP, Moore N, Tran G, et al. Acamprosate appears to decrease alcohol intake in weaned alcoholics. *Alcohol Alcohol.* 1990;25(6):613-622.
96. Likhitsathian S, Uttawichai K, Booncharoen H, Wittayanookulluk A, Angkurawaranon C, Srisurapanont M. Topiramate treatment for alcoholic outpatients recently receiving residential treatment programs: A 12-week, randomized, placebo-controlled trial. *Drug Alcohol Depend.* 2013;133(2):440-446.
97. Ling W, Weiss DG, Charuvastra VC, O'Brien CP. Use of disulfiram for alcoholics in methadone maintenance programs. A Veterans Administration Cooperative Study. *Arch. Gen. Psychiatry.* Aug 1983;40(8):851-854.

98. Litten RZ, Ryan ML, Fertig JB, et al. A double-blind, placebo-controlled trial assessing the efficacy of varenicline tartrate for alcohol dependence. *J. Addict. Med.* Jul-Aug 2013;7(4):277-286.
99. Longabaugh R, Wirtz PW, Gulliver SB, Davidson D. Extended naltrexone and broad spectrum treatment or motivational enhancement therapy. *Psychopharmacology (Berl)*. Oct 2009;206(3):367-376.
100. Malcolm R, Anton RF, Randall CL, Johnston A, Brady K, Thevos A. A placebo-controlled trial of buspirone in anxious inpatient alcoholics. *Alcohol. Clin. Exp. Res.* Dec 1992;16(6):1007-1013.
101. Malec E, Malec T, Gagne MA, Dongier M. Buspirone in the treatment of alcohol dependence: a placebo-controlled trial. *Alcohol. Clin. Exp. Res.* Apr 1996;20(2):307-312.
102. Mann K, Lemenager T, Hoffmann S, et al; PREDICT Study Team. Results of a double-blind, placebo-controlled pharmacotherapy trial in alcoholism conducted in Germany and comparison with the US COMBINE study. *Addict Biol.* 2013;18(6):937-946.
103. Mann K, Bladström A, Torup L, Gual A, van den Brink W. Extending the treatment options in alcohol dependence: A randomized controlled study of as-needed nalmefene. *Biol. Psychiatry.* 2013;73(8):706-713.
104. Martinotti G, Di Nicola M, Di Giannantonio M, Janiri L. Aripiprazole in the treatment of patients with alcohol dependence: a double-blind, comparison trial vs. naltrexone. *J Psychopharmacol.* Mar 2009;23(2):123-129.
105. Mason BJ, Ritvo EC, Morgan RO, et al. A double-blind, placebo-controlled pilot study to evaluate the efficacy and safety of oral nalmefene HCl for alcohol dependence. *Alcohol.* 1994;18(5):1162-1167.
106. Mason BJ, Kocsis JH, Ritvo EC, Cutler RB. A double-blind, placebo-controlled trial of desipramine for primary alcohol dependence stratified on the presence or absence of major depression. *JAMA.* Mar 13 1996;275(10):761-767.
107. Mason BJ, Salvato FR, Williams LD, Ritvo EC, Robert B C. A double-blind, placebo-controlled study of oral nalmefene for alcohol dependence. *Arch. Gen. Psychiatry.* 1999;56(8):719-724.
108. Mason BJ, Goodman AM, Chabac S, Lehert P. Effect of oral acamprosate on abstinence in patients with alcohol dependence in a double-blind, placebo-controlled trial: the role of patient motivation. *J. Psychiatr. Res.* Aug 2006;40(5):383-393.
109. Mason BJ, Quello S, Goodell V, Shadan F, Kyle M, Begovic A. Gabapentin treatment for alcohol dependence: a randomized clinical trial. *JAMA Intern Med.* Jan 2014;174(1):70-77.
110. McGrath PJ, Nunes EV, Stewart JW, et al. Imipramine treatment of alcoholics with primary depression: A placebo-controlled clinical trial. *Arch. Gen. Psychiatry.* Mar 1996;53(3):232-240.
111. Moak DH, Anton RF, Latham PK, Voronin KE, Waid RL, Durazo-Arvizu R. Sertraline and cognitive behavioral therapy for depressed alcoholics: results of a placebo-controlled trial. *J. Clin. Psychopharmacol.* Dec 2003;23(6):553-562.
112. Monterosso JR, Flannery BA, Pettinati HM, et al. Predicting treatment response to naltrexone: the influence of craving and family history. *Am. J. Addict.* Summer 2001;10(3):258-268.
113. Monti PM, Rohsenow DJ, Swift RM, et al. Naltrexone and cue exposure with coping and communication skills training for alcoholics: treatment process and 1-year outcomes. *Alcohol. Clin. Exp. Res.* Nov 2001;25(11):1634-1647.

114. Rohsenow DJ, Miranda R, Jr., McGeary JE, Monti PM. Family history and antisocial traits moderate naltrexone's effects on heavy drinking in alcoholics. *Exp Clin Psychopharmacol*. Jun 2007;15(3):272-281.
115. Rohsenow DJ, Colby SM, Monti PM, et al. Predictors of compliance with naltrexone among alcoholics. *Alcohol*. 2000;24(10):1542-1549.
116. Morgenstern J, Kuerbis AN, Chen AC, Kahler CW, Bux DA, Jr., Kranzler HR. A randomized clinical trial of naltrexone and behavioral therapy for problem drinking men who have sex with men. *J. Consult. Clin. Psychol*. 2012;80(5):863-875.
117. Morley KC, Teesson M, Reid SC, et al. Naltrexone versus acamprosate in the treatment of alcohol dependence: A multi-centre, randomized, double-blind, placebo-controlled trial. *Addiction*. Oct 2006;101(10):1451-1462.
118. Morley KC, Teesson M, Sannibale C, Baillie A, Haber PS. Clinical predictors of outcome from an Australian pharmacological relapse prevention trial. *Alcohol Alcohol*. Nov-Dec 2010;45(6):520-526.
119. Morris PL, Hopwood M, Whelan G, Gardiner J, Drummond E. Naltrexone for alcohol dependence: a randomized controlled trial. *Addiction*. Nov 2001;96(11):1565-1573.
120. Naranjo CA, Bremner KE, Lanctot KL. Effects of citalopram and a brief psycho-social intervention on alcohol intake, dependence and problems. *Addiction*. Jan 1995;90(1):87-99.
121. Narayana PL, Gupta AK, Sharma PK. Use of anti-craving agents in soldiers with alcohol dependence syndrome. *Medical Journal Armed Forces India*. 2008;64(4):320-324.
122. Nava F, Premi S, Manzato E, Lucchini A. Comparing treatments of alcoholism on craving and biochemical measures of alcohol consumption. *J. Psychoactive Drugs*. Sep 2006;38(3):211-217.
123. O'Malley SS, Jaffe AJ, Chang G, Schottenfeld RS, Meyer RE, Rounsaville B. Naltrexone and coping skills therapy for alcohol dependence. A controlled study. *Arch. Gen. Psychiatry*. Nov 1992;49(11):881-887.
124. O'Malley SS, Jaffe AJ, Chang G, et al. Six-month follow-up of naltrexone and psychotherapy for alcohol dependence. *Arch. Gen. Psychiatry*. Mar 1996;53(3):217-224.
125. O'Malley SS, Sinha R, Grilo CM, et al. Naltrexone and cognitive behavioral coping skills therapy for the treatment of alcohol drinking and eating disorder features in alcohol-dependent women: a randomized controlled trial. *Alcohol. Clin. Exp. Res*. Apr 2007;31(4):625-634.
126. O'Malley SS, Robin RW, Levenson AL, et al. Naltrexone alone and with sertraline for the treatment of alcohol dependence in Alaska natives and non-natives residing in rural settings: a randomized controlled trial. *Alcohol. Clin. Exp. Res*. Jul 2008;32(7):1271-1283.
127. Oslin D, Liberto JG, O'Brien J, Krois S, Norbeck J. Naltrexone as an adjunctive treatment for older patients with alcohol dependence. *Am. J. Geriatr. Psychiatry*. Fall 1997;5(4):324-332.
128. Oslin DW, Lynch KG, Pettinati HM, et al. A placebo-controlled randomized clinical trial of naltrexone in the context of different levels of psychosocial intervention. *Alcohol. Clin. Exp. Res*. Jul 2008;32(7):1299-1308.
129. Paille FM, Guelfi JD, Perkins AC, Royer RJ, Steru L, Parot P. Double-blind randomized multicentre trial of acamprosate in maintaining abstinence from alcohol. *Alcohol Alcohol*. Mar 1995;30(2):239-247.
130. Pelc I, Le Bon O, Lehert P, Verbanck P. Acamprosate in the Treatment of Alcohol Dependence: A 6-Month Postdetoxification Study. In: Soyka M, ed. *Acamprosate in Relapse Prevention of Alcoholism*: Springer Berlin Heidelberg; 1996:133-142.

131. Pelc I, Le Bon O, Verbanck P, Lehert P, Opsomer L. Calciumacetylhomotaurinate for maintaining abstinence in weaned alcoholic patients: a placebo-controlled double-blind multi-centre study. In: Naranjo CA, Sellers EM, eds. *Novel Pharmacological Interventions for Alcoholism*. New York: Springer-Verlag; 1992.
132. Pelc I, Verbanck P, Le Bon O, Gavrilovic M, Lion K, Lehert P. Efficacy and safety of acamprosate in the treatment of detoxified alcohol-dependent patients. A 90-day placebo-controlled dose-finding study. *Br. J. Psychiatry*. Jul 1997;171:73-77.
133. Petrakis IL, O'Malley S, Rounsaville B, Poling J, McHugh-Strong C, Krystal JH. Naltrexone augmentation of neuroleptic treatment in alcohol abusing patients with schizophrenia. *Psychopharmacology (Berl)*. Mar 2004;172(3):291-297.
134. Ralevski E, Balachandra K, Gueorguieva R, Limoncelli D, Petrakis I. Effects of naltrexone on cognition in a treatment study of patients with schizophrenia and comorbid alcohol dependence. *J. Dual Diagn*. 2006;2(4):53-69.
135. Petrakis IL, Poling J, Levinson C, Nich C, Carroll K, Rounsaville B. Naltrexone and disulfiram in patients with alcohol dependence and comorbid psychiatric disorders. *Biol. Psychiatry*. May 15 2005;57(10):1128-1137.
136. Ralevski E, Ball S, Nich C, Limoncelli D, Petrakis I. The impact of personality disorders on alcohol-use outcomes in a pharmacotherapy trial for alcohol dependence and comorbid Axis I disorders. *Am. J. Addict*. Nov-Dec 2007;16(6):443-449.
137. Petrakis I, Ralevski E, Nich C, et al. Naltrexone and disulfiram in patients with alcohol dependence and current depression. *J. Clin. Psychopharmacol*. Apr 2007;27(2):160-165.
138. Petrakis IL, Poling J, Levinson C, et al. Naltrexone and disulfiram in patients with alcohol dependence and comorbid post-traumatic stress disorder. *Biol. Psychiatry*. Oct 1 2006;60(7):777-783.
139. Petrakis IL, Ralevski E, Desai N, et al. Noradrenergic vs serotonergic antidepressant with or without naltrexone for veterans with PTSD and comorbid alcohol dependence. *Neuropsychopharmacology*. Mar 2012;37(4):996-1004.
140. Pettinati HM, Volpicelli JR, Luck G, Kranzler HR, Rukstalis MR, Cnaan A. Double-blind clinical trial of sertraline treatment for alcohol dependence. *J. Clin. Psychopharmacol*. Apr 2001;21(2):143-153.
141. Pettinati HM, Kampman KM, Lynch KG, et al. Gender differences with high-dose naltrexone in patients with co-occurring cocaine and alcohol dependence. *J. Subst. Abuse Treat*. Jun 2008;34(4):378-390.
142. Pettinati HM, Oslin DW, Kampman KM, et al. A double-blind, placebo-controlled trial combining sertraline and naltrexone for treating co-occurring depression and alcohol dependence. *A. J. Psychiatry*. Jun 2010;167(6):668-675.
143. Plebani JG, Lynch KG, Rennert L, Pettinati HM, O'Brien CP, Kampman KM. Results from a pilot clinical trial of varenicline for the treatment of alcohol dependence. *Drug Alcohol Depend*. 2013;133(2):754-758.
144. Poldrugo F. Acamprosate treatment in a long-term community-based alcohol rehabilitation programme. *Addiction*. Nov 1997;92(11):1537-1546.
145. Ralevski E, O'Brien E, Jane JS, Dean E, Dwan R, Petrakis I. Effects of acamprosate on cognition in a treatment study of patients with schizophrenia spectrum disorders and comorbid alcohol dependence. *J. Nerv. Ment. Dis*. Jul 2011;199(7):499-505.
146. Ralevski E, O'Brien E, Jane JS, et al. Treatment with acamprosate in patients with schizophrenia spectrum disorders and comorbid alcohol dependence. *J. Dual Diagn*. 2011;7(1-2):64-73.

147. Rubio G, Jimenez-Arriero MA, Ponce G, Palomo T. Naltrexone versus acamprosate: one year follow-up of alcohol dependence treatment. *Alcohol Alcohol.* Sep-Oct 2001;36(5):419-425.
148. Rubio G, Martinez-Gras I, Manzanares J. Modulation of impulsivity by topiramate: implications for the treatment of alcohol dependence. *J. Clin. Psychopharmacol.* Dec 2009;29(6):584-589.
149. Salloum IM, Cornelius JR, Daley DC, Kirisci L, Himmelhoch JM, Thase ME. Efficacy of valproate maintenance in patients with bipolar disorder and alcoholism: a double-blind placebo-controlled study. *Arch. Gen. Psychiatry.* Jan 2005;62(1):37-45.
150. Sass H, Soyka M, Mann K, Zieglsangberger W. Relapse prevention by acamprosate. Results from a placebo-controlled study on alcohol dependence. *Arch. Gen. Psychiatry.* Aug 1996;53(8):673-680.
151. Schmitz JM, Stotts AL, Sayre SL, DeLaune KA, Grabowski J. Treatment of cocaine-alcohol dependence with naltrexone and relapse prevention therapy. *Am. J. Addict.* Jul-Sep 2004;13(4):333-341.
152. Schmitz JM, Lindsay JA, Green CE, Herin DV, Stotts AL, Moeller FG. High-dose naltrexone therapy for cocaine-alcohol dependence. *Am. J. Addict.* Sep-Oct 2009;18(5):356-362.
153. Safety and Efficacy of Nalmefene in Patients with Alcohol Dependence (SENSE). 2013.
154. Stedman M, Pettinati HM, Brown ES, Kotz M, Calabrese JR, Raines S. A double-blind, placebo-controlled study with quetiapine as adjunct therapy with lithium or divalproex in bipolar I patients with coexisting alcohol dependence. *Alcohol. Clin. Exp. Res.* Oct 2010;34(10):1822-1831.
155. Tempesta E, Janiri L, Bignamini A, Chabac S, Potgieter A. Acamprosate and relapse prevention in the treatment of alcohol dependence: a placebo-controlled study. *Alcohol Alcohol.* Mar-Apr 2000;35(2):202-209.
156. Tiihonen J, Ryynanen OP, Kauhanen J, Hakola HP, Salaspuro M. Citalopram in the treatment of alcoholism: a double-blind placebo-controlled study. *Pharmacopsychiatry.* Jan 1996;29(1):27-29.
157. Tollefson GD, Lancaster SP, Montague-Clouse J. The association of buspirone and its metabolite 1-pyrimidinylpiperazine in the remission of comorbid anxiety with depressive features and alcohol dependency. *Psychopharmacol. Bull.* 1991;27(2):163-170.
158. Tollefson GD, Montague-Clouse J, Tollefson SL. Treatment of comorbid generalized anxiety in a recently detoxified alcoholic population with a selective serotonergic drug (buspirone). *J. Clin. Psychopharmacol.* Feb 1992;12(1):19-26.
159. Volpicelli JR, Clay KL, Watson NT, O'Brien CP. Naltrexone in the treatment of alcoholism: predicting response to naltrexone. *J. Clin. Psychiatry.* 1995;56 Suppl 7:39-44.
160. Volpicelli JR, Alterman AI, Hayashida M, O'Brien CP. Naltrexone in the treatment of alcohol dependence. *Arch. Gen. Psychiatry.* Nov 1992;49(11):876-880.
161. Volpicelli JR, Rhines KC, Rhines JS, Volpicelli LA, Alterman AI, O'Brien CP. Naltrexone and alcohol dependence. Role of subject compliance. *Arch. Gen. Psychiatry.* Aug 1997;54(8):737-742.
162. Whitworth AB, Fischer F, Lesch OM, et al. Comparison of acamprosate and placebo in long-term treatment of alcohol dependence. *Lancet.* May 25 1996;347(9013):1438-1442.
163. Wilens TE, Adler LA, Weiss MD, et al. Atomoxetine treatment of adults with ADHD and comorbid alcohol use disorders. *Drug Alcohol Depend.* Jul 1 2008;96(1-2):145-154.

164. Wolwer W, Frommann N, Janner M, et al. The effects of combined acamprosate and integrative behaviour therapy in the outpatient treatment of alcohol dependence: a randomized controlled trial. *Drug Alcohol Depend.* Nov 1 2011;118(2-3):417-422.
165. Kranzler HR, Armeli S, Feinn R, Tennen H. Targeted naltrexone treatment moderates the relations between mood and drinking behavior among problem drinkers. *J. Consult. Clin. Psychol.* Apr 2004;72(2):317-327.
166. Oslin D, Liberto JG, O'Brien J, Krois S. Tolerability of naltrexone in treating older, alcohol-dependent patients. *Am. J. Addict.* Summer 1997;6(3):266-270.
167. Pettinati HM, Kampman KM, Lynch KG, et al. A double blind, placebo-controlled trial that combines disulfiram and naltrexone for treating co-occurring cocaine and alcohol dependence. *Addict. Behav.* May 2008;33(5):651-667.
168. Kiritze-Topor P, Huas D, Rosenzweig C, Comte S, Paille F, Lehert P. A pragmatic trial of acamprosate in the treatment of alcohol dependence in primary care. *Alcohol Alcohol.* Nov-Dec 2004;39(6):520-527.
169. O'Malley SS, Rounsaville BJ, Farren C, et al. Initial and maintenance naltrexone treatment for alcohol dependence using primary care vs specialty care: a nested sequence of 3 randomized trials. *Arch. Intern. Med.* Jul 28 2003;163(14):1695-1704.
170. Oslin DW, Lynch KG, Maisto SA, et al. A Randomized Clinical Trial of Alcohol Care Management Delivered in Department of Veterans Affairs Primary Care Clinics Versus Specialty Addiction Treatment. *J. Gen. Intern. Med.* Sep 20 2013.
171. Saitz R, Cheng DM, Winter M, et al. Chronic care management for dependence on alcohol and other drugs: the AHEAD randomized trial. *JAMA.* Sep 18 2013;310(11):1156-1167.