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ORIGINAL RESEARCH

The effects of memantine on behavioral disturbances in patients with Alzheimer's disease: a meta-analysis

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Background: Memantine is effective in the treatment of behavioral disturbances in patients with Alzheimer's disease. It has not yet been fully determined which behavioral disturbances respond best to memantine.

Methods: We conducted a meta-analysis of memantine vs control (placebo or usual care) for the treatment of individual behavioral disturbances (delusion, hallucination, agitation/aggression, dysphoria, anxiety/phobia, euphoria, apathy, disinhibition, irritability/lability, aberrant motor activity/activity disturbances, nighttime disturbance/diurnal rhythm disturbances, and eating disturbances). Randomized controlled studies of memantine in patients with Alzheimer's disease were included in this study. To evaluate these outcomes, standardized mean difference (SMD), with 95% confidence intervals (95% CIs), based upon a random-effects model was evaluated in the meta-analysis.

Results: A total of 11 studies (n=4,261; memantine vs placebo: N=4, n=1,500; memantine + cholinesterase inhibitors [M + ChEIs] vs ChEIs: N=7, n=2,761) were included in the meta-analysis. Compared to control, memantine showed significant improvement in agitation/aggression (SMD =−0.11; 95% CIs =−0.20, −0.03; P=0.01; I^2=47%), delusion (SMD =−0.12; 95% CIs =−0.18, −0.06; P=0.0002; I^2=0%), disinhibition (SMD =−0.08; 95% CIs =−0.15, −0.00; P=0.04; I^2=0%), and nighttime disturbance/diurnal rhythm disturbances (SMD =−0.10; 95% CIs =−0.18, −0.02; P=0.02; I^2=36%). Memantine was also marginally superior to control in hallucination (SMD =−0.06; 95% CIs =−0.12, 0.01; P=0.07; I^2=0%) and irritability/lability (SMD =−0.09; 95% CIs =−0.19, 0.01; P=0.07; I^2=42%). Memantine is similar to control in dysphoria, anxiety/phobia, euphoria, apathy, and eating disturbance.

Conclusion: The meta-analysis suggest that memantine has benefits for the treatment of most of the behavioral disturbances in patients with Alzheimer's disease. Memantine does not deteriorate negative symptoms as behavioral disturbances in patients with Alzheimer's disease.

Keywords: memantine, Alzheimer's disease, behavioral disturbances, meta-analysis

Introduction

Alzheimer's disease is a neurodegenerative disease.[1] The percentage of people with Alzheimer's disease increases with age: 3% of people aged 65–74 years, 17% of people aged 75–84 years, and 32% of people aged 85 years and older have Alzheimer's disease.[2] It has an insidious onset, with gradual progression of cognitive symptoms and behavioral disturbances.[1]

There are the following four approved drugs for the treatment of Alzheimer's disease worldwide: memantine and three cholinesterase inhibitors (donepezil, galantamine, and rivastigmine).[1] Memantine has been approved worldwide for treating

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moderate-to-severe Alzheimer's disease. It is postulated that memantine exerts its therapeutic effect through its action as a low-to-moderate affinity, noncompetitive (open channel), nonselective, voltage-dependent, *N*-methyl-D-aspartic acid (NMDA) receptor antagonist, which binds preferentially to NMDA receptor-operated calcium channels.[3] Memantine blocks the effects of sustained, pathologically elevated levels of glutamate, which could otherwise lead to neuronal dysfunction.[4–6] In addition, memantine may also upregulate NMDA receptor expression, causing activation in the presence of a strong stimulus.[7]

Our previous meta-analysis showed that memantine monotherapy was superior to placebo in cognitive impairment (standardized mean difference [SMD] =−0.27; 95% confidence intervals [95% CIs] =−0.39 to −0.14) and behavioral disturbances (SMD =−0.12; 95% CIs =−0.22 to −0.01).[8] We did an additional meta-analysis to show that although there was a trend favoring the combination therapy with memantine and cholinesterase inhibitors compared to cholinesterase inhibitor monotherapy for treating cognitive impairment (SMD =−0.13; 95% CIs =−0.26 to 0.01), memantine was superior to placebo in behavioral disturbances (SMD =−0.13; 95% CIs =−0.24 to −0.02).[9] Thus, there was evidence on the efficacy of memantine for cognitive impairment and behavioral disturbances on patients with Alzheimer's disease to date.

However, there are various symptoms of behavioral disturbances, such as delusion, hallucination, agitation/aggression, dysphoria, anxiety/phobia, euphoria, apathy, disinhibition, irritability/lability, aberrant motor activity/activity disturbances, nighttime disturbance/diurnal rhythm disturbances, and eating disturbances.[10] For example, although a drug, which has sedative effect, seems to be effective for positive symptoms, such as agitation and irritability, this drug seems to exasperate negative symptoms, such as apathy.[10] There has not been robust evidence on the efficacy of memantine for individual behavioral disturbances in patients with Alzheimer's disease. The effect size of anti-dementia drugs for individual behavioral disturbances in patients with Alzheimer's disease in randomized trials has been extremely small, due to the need to manage subscale scores of behavioral disturbance scale. Therefore, because a meta-analysis can increase the statistical power for group comparisons and can overcome the limitation of sample size in underpowered studies,[11] we conducted a meta-analysis to achieve conclusive evidence for the efficacy of memantine on individual behavioral disturbances in patients with Alzheimer's disease.

Methods

This meta-analysis was performed based upon the Preferred Reporting Items for Systematic Reviews and Meta-Analysis guidelines (International prospective register of systematic reviews [PROSPERO]: CRD42017059245).[12] We combined with the data from the studies of memantine monotherapy and the studies of combination therapy with memantine and cholinesterase inhibitors, because studies of the combination therapy included the patients who had several dementia symptoms at the baseline despite taking some cholinesterase inhibitors.

Search strategy and inclusion criteria

To identify relevant studies, two of the authors (TK and SM) independently searched MEDLINE, Cochrane library, Scopus, and PsycINFO without language restrictions from the inception of their databases to April 25, 2017, using the following search strategy: ("Alzheimer Disease" [Mesh] OR "Alzheimer disease" OR "Alzheimer's disease") AND ("Memantine"[Mesh] OR "memantine") AND ("randomized" OR "random" OR "randomly"). The authors also searched ClinicalTrials.gov (http://ClinicalTrials.gov/), ISRCTN registry (https://www.isrctn.com/), and the International Clinical Trials Registry Platform (http://www.who.int/ictrp/en/) to include randomized controlled trials as comprehensively as possible and to minimize the possibility of publication bias. Only randomized placebo- or usual care-controlled trials of memantine treatment in patients with Alzheimer's disease lasting ≥2 weeks were included. The studies that included more than 50% patients who received the combination therapy were classified as a combination therapy group in this study (Table 1). Two authors (TK and SM) independently assessed inclusion/exclusion criteria and selected the studies. The references of the included articles and review articles were also searched for citations of additional relevant published and unpublished studies, including conference abstracts.

Data synthesis and outcome measures

The primary outcomes were individual behavioral disturbances as follows: delusion, hallucination, agitation/aggression, dysphoria, anxiety/phobia, euphoria, apathy, disinhibition, irritability/lability, aberrant motor activity/activity disturbances, nighttime disturbance/diurnal rhythm disturbances, and eating disturbances. Nine of 11 studies included in the meta-analysis used Neuropsychiatric Inventory,[13] and the other two studies[14,15] used the Behavioral Pathology in Alzheimer's Disease Rating Scale.[16] For three-arm (memantine 10 mg/day arm, memantine 20 mg/day arm, and placebo arm) studies,[17] we combined the data of the

memantine 10 mg/day arm with that of memantine 20 mg/day. For four-arm (memantine monotherapy arm, combination therapy with memantine and donepezil arm, donepezil monotherapy arm, and placebo arm) studies,[18] we combined the data of the memantine monotherapy arm with that of the combination therapy with memantine (ie, memantine group) and donepezil arm and the data of donepezil monotherapy arm with that of placebo arm (ie, non-memantine group).

Data extraction

Two authors (TK and SM) independently extracted the data from the included studies. Where possible, we used intention-to-treat (ITT) or a full analysis set (FAS) population. When such data were unavailable, the results for observed case (OC) analysis were extracted from each study. When the data required for meta-analysis were missing, we contacted the investigators (or the industries) of the relevant study and requested unpublished data.

Meta-analysis methods

The meta-analysis was conducted using Review Manager software.[19] The random-effects model was selected for this meta-analysis due to the potential heterogeneity across studies. To evaluate these outcomes, SMD, with 95% CIs, based upon a random-effects model, was evaluated in the meta-analysis. We assessed the methodological quality of the trials, according to the Cochrane risk-of-bias criteria in the Cochrane Handbook.[11] Study heterogeneity was tested using the I^2 statistic, considering $I^2 \geq 50\%$ to reflect considerable heterogeneity.[11] We did not find considerable heterogeneity with respect to all meta-analysis. To detect the confounding factors for the result of primary outcomes for efficacy, two subgroup analysis (including a test for subgroup differences) were performed for the following: severity of disease (mild-to-moderate vs moderate and moderate-to-severe) and therapeutic strategy (memantine monotherapy vs combination therapy with memantine and cholinesterase inhibitors). Finally, we utilized funnel plots to explore potential publication bias.

Results
Study characteristics

Of the 2,239 results obtained in our literature search, we excluded the following: 1,498 as duplicates, 693 after a review of the abstract or title review, and 28 articles after a review of the full text (22 review articles, four single-arm studies, and two same studies). We did not retrieve 10 studies by searching through the review articles and clinical trial registries (Figure S1). Although 30 studies were identified though the

literature search, only 11 studies (memantine monotherapy vs placebo: four studies,[14,17,20,21] n=1,500; combination therapy with memantine and cholinesterase inhibitors vs cholinesterase inhibitors: seven studies,[15,18,22–26] n=2,761) were included in the meta-analysis, since the other 20 studies did not report any available data for performing a meta-analysis.

The main characteristics of studies and patients are summarized in Table 1. The mean duration of the studies was 26.5 weeks (one study was 52 weeks, other studies were 24 weeks), the mean patient age was 76.3 years, and the percentage of males was 34.6%. Although one of the 11 studies was an open-label study (ie, not placebo-controlled study),[22] the other 10 studies were double-blinded, randomized, placebo-controlled trials. One study was a memantine extended-release study.[23] The dose of memantine was 20 mg/day in all studies, other than Kitamura et al's[17] study (three arms: memantine 10 mg/day arm, memantine 20 mg/day arm, and placebo arm). The Howard et al's[18] study used OC populations in their analysis. Because this study was a four-arm study (memantine monotherapy arm, combination therapy with memantine and donepezil arm, donepezil monotherapy arm, and placebo arm),[18] we combined the data of memantine monotherapy arm with that of combination therapy with memantine (ie, memantine group) and donepezil arm and data of donepezil monotherapy arm with that of placebo arm (ie, non-memantine group). Two studies were not sponsored by a pharmaceutical company.[18,22] Most of all studies included in the study excluded the patients who had psychiatric disorders other than Alzheimer's disease.

Evaluations on the methodological quality of the included studies were performed based upon the Cochrane risk-of-bias criteria and are shown in Figures S2 and S3.

Results of the meta-analysis

Memantine showed significant improvement in agitation/aggression (SMD =−0.11; 95% CIs =−0.20, −0.03; P=0.01, I^2=47%; Figure 1), delusion (SMD =−0.12; 95% CIs =−0.18, −0.06; P=0.0002; I^2=0%; Figure 2), disinhibition (SMD =−0.08; 95% CIs =−0.15, −0.00; P=0.04; I^2=0%; Figure 3), and nighttime disturbance/diurnal rhythm disturbances (SMD =−0.10; 95% CIs =−0.18, −0.02; P=0.02; I^2=36%; Figure 4) compared to control. Memantine was also marginally superior to control in hallucination (SMD =−0.06; 95% CIs =−0.12, 0.01; P=0.07; I^2=0%; Figure 5) and irritability/lability (SMD =−0.09; 95% CIs =−0.19, 0.01; P=0.07; I^2=42%; Figure 6). Memantine is similar to control in aberrant motor activity/activity disturbances, anxiety/phobia, apathy, dysphoria, eating disturbances, and euphoria

Table 1 Characteristics of included randomized controlled trials

Study, country, sponsorship	Total (n)	Methods: 1. Study design 2. Duration 3. Analyzed population	Patients 1. Diagnosis 2. Inclusion criteria 3. Study-defined disease severity 4. Mental disorder comorbidities 5. Concomitant drug	Age mean ± SD, years
Monotherapy				
Kitamura et al,[17] Japan, industry	315	1. DB-RCT 2. 24 weeks 3. FAS	1. AD, DSM-IV, and NINCDS-ADRDA 2. Age ≥50 years, MMSE 5–14, FAST 6a–7a 3. Moderate to severe 4. NR 5. Not allowed concomitant drug use: AE, AP, APD, DON, MR, NMDARI, S/H, TD; allowed concomitant use within 2 weeks: BRO, LOR, RIL, TIA	73.3±9.4
Nakamura et al,[14] Japan, industry	432	1. DB-RCT 2. 24 weeks 3. FAS	1. AD, DSM-IV, and NINCDS-ADRDA 2. Age ≥50 years, MMSE 5–14, FAST 6a–7a 3. Moderate to severe 4. Exclusion: severe psychiatric disorder other than probable AD 5. Not allowed concomitant drug use: AE, AP, APD, DON, MR, NMDARI, S/H, TD; allowed concomitant use: BRO, LOR, RIL, TIA ≤150 mg/day	74.6±8.4
Peskind et al,[20] USA, industry	403	1. DB-RCT 2. 24 weeks 3. ITT	1. AD, NINCDS-ADRDA 2. Age ≥50 years, MMSE 10–22 3. Mild to moderate 4. Exclusion: psychiatric disorder other than probable AD 5. Allowed concomitant drug use: ADD, AH, AI, GB, GIN, OLA, RIS, TD, TOC	77.5
van Dyck et al,[21] USA, industry	350	1. DB-RCT 2. 24 weeks 3. ITT	1. AD, NINCDS-ADRDA 2. Age ≥50 years, MMSE 5–14 3. Moderate to severe 4. Exclusion: psychiatric disorder other than probable AD 5. Allowed concomitant drug use: AAPD, ADD, AH, AI, LAX, TD, TOC	78.2
Combination therapy				
Araki et al,[22] Japan, nonindustry	37	1. O-RCT 2. 24 weeks 3. FAS	1. AD, DSM-IV, and ICD-10 2. HDS-R 3–16 3. Moderate to severe 4. NR 5. NR	78.8±7.7
Grossberg et al,[23] international, industry	677	1. DB-RCT 2. 24 weeks 3. ITT	1. AD, DSM-IV-TR, and NINCDS-ADRDA 2. Age ≥50 years, MMSE 3–14 3. Moderate to severe 4. Exclusion: DSM-IV Axis I disorder other than AD 5. NR	76.5
Herrmann et al,[24] Canada, industry	369	1. DB-RCT 2. 24 weeks 3. FAS	1. AD, NINCDS-ADRDA 2. Age ≥50 years, MMSE 5–15, NPI ≥13, NPI agitation/aggression score ≥1 3. Moderate to severe 4. Exclusion: psychiatric disorder other than probable AD 5. Concomitant drug use: ADD 23.6%, ANX 3.3%, APD 22.2%	74.9

Male (%)	Race (%)	Baseline cognitive function scales (mean ± SD)	Intervention, dose (mg/day)	n	Efficacy outcomes[a]
29.3	Japanese: 100	MMSE: 10.1±3.0; SIB: 71.1±17.8	MEM 20 mg (Fi) MEM 10 mg (Fi) PLA	100 107 108	MEM > PLA: FAST (20 mg), MMSE (20 mg), **SIB** (20 mg); MEM = PLA: **ADCS-ADL19**, CIBIC-Plus, FAST (10 mg), MMSE (10 mg), NPI10, **SIB** (10 mg)
35.7	Japanese: 100	MMSE: 9.9±3.0; SIB: 71.0±17.9	MEM 20 mg (Fi) PLA	221 211	MEM > PLA: Behave-AD, **SIB**; MEM = PLA: **CIBIC-Plus**, FAST, MENFIS
41.2	Caucasian: 91.3, others: 8.7	ADAS-cog: 27.3; MMSE: 17.3	MEM 20 mg (Fi) PLA	201 202	MEM > PLA: **ADAS-cog**, CIBIC-Plus, NPI12; MEM = PLA: ADCS-ADL23
28.6	Caucasian: 80.9; others: 19.1	MMSE: 10.1; SIB: 76.4	MEM 20 mg (Fi)	178	MEM = PLA: **ADCS-ADL19**, BGP, CIBIC-Plus, FAST, NPI12, **SIB**
48.6	Japanese: 100	MMSE: 16.1	MEM 20 mg (Fi) + DON (100%, NR) DON (100%, NR)	19 18	MEM + DON > DON: CDT, **CGI-I**, **MMSE**, **NPI10**, **ZBI**; MEM + DON = DON: NIRS (mean of all channels)
28.0	Caucasian: 94.1; others: 5.9	MMSE: 10.8; SIB: 76.0	MEM-ER 28 mg (Fi) + ChEIs (DON [69%, 8.0 mg], GAL [21%, 13.5 mg], RIV [9%, 6.8 mg]) PLA + ChEIs (DON [63%, 7.8 mg], GAL [20%, 13.5 mg], RIV [12%, 6.8 mg])	342 335	MEM (ER) + ChEIs > PLA + ChEIs: **CIBIC-Plus**, NPI12, **SIB**, VFT; MEM (ER) + ChEIs = PLA + ChEIs: ADCS-ADL19
41.7	NR	MMSE: 11.8; SIB: 82.1	MEM 20 mg (Fi) + ChEIs (combination therapy 95%) PLA + ChEIs (combination therapy 97%)	182 187	MEM + ChEIs = PLA + ChEIs: ADCS-ADL19, CIBIC-Plus, CMAI, **NPI12**, **SIB**

(Continued)

Table 1 (*Continued*)

Study, country, sponsorship	Total (n)	Methods: 1. Study design 2. Duration 3. Analyzed population	Patients 1. Diagnosis 2. Inclusion criteria 3. Study-defined disease severity 4. Mental disorder comorbidities 5. Concomitant drug	Age mean ± SD, years
Howard et al,[18] UK, nonindustry	295	1. DB-RCT 2. 52 weeks 3. OC	1. AD, NINCDS-ADRDA 2. Age ≥50 years, MMSE 5–13 3. Moderate to severe 4. NR 5. NR	77.1±8.4
Nakamura et al,[15] Japan, industry	546	1. DB-RCT 2. 24 weeks 3. FAS	1. AD, DSM-IV-TR, and NINCDS-ADRDA 2. Age ≥50 years, MMSE 1–14, SIB 30–85 3. Moderate to severe 4. Exclusion: severe psychiatric disorder other than probable AD 5. Not allowed concomitant use: AP, APD, CD, GAL, MR, NMDARI, RIV, S/H, TD; allowed concomitant drug use: BRO, ESZ, LOR, RAM, RIL, SUV, TIA, ZOP	78.5
Porsteinsson et al,[25] USA, industry	433	1. DB-RCT 2. 24 weeks 3. ITT	1. AD, NINCDS-ADRDA 2. Age ≥50 years, MMSE 10–22 3. Mild to moderate 4. Exclusion: psychiatric disorder other than probable AD 5. NR	75.4
Tariot et al,[26] USA, industry	404	1. DB-RCT 2. 24 weeks 3. ITT	1. AD, NINCDS-ADRDA 2. Age ≥50 years, MMSE 5–14 3. Moderate to severe 4. Exclusion: psychiatric disorder other than probable AD 5. Concomitant drug use: ACE 37.0%, ASC 19.4%, CAL 11.4%, GB 13.6%, MV 39.2%, PAR 14.1%, TOC 62.3%	75.5

Note: [a]Primary outcomes in each study are given in bold.
Abbreviations: AA, African-American; AAPD, atypical antipsychotic drugs; ACE, acetylsalicylic acid; AD, Alzheimer disease; ADAS-cog, Alzheimer's Disease Assessment Scale-Cognitive Subscale; ADCS-ADL, Alzheimer's Disease Cooperative Study-Activities of Daily Living; ADD, antidepressant drugs; AE, antiepileptics; AH, antihypertensives; AI, anti-inflammatories; ANX, anxiolytics; APD, antipsychotic drugs; AP, anti-Parkinson; ASC, ascorbic acid; BADLS, Bristol Activities of Daily Living Scale; Behave-AD, Behavioral Pathology in Alzheimer's Disease Rating Scale; BGP, Behavioral Rating Scale for Geriatric Patients; BRO, brotizolam; CAL, calcium; CD, cholinergic drugs; CDT, clock drawing test; CGBRS, Crichton Geriatric Behavioral Rating Scale; CGI-I, Clinical Global Impression-Improvement scale; ChEI, cholinesterase inhibitors; CIBIC-Plus, Clinician's Interview-based Impression of Change Plus Caregiver Input; CMAI, Cohen-Mansfield Agitation Inventory; DB-RCT, double-blind randomized controlled trial; DON, donepezil; DSM-IV, *Diagnostic and Statistical Manual of Mental Disorders*, fourth edition; TR, Text Revision; ER, extended release; ESZ, eszopiclone; FAS, full analysis set; FAST, functional assessment staging instrument; Fi, fixed dose; GAL, galantamine; GB, *Ginkgo biloba*; GHQ-12, General Health Questionnaire 12; GIN, ginseng; HDS-R, Hasegawa's Dementia Scale-Revision; ICD-10, International Classification Of Diseases, 10th edition; ITT, intention to treat; LAX, laxatives; LOR, lormetazepam; MEM, memantine; MENFIS, Mental Function Impairment Scale; MMSE, mini-mental state examination; MR, muscle relaxant; MV, multi-vitamins; n, number of patients; NINCDS-ADRDA, National Institute of Neurological and Communicative Disorders and Stroke and the Alzheimer's Disease and Related Disorders Association; NIRS, near-infrared spectroscopy; NMDARI, *N*-methyl-D-aspartate receptor inhibitor; NPI, Neuropsychiatric Inventory; NR, not reported; OC, observed case; OLA, olanzapine; O-RCT, open-label randomized controlled trial; PAR, paracetamol; PLA, placebo; RAM, ramelteon; RIL, rilmazafone; RIS, risperidone; RIV, rivastigmine; SIB, severe impairment battery; S/H, sedatives/hypnotics; SUV, suvorexant; TD, thiazide diuretics; TIA, tiapride; TOC, tocopherol; VFT, verbal fluency test; ZBI, Zarit Burden Interview; ZOP, zopiclone.

(Figures 7–12). The data for individual behavioral disturbances scores were simulated with no publication bias.

Subgroup analysis divided by therapeutic strategy

We did not find considerable heterogeneity with respect to all meta-analysis (Figures 1–12). We also did not find any significant subgroup differences in all subgroup analysis.

Delusion was the outcome, where memantine was superior to control in the monotherapy subgroup and the combination therapy subgroup (Figure 2). Agitation/aggression and disinhibition were the outcomes, where memantine was superior to control in the combination therapy subgroup but not in the monotherapy subgroup (Figures 1 and 3).

Subgroup analysis divided by the severity of disease

We also did not find considerable heterogeneity with respect to all meta-analysis (Figures 1–12). We also did not find any significant subgroup differences in all subgroup analysis. Although we found marginally subgroup differences in subgroup analysis divided by the severity of disease with

Male (%)	Race (%)	Baseline cognitive function scales (mean ± SD)	Drug, dose (mg/day)	n	Efficacy outcomes[a]
35	Caucasian: 95; AA: 3; others: 2	MMSE: 9.1±2.6	MEM 20 mg (Fi) + DON (50%, 10 mg)	149	MEM + DON = PLA + DON: **BADLS**, DEMQOL-proxy, GHQ-12, **MMSE**, NPI12
			PLA + DON (50%, 10 mg)	146	
27.2	Japanese: 100	MMSE: 10.8; SIB: 77.0	MEM 20 mg (Fi) + DON (100%, 6.9 mg)	273	MEM + DON = PLA + DON: Behave-AD, CGBRS, **SIB**
			PLA + DON (100%, 6.9 mg)	273	
47.8	NR	ADAS-cog: 27.4; MMSE: 16.8	MEM 20 mg (Fi) + ChEIs (DON [71%, 9.5 mg], GAL [14%, 19.7 mg], RIV [15%, 9.2 mg])	217	MEM + ChEIs = PLA + ChEIs: **ADAS-cog**, **CIBIC-Plus**, ADCS-ADL, NPI12, MMSE
			PLA + ChEIs (DON [63%, 8.9 mg], GAL [16%, 19.4 mg], RIV [20%, 10.0 mg])	216	
35.0	Caucasian: 91.3; others: 8.7	MMSE: 10.0; SIB: 79.0	MEM 20 mg (Fi) + DON (100%, 9.3 mg)	203	MEM + DON > PLA + DON: **ADCS-ADL**, BGP, CIBIC-Plus, NPI12, **SIB**
			PLA + DON (100%, 9.5 mg)	201	

Therapeutic strategy

Study or subgroup	Memantine Mean	SD	Total	Control Mean	SD	Total	Weight (%)	SMD IV, random, 95% CI	SMD IV, random, 95% CI
Monotherapy									
Kitamura et al (2011)[17]	0.003	2.06	207	−0.33	2.13	107	8.5	0.16 (−0.07, 0.39)	
Nakamura et al (2011)[14]	−0.05	1.19	217	0.26	1.55	208	10.5	−0.22 (−0.42, −0.03)	
Peskind et al (2006)[20]	0	1.3	191	0.1	1.63	190	10.0	−0.07 (−0.27, 0.13)	
van Dyck et al (2007)[21]	−0.1	2.9	161	0.3	2.8	154	9.0	−0.14 (−0.36, 0.08)	
Subtotal (95% CI)			776			659	37.9	−0.08 (−0.23, 0.08)	
Heterogeneity: τ^2=0.01; χ^2=6.52, df=3 (P=0.09); I^2=54%									
Test for overall effect: Z=0.96 (P=0.34)									
Combination therapy									
Araki et al (2014)[22]	0.42	1.16	12	3.08	3.15	13	1.0	−1.07 (−1.91, −0.22)	
Grossberg et al (2013)[23]	−0.4	2.44	318	0	2.5	321	12.5	−0.16 (−0.32, −0.01)	
Herrmann et al (2013)[24]	−1.5	3.16	159	−1.71	2.91	165	9.2	0.07 (−0.15, 0.29)	
Howard et al (2012)[18]	0.36697248	3.1140677	109	1.1743119	2.9933384	109	7.2	−0.26 (−0.53, 0.00)	
Nakamura et al (2016)[15]	−0.1	1.31	267	0.1	1.43	267	11.7	−0.15 (−0.32, 0.02)	
Porsteinsson et al (2008)[25]	0	1.97	212	0	1.97	209	10.5	0.00 (−0.19, 0.19)	
Tariot et al (2004)[26]	−0.1	2.15	193	0.5	2.83	189	10.0	−0.24 (−0.44, −0.04)	
Subtotal (95% CI)			1,270			1,273	62.1	−0.14 (−0.25, −0.02)	
Heterogeneity: τ^2=0.01; χ^2=11.94, df=6 (P=0.06); I^2=50%									
Test for overall effect: Z=2.31 (P=0.02)									
Total (95% CI)			2,046			1,932	100	−0.11 (−0.20, −0.03)	
Heterogeneity: τ^2=0.01; χ^2=18.90, df=10 (P=0.04); I^2=47%									
Test for overall effect: Z=2.52 (P=0.01)									
Test for subgroup differences: χ^2=0.38, df=1 (P=0.54); I^2=0%									

−2 −1 0 1 2
Favors (memantine)   Favors (control)

Figure 1 (Continued)

Severity of disease

Study or subgroup	Memantine Mean	SD	Total	Control Mean	SD	Total	Weight (%)	SMD IV, random, 95% CI
Mild-to-moderate								
Peskind et al (2006)[20]	0	1.3	191	0.1	1.63	190	10.0	−0.07 (−0.27, 0.13)
Porsteinsson et al (2008)[25]	0	1.97	212	0	1.97	209	10.5	0.00 (−0.19, 0.19)
Subtotal (95% CI)			403			399	20.5	−0.03 (−0.17, 0.11)
Heterogeneity: τ²=0.00; χ²=0.23, df=1 (P=0.63); I²=0%								
Test for overall effect: Z=0.46 (P=0.65)								
Moderate-to-severe								
Araki et al (2014)[22]	0.42	1.16	12	3.08	3.15	13	1.0	−1.07 (−1.91, −0.22)
Grossberg et al (2013)[23]	−0.4	2.44	318	0	2.5	321	12.5	−0.16 (−0.32, −0.01)
Herrmann et al (2013)[24]	−1.5	3.16	159	−1.71	2.91	165	9.2	0.07 (−0.15, 0.29)
Howard et al (2012)[18]	0.36697248	3.1140677	109	1.1743119	2.9933384	109	7.2	−0.26 (−0.53, 0.00)
Kitamura et al (2011)[17]	0.003	2.06	207	−0.33	2.13	107	8.5	0.16 (−0.07, 0.39)
Nakamura et al (2011)[14]	−0.05	1.19	217	0.26	1.55	208	10.5	−0.22 (−0.42, −0.03)
Nakamura et al (2016)[15]	−0.1	1.13	267	0.1	1.43	267	11.7	−0.15 (−0.32, 0.02)
Tariot et al (2004)[26]	−0.1	2.15	193	0.5	2.83	189	10.0	−0.24 (−0.44, −0.04)
van Dyck et al (2007)[21]	−0.1	2.9	161	0.3	2.8	154	9.0	−0.14 (−0.36, 0.08)
Subtotal (95% CI)			1,643			1,533	79.5	−0.14 (−0.24, −0.03)
Heterogeneity: τ²=0.01; χ²=16.98, df=8 (P=0.03); I²=53%								
Test for overall effect: Z=2.49 (P=0.01)								
Total (95% CI)			2,046			1,932	100	−0.11 (−0.20, −0.03)
Heterogeneity: τ²=0.01; χ²=18.90, df=10 (P=0.04); I²=47%								
Test for overall effect: Z=2.52 (P=0.01)								
Test for subgroup differences: χ²=1.36, df=1 (P=0.24); I²=26.2%								

SMD IV, random, 95% CI — scale: −2 −1 0 1 2
Favors (memantine)   Favors (control)

Figure 1 Forest plot of agitation/aggression scores.
Abbreviations: 95% CI, 95% confidence interval; IV, inverse variance; SD, standard deviation; SMD, standardized mean difference.

Therapeutic strategy

Study or subgroup	Memantine Mean	SD	Total	Control Mean	SD	Total	Weight (%)	SMD IV, random, 95% CI
Monotherapy								
Kitamura et al (2011)[17]	−0.246	1.679	207	0.07	1.9	107	7.2	−0.18 (−0.41, 0.05)
Nakamura et al (2011)[14]	−0.15	1.68	217	0.13	1.9	208	10.8	−0.16 (−0.35, 0.03)
Peskind et al (2006)[20]	−0.1	1.27	191	0.2	1.36	190	9.7	−0.23 (−0.43, −0.03)
van Dyck et al (2007)[21]	−0.2	3.4	161	0.2	3	154	8.0	−0.12 (−0.35, 0.10)
Subtotal (95% CI)			776			659	35.7	−0.17 (−0.28, −0.07)
Heterogeneity: τ²=0.00; χ²=0.50, df=3 (P=0.92); I²=0%								
Test for overall effect: Z=3.23 (P=0.001)								
Combination therapy								
Araki et al (2014)[22]	0	0	12	2.69	2.14	13		Not estimable
Grossberg et al (2013)[23]	−0.4	2.47	318	−0.1	2.4	321	16.3	−0.12 (−0.28, 0.03)
Herrmann et al (2013)[24]	−0.65	2.85	159	−0.5	2.67	165	8.3	−0.05 (−0.27, 0.16)
Howard et al (2012)[18]	−0.27522936	2.9560721	109	−0.11009174	2.9006367	109	5.6	−0.06 (−0.32, 0.21)
Nakamura et al (2016)[15]	−0.1	1.16	267	0.1	1.5	267	13.6	−0.15 (−0.32, 0.02)
Porsteinsson et al (2008)[25]	−0.1	1.4	212	−0.1	1.64	209	10.8	0.00 (−0.19, 0.19)
Tariot et al (2004)[26]	0.2	2.02	193	0.4	2.55	189	9.8	−0.09 (−0.29, 0.11)
Subtotal (95% CI)			1,270			1,273	64.3	−0.09 (−0.17, −0.01)
Heterogeneity: τ²=0.00; χ²=1.65, df=5 (P=0.89); I²=0%								
Test for overall effect: Z=2.20 (P=0.03)								
Total (95% CI)			2,046			1,932	100	−0.12 (−0.18, −0.06)
Heterogeneity: τ²=0.00; χ²=3.78, df=9 (P=0.93); I²=0%								
Test for overall effect: Z=3.70 (P=0.0002)								
Test for subgroup differences: χ²=1.63, df=1 (P=0.20); I²=38.6%								

SMD IV, random, 95% CI — scale: −0.5 −0.25 0 0.25 0.5
Favors (memantine)   Favors (control)

Severity of disease

Study or subgroup	Memantine Mean	SD	Total	Control Mean	SD	Total	Weight (%)	SMD IV, random, 95% CI
Mild-to-moderate								
Peskind et al (2006)[20]	−0.1	1.27	191	0.2	1.36	190	9.7	−0.23 (−0.43, −0.03)
Porsteinsson et al (2008)[25]	−0.1	1.4	212	−0.1	1.64	209	10.8	0.00 (−0.19, 0.19)
Subtotal (95% CI)			403			399	20.4	−0.11 (−0.33, 0.11)
Heterogeneity: τ²=0.02; χ²=2.58, df=1 (P=0.11); I²=61%								
Test for overall effect: Z=0.98 (P=0.33)								
Moderate-to-severe								
Araki et al (2014)[22]	0	0	12	2.69	2.14	13		Not estimable
Grossberg et al (2013)[23]	−0.4	2.47	318	−0.1	2.4	321	16.3	−0.12 (−0.28, 0.03)
Herrmann et al (2013)[24]	−0.65	2.85	159	−0.5	2.67	165	8.3	−0.05 (−0.27, 0.16)
Howard et al (2012)[18]	−0.27522936	2.9560721	109	−0.11009174	2.9006367	109	5.6	−0.06 (−0.32, 0.21)
Kitamura et al (2011)[17]	−0.246	1.679	207	0.07	1.9	107	7.2	−0.18 (−0.41, 0.05)
Nakamura et al (2011)[14]	−0.15	1.68	217	0.13	1.9	208	10.8	−0.16 (−0.35, 0.03)
Nakamura et al (2016)[15]	−0.1	1.16	267	0.1	1.5	267	13.6	−0.15 (−0.32, 0.02)
Tariot et al (2004)[26]	0.2	2.02	193	0.4	2.55	189	9.8	−0.09 (−0.29, 0.11)
van Dyck et al (2007)[21]	−0.2	3.4	161	0.2	3	154	8.0	−0.12 (−0.35, 0.10)
Subtotal (95% CI)			1,643			1,533	79.6	−0.12 (−0.19, −0.05)
Heterogeneity: τ²=0.00; χ²=1.18, df=7 (P=0.99); I²=0%								
Test for overall effect: Z=3.37 (P=0.0007)								
Total (95% CI)			2,046			1,932	100	−0.12 (−0.18, −0.06)
Heterogeneity: τ²=0.00; χ²=3.78, df=9 (P=0.93); I²=0%								
Test for overall effect: Z=3.70 (P=0.0002)								
Test for subgroup differences: χ²=0.01, df=1 (P=0.94); I²=0%								

SMD IV, random, 95% CI — scale: −0.5 −0.25 0 0.25 0.5
Favors (memantine)   Favors (control)

Figure 2 Forest plot of delusion scores.
Abbreviations: 95% CI, 95% confidence interval; IV, inverse variance; SD, standard deviation; SMD, standardized mean difference.

Therapeutic strategy

Study or subgroup	Memantine Mean	SD	Total	Control Mean	SD	Total	Weight (%)	SMD IV, random, 95% CI	SMD IV, random, 95% CI
Monotherapy									
Kitamura et al (2011)[17]	0.0207	1.881	207	0.36	1.77	107	9.4	−0.18 (−0.42, 0.05)	
Peskind et al (2006)[20]	0	1.01	191	−0.1	1.02	190	12.8	0.10 (−0.10, 0.30)	
van Dyck et al (2007)[21]	0	2.3	161	0.2	2.2	154	10.6	−0.09 (−0.31, 0.13)	
Subtotal (95% CI)			559			451	32.7	−0.05 (−0.21, 0.12)	

Heterogeneity: τ²=0.01; χ²=3.45, df=2 (P=0.18); I²=42%
Test for overall effect: Z=0.59 (P=0.56)

Combination therapy									
Araki et al (2014)[22]	−0.08	0.29	12	1.54	2.67	13	0.8	−0.81 (−1.63, 0.01)	
Grossberg et al (2013)[23]	−0.2	1.85	318	0	1.71	321	21.4	−0.11 (−0.27, 0.04)	
Herrmann et al (2013)[24]	−0.39	2.28	159	−0.1	2.12	165	10.8	−0.13 (−0.35, 0.09)	
Howard et al (2012)[18]	0.05504587	2.5813966	109	0.12844037	2.2073149	109	7.3	−0.03 (−0.30, 0.24)	
Porsteinsson et al (2008)[25]	0	1.49	212	0.1	1.6	209	14.1	−0.06 (−0.26, 0.13)	
Tariot et al (2004)[26]	0	2.01	193	0.1	1.81	189	12.8	−0.05 (−0.25, 0.15)	
Subtotal (95% CI)			1,003			1,006	67.3	−0.09 (−0.18, −0.01)	

Heterogeneity: τ²=0.00; χ²=3.54, df=5 (P=0.62); I²=0%
Test for overall effect: Z=2.08 (P=0.04)

| Total (95% CI) | | | 1,562 | | | 1,457 | 100 | −0.08 (−0.15, −0.00) | |

Heterogeneity: τ²=0.00; χ²=7.40, df=8 (P=0.49); I²=0%
Test for overall effect: Z=2.09 (P=0.04)
Test for subgroup differences: χ²=0.21, df=1 (P=0.65); I²=0%

−1 −0.5 0 0.5 1
Favors (memantine) Favors (control)

Severity of disease

Study or subgroup	Memantine Mean	SD	Total	Control Mean	SD	Total	Weight (%)	SMD IV, random, 95% CI	SMD IV, random, 95% CI
Mild-to-moderate									
Peskind et al (2006)[20]	0	1.01	191	−0.1	1.02	190	12.8	0.10 (−0.10, 0.30)	
Porsteinsson et al (2008)[25]	0	1.49	212	0.1	1.6	209	14.1	−0.06 (−0.26, 0.13)	
Subtotal (95% CI)			403			399	26.9	0.01 (−0.15, 0.17)	

Heterogeneity: τ²=0.00; χ²=1.33, df=1 (P=0.25); I²=25%
Test for overall effect: Z=0.17 (P=0.87)

Moderate-to-severe									
Araki et al (2014)[22]	−0.08	0.29	12	1.54	2.67	13	0.8	−0.81 (−1.63, 0.01)	
Grossberg et al (2013)[23]	−0.2	1.85	318	0	1.71	321	21.4	−0.11 (−0.27, 0.04)	
Herrmann et al (2013)[24]	−0.39	2.28	159	−0.1	2.12	165	10.8	−0.13 (−0.35, 0.09)	
Howard et al (2012)[18]	0.05504587	2.5813966	109	0.12844037	2.2073149	109	7.3	−0.03 (−0.30, 0.24)	
Kitamura et al (2011)[17]	0.0207	1.881	207	0.36	1.77	107	9.4	−0.18 (−0.42, 0.05)	
Tariot et al (2004)[26]	0	2.01	193	0.1	1.81	189	12.8	−0.05 (−0.25, 0.15)	
van Dyck et al (2007)[21]	0	2.3	161	0.2	2.2	154	10.6	−0.09 (−0.31, 0.13)	
Subtotal (95% CI)			1,159			1,058	73.1	−0.11 (−0.19, −0.03)	

Heterogeneity: τ²=0.00; χ²=3.89, df=6 (P=0.69); I²=0%
Test for overall effect: Z=2.55 (P=0.01)

| Total (95% CI) | | | 1,562 | | | 1,457 | 100 | −0.08 (−0.15, −0.00) | |

Heterogeneity: τ²=0.00; χ²=7.40, df=8 (P=0.49); I²=0%
Test for overall effect: Z=2.09 (P=0.04)
Test for subgroup differences: χ²=1.79, df=1 (P=0.18); I²=44.3%

−1 −0.5 0 0.5 1
Favors (memantine) Favors (control)

Figure 3 Forest plot of disinhibition scores.
Abbreviations: 95% CI, 95% confidence interval; IV, inverse variance; SD, standard deviation; SMD, standardized mean difference.

Therapeutic strategy

Study or subgroup	Memantine Mean	SD	Total	Control Mean	SD	Total	Weight (%)	SMD IV, random, 95% CI	SMD IV, random, 95% CI
Monotherapy									
Nakamura et al (2011)[14]	0	0.53	217	0.06	0.61	208	11.7	−0.10 (−0.30, 0.09)	
Peskind et al (2006)[20]	0.1	1.05	191	0.2	1.25	190	10.9	−0.09 (−0.29, 0.11)	
van Dyck et al (2007)[21]	−0.2	3.1	161	−0.1	2.9	154	9.6	−0.03 (−0.25, 0.19)	
Subtotal (95% CI)			569			552	32.2	−0.08 (−0.20, 0.04)	

Heterogeneity: τ²=0.00; χ²=0.24, df=2 (P=0.89); I²=0%
Test for overall effect: Z=1.31 (P=0.19)

Combination therapy									
Grossberg et al (2013)[23]	−0.5	2.67	317	0.1	2.83	321	14.8	−0.22 (−0.37, −0.06)	
Herrmann et al (2013)[24]	−0.36	3.19	159	−0.12	3.11	165	9.8	−0.08 (−0.29, 0.14)	
Howard et al (2012)[18]	0.09174312	3.5159459	109	0.83486239	3.9639901	109	7.3	−0.20 (−0.46, 0.07)	
Nakamura et al (2016)[15]	−0.1	0.57	267	0	0.6	267	13.4	−0.17 (−0.34, −0.00)	
Porsteinsson et al (2008)[25]	0.4	2.32	212	0	2.05	209	11.6	0.18 (−0.01, 0.37)	
Tariot et al (2004)[26]	0.2	2.57	193	0.6	2.48	189	10.9	−0.16 (−0.36, 0.04)	
Subtotal (95% CI)			1,257			1,260	67.8	−0.11 (−0.23, 0.02)	

Heterogeneity: τ²=0.01; χ²=12.01, df=5 (P=0.03); I²=58%
Test for overall effect: Z=1.68 (P=0.09)

| Total (95% CI) | | | 1,826 | | | 1,812 | 100 | −0.10 (−0.18, −0.02) | |

Heterogeneity: τ²=0.01; χ²=12.47, df=8 (P=0.13); I²=36%
Test for overall effect: Z=2.33 (P=0.02)
Test for subgroup differences: χ²=0.10, df=1 (P=0.75); I²=0%

−0.5 −0.25 0 0.25 0.5
Favors (memantine) Favors (control)

Figure 4 (*Continued*)

Severity of disease

Study or subgroup	Memantine Mean	SD	Total	Control Mean	SD	Total	Weight (%)	SMD IV, random, 95% CI	SMD IV, random, 95% CI
Mild-to-moderate									
Peskind et al (2006)[20]	0.1	1.05	191	0.2	1.25	190	10.9	−0.09 (−0.29, 0.11)	
Porsteinsson et al (2008)[25]	0.4	2.32	212	0	2.05	209	11.6	0.18 (−0.01, 0.37)	
Subtotal (95% CI)			403			399	22.5	0.05 (−0.21, 0.31)	
Heterogeneity: τ²=0.03; χ²=3.60, df=1 (P=0.06); I²=72%									
Test for overall effect: Z=0.37 (P=0.71)									
Moderate-to-severe									
Grossberg et al (2013)[23]	−0.5	2.67	317	0.1	2.83	321	14.8	−0.22 (−0.37, −0.06)	
Herrmann et al (2013)[24]	−0.36	3.19	159	−0.12	3.11	165	9.8	−0.08 (−0.29, 0.14)	
Howard et al (2012)[18]	0.09174312	3.5159459	109	0.83486239	3.9639901	109	7.3	−0.20 (−0.46, 0.07)	
Nakamura et al (2011)[14]	0	0.53	217	0.06	0.61	208	11.7	−0.10 (−0.30, 0.09)	
Nakamura et al (2016)[15]	−0.1	0.57	267	0	0.6	267	13.4	−0.17 (−0.34, −0.00)	
Tariot et al (2004)[26]	0.2	2.57	193	0.6	2.48	189	10.9	−0.16 (−0.36, 0.04)	
van Dyck et al (2007)[21]	−0.2	3.1	161	−0.1	2.9	154	9.6	−0.03 (−0.25, 0.19)	
Subtotal (95% CI)			1,423			1,413	77.5	−0.15 (−0.22, −0.07)	
Heterogeneity: τ²=0.00; χ²=2.63, df=6 (P=0.85); I²=0%									
Test for overall effect: Z=3.87 (P=0.0001)									
Total (95% CI)			1,826			1,812	100	−0.10 (−0.18, −0.02)	
Heterogeneity: τ²=0.01; χ²=12.47, df=8 (P=0.13); I²=36%									
Test for overall effect: Z=2.33 (P=0.02)									
Test for subgroup differences: χ²=1.96, df=1 (P=0.16); I²=49.0%									

−0.5   −0.25   0   0.25   0.5
Favors (memantine)   Favors (control)

Figure 4 Forest plot of nighttime disturbance/diurnal rhythm disturbance scores.
Abbreviations: 95% CI, 95% confidence interval; IV, inverse variance; SD, standard deviation; SMD, standardized mean difference.

Therapeutic strategy

Study or subgroup	Memantine Mean	SD	Total	Control Mean	SD	Total	Weight (%)	SMD IV, random, 95% CI	SMD IV, random, 95% CI
Monotherapy									
Kitamura et al (2011)[17]	−0.109	1.489	207	0.15	1.36	107	7.1	−0.18 (−0.41, 0.06)	
Nakamura et al (2011)[14]	0	0.93	217	0.05	1.19	208	10.8	−0.05 (−0.24, 0.14)	
Peskind et al (2006)[20]	0	1.02	191	0.2	1.07	190	9.6	−0.19 (−0.39, 0.01)	
van Dyck et al (2007)[21]	0	2.1	161	0	2	154	8.0	0.00 (−0.22, 0.22)	
Subtotal (95% CI)			776			659	35.5	−0.10 (−0.21, 0.00)	
Heterogeneity: τ²=0.00; χ²=2.30, df=3 (P=0.51); I²=0%									
Test for overall effect: Z=1.90 (P=0.06)									
Combination therapy									
Araki et al (2014)[22]	0.17	0.39	12	1.85	2.48	13	0.6	−0.90 (−1.73, −0.07)	
Grossberg et al (2013)[23]	0	1.83	318	−0.1	1.89	321	16.2	0.05 (−0.10, 0.21)	
Herrmann et al (2013)[24]	0.03	2.43	159	0.08	2.12	165	8.2	−0.02 (−0.24, 0.20)	
Howard et al (2012)[18]	0.17431193	1.8799332	109	0.53211009	2.692819	109	5.5	−0.15 (−0.42, 0.11)	
Nakamura et al (2016)[15]	0.1	0.73	267	0.1	0.81	267	3.6	0.00 (−0.17, 0.17)	
Porsteinsson et al (2008)[25]	0.1	1.47	212	0.2	1.16	209	10.7	−0.08 (−0.27, 0.12)	
Tariot et al (2004)[26]	0.1	1.59	193	0.2	1.49	189	9.7	−0.06 (−0.27, 0.14)	
Subtotal (95% CI)			1,270			1,273	64.5	−0.04 (−0.12, 0.05)	
Heterogeneity: τ²=0.00; χ²=6.59, df=6 (P=0.36); I²=9%									
Test for overall effect: Z=0.84 (P=0.40)									
Total (95% CI)			2,046			1,932	100	−0.06 (−0.12, 0.01)	
Heterogeneity: τ²=0.00; χ²=9.98, df=10 (P=0.44); I²=0%									
Test for overall effect: Z=1.79 (P=0.07)									
Test for subgroup differences: χ²=0.95, df=1 (P=0.33); I²=0%									

−2   −1   0   1   2
Favors (memantine)   Favors (control)

Severity of disease

Study or subgroup	Memantine Mean	SD	Total	Control Mean	SD	Total	Weight (%)	SMD IV, random, 95% CI	SMD IV, random, 95% CI
Mild-to-moderate									
Peskind et al (2006)[20]	0	1.02	191	0.2	1.07	190	9.6	−0.19 (−0.39, 0.01)	
Porsteinsson et al (2008)[25]	0.1	1.47	212	0.2	1.16	209	10.7	−0.08 (−0.27, 0.12)	
Subtotal (95% CI)			403			399	20.3	−0.13 (−0.27, 0.01)	
Heterogeneity: τ²=0.00; χ²=0.67, df=1 (P=0.41); I²=0%									
Test for overall effect: Z=1.84 (P=0.07)									
Moderate-to-severe									
Araki et al (2014)[22]	0.17	0.39	12	1.85	2.48	13	0.6	−0.90 (−1.73, −0.07)	
Grossberg et al (2013)[23]	0	1.83	318	−0.1	1.89	321	16.2	0.05 (−0.10, 0.21)	
Herrmann et al (2013)[24]	0.03	2.43	159	0.08	2.12	165	8.2	−0.02 (−0.24, 0.20)	
Howard et al (2012)[18]	0.17431193	1.8799332	109	0.5321009	2.692819	109	5.5	−0.15 (−0.42, 0.11)	
Kitamura et al (2011)[17]	−0.109	1.489	207	0.15	1.36	107	7.1	−0.18 (−0.41, 0.06)	
Nakamura et al (2011)[14]	0	0.93	217	0.05	1.19	208	10.8	−0.05 (−0.24, 0.14)	
Nakamura et al (2016)[15]	0.1	0.73	267	0.1	0.81	267	13.6	0.00 (−0.17, 0.17)	
Tariot et al (2004)[26]	0.1	1.59	193	0.2	1.49	189	9.7	−0.06 (−0.27, 0.14)	
van Dyck et al (2007)[21]	0	2.1	161	0	2	154	8.0	0.00 (−0.22, 0.22)	
Subtotal (95% CI)			1,643			1,533	79.7	−0.04 (−0.11, 0.03)	
Heterogeneity: τ²=0.00; χ²=7.97, df=8 (P=0.44); I²=0%									
Test for overall effect: Z=1.08 (P=0.28)									
Total (95% CI)			2,046			1,932	100	−0.06 (−0.12, 0.01)	
Heterogeneity: τ²=0.00; χ²=9.98, df=10 (P=0.44); I²=0%									
Test for overall effect: Z=1.79 (P=0.07)									
Test for subgroup differences: χ²=1.34, df=1 (P=0.25); I²=25.2%									

−2   −1   0   1   2
Favors (memantine)   Favors (control)
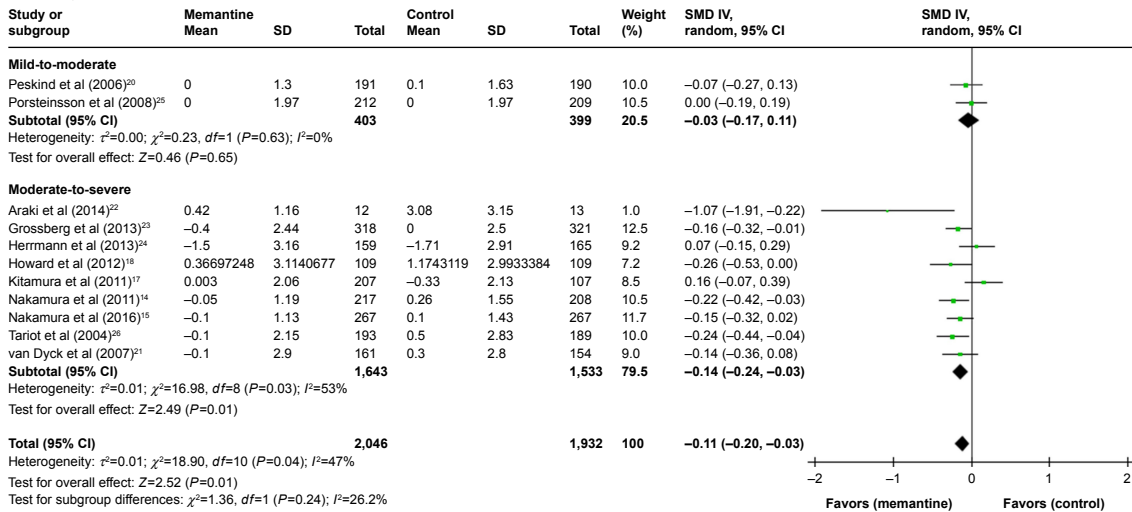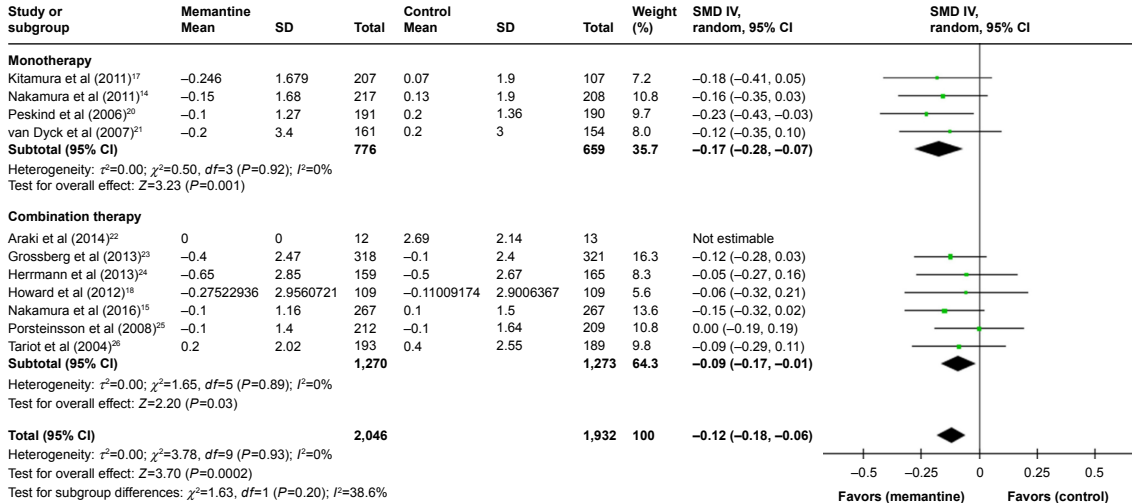
Figure 5 Forest plot of hallucination scores.
Abbreviations: 95% CI, 95% confidence interval; IV, inverse variance; SD, standard deviation; SMD, standardized mean difference.
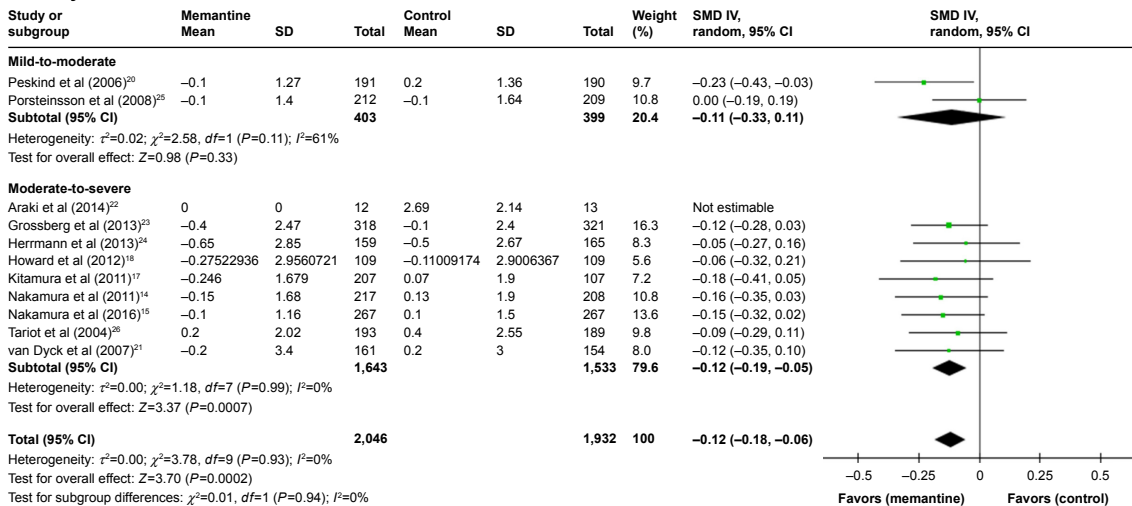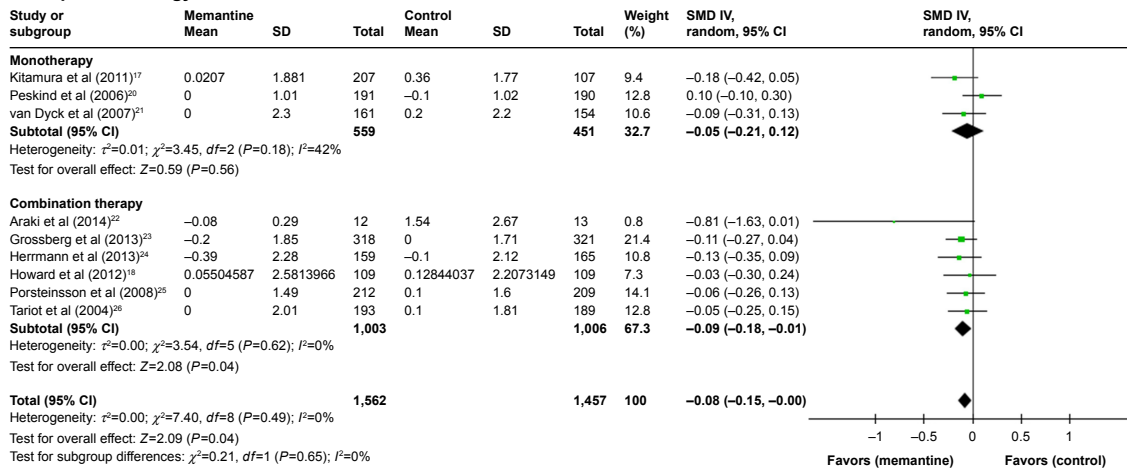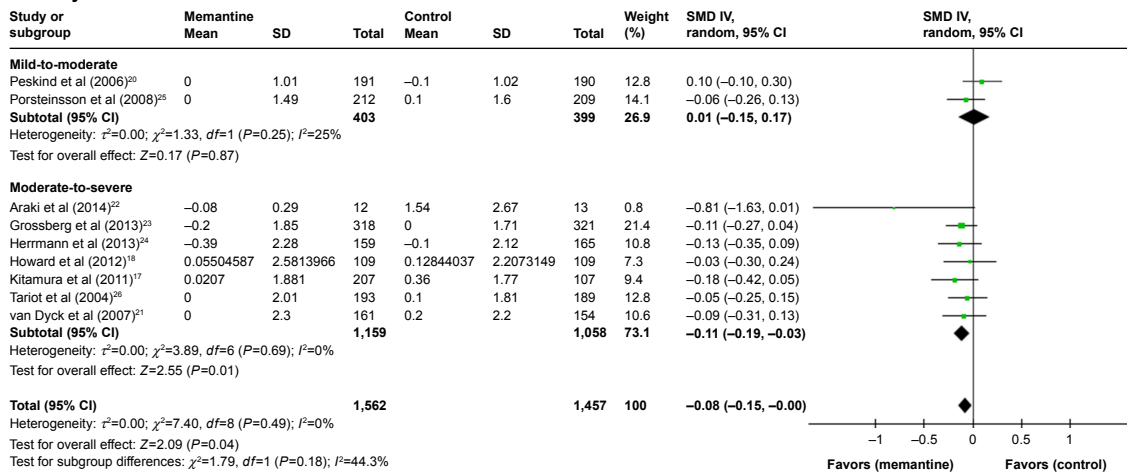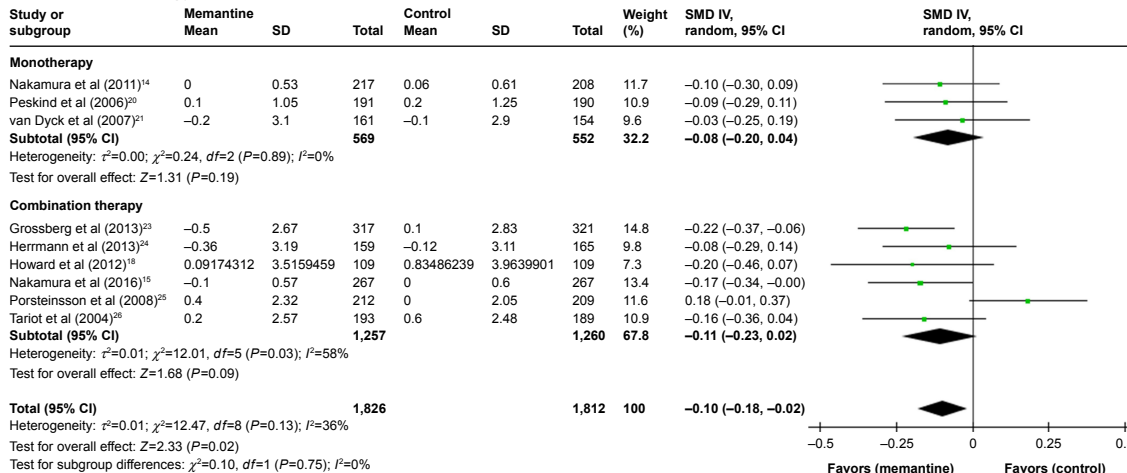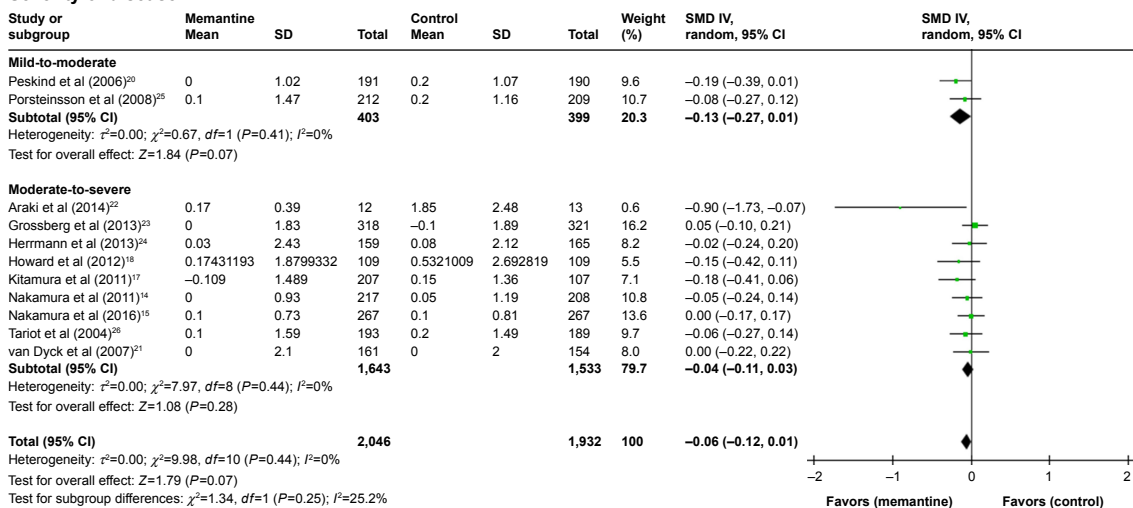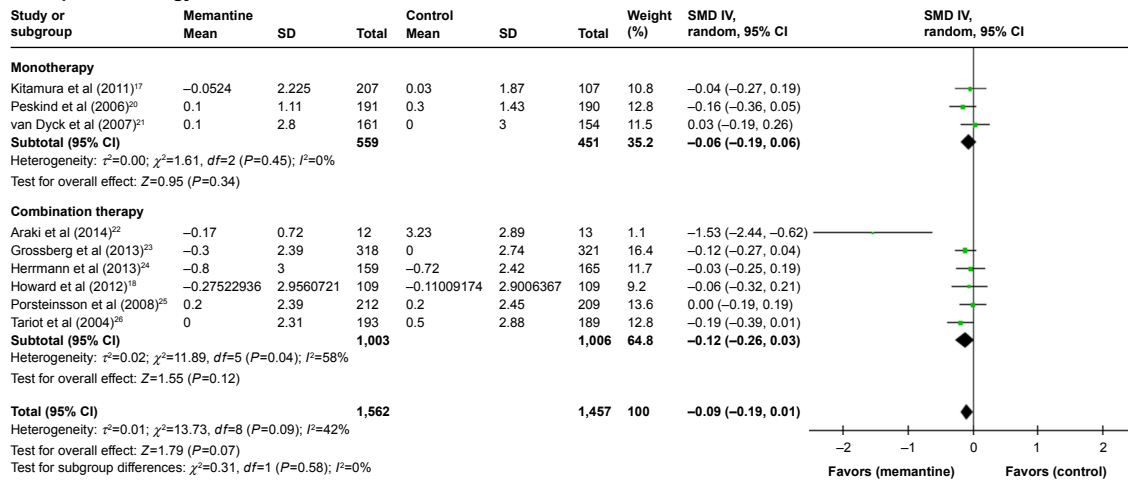
Therapeutic strategy

Study or subgroup	Memantine Mean	SD	Total	Control Mean	SD	Total	Weight (%)	SMD IV, random, 95% CI	SMD IV, random, 95% CI
Monotherapy									
Kitamura et al (2011)[17]	−0.0524	2.225	207	0.03	1.87	107	10.8	−0.04 (−0.27, 0.19)	
Peskind et al (2006)[20]	0.1	1.11	191	0.3	1.43	190	12.8	−0.16 (−0.36, 0.05)	
van Dyck et al (2007)[21]	0.1	2.8	161	0	3	154	11.5	0.03 (−0.19, 0.26)	
Subtotal (95% CI)			559			451	35.2	−0.06 (−0.19, 0.06)	
Heterogeneity: τ^2=0.00; χ^2=1.61, df=2 (P=0.45); I^2=0%									
Test for overall effect: Z=0.95 (P=0.34)									
Combination therapy									
Araki et al (2014)[22]	−0.17	0.72	12	3.23	2.89	13	1.1	−1.53 (−2.44, −0.62)	
Grossberg et al (2013)[23]	−0.3	2.39	318	0	2.74	321	16.4	−0.12 (−0.27, 0.04)	
Herrmann et al (2013)[24]	−0.8	3	159	−0.72	2.42	165	11.7	−0.03 (−0.25, 0.19)	
Howard et al (2012)[18]	−0.27522936	2.9560721	109	−0.11009174	2.9006367	109	9.2	−0.06 (−0.32, 0.21)	
Porsteinsson et al (2008)[25]	0.2	2.39	212	0.2	2.45	209	13.6	0.00 (−0.19, 0.19)	
Tariot et al (2004)[26]	0	2.31	193	0.5	2.88	189	12.8	−0.19 (−0.39, 0.01)	
Subtotal (95% CI)			1,003			1,006	64.8	−0.12 (−0.26, 0.03)	
Heterogeneity: τ^2=0.02; χ^2=11.89, df=5 (P=0.04); I^2=58%									
Test for overall effect: Z=1.55 (P=0.12)									
Total (95% CI)			1,562			1,457	100	−0.09 (−0.19, 0.01)	
Heterogeneity: τ^2=0.01; χ^2=13.73, df=8 (P=0.09); I^2=42%									
Test for overall effect: Z=1.79 (P=0.07)									
Test for subgroup differences: χ^2=0.31, df=1 (P=0.58); I^2=0%									

Favors (memantine)   Favors (control)

Severity of disease

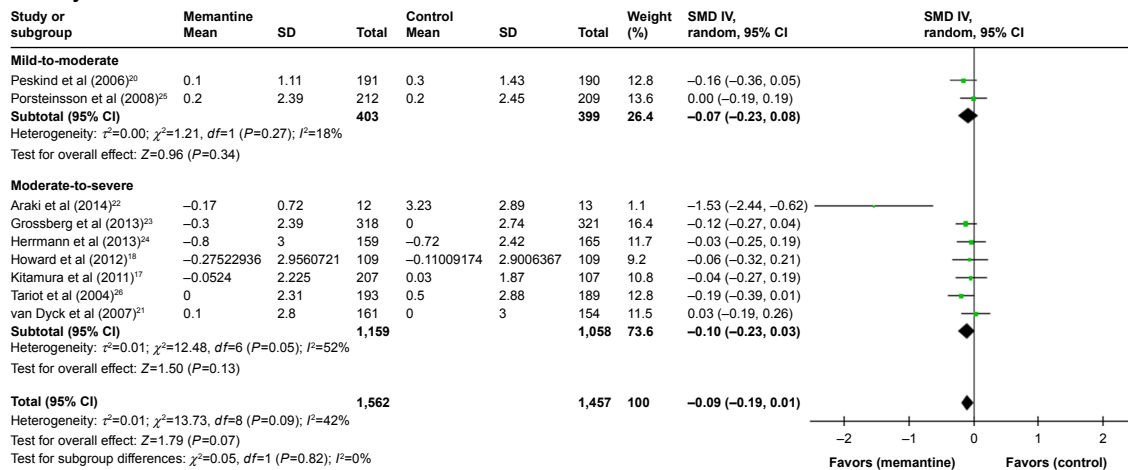| Study or subgroup	Memantine Mean	SD	Total	Control Mean	SD	Total	Weight (%)	SMD IV, random, 95% CI	SMD IV, random, 95% CI
Mild-to-moderate									
Peskind et al (2006)[20]	0.1	1.11	191	0.3	1.43	190	12.8	−0.16 (−0.36, 0.05)	
Porsteinsson et al (2008)[25]	0.2	2.39	212	0.2	2.45	209	13.6	0.00 (−0.19, 0.19)	
Subtotal (95% CI)			403			399	26.4	−0.07 (−0.23, 0.08)	
Heterogeneity: τ^2=0.00; χ^2=1.21, df=1 (P=0.27); I^2=18%									
Test for overall effect: Z=0.96 (P=0.34)									
Moderate-to-severe									
Araki et al (2014)[22]	−0.17	0.72	12	3.23	2.89	13	1.1	−1.53 (−2.44, −0.62)	
Grossberg et al (2013)[23]	−0.3	2.39	318	0	2.74	321	16.4	−0.12 (−0.27, 0.04)	
Herrmann et al (2013)[24]	−0.8	3	159	−0.72	2.42	165	11.7	−0.03 (−0.25, 0.19)	
Howard et al (2012)[18]	−0.27522936	2.9560721	109	−0.11009174	2.9006367	109	9.2	−0.06 (−0.32, 0.21)	
Kitamura et al (2011)[17]	−0.0524	2.225	207	0.03	1.87	107	10.8	−0.04 (−0.27, 0.19)	
Tariot et al (2004)[26]	0	2.31	193	0.5	2.88	189	12.8	−0.19 (−0.39, 0.01)	
van Dyck et al (2007)[21]	0.1	2.8	161	0	3	154	11.5	0.03 (−0.19, 0.26)	
Subtotal (95% CI)			1,159			1,058	73.6	−0.10 (−0.23, 0.03)	
Heterogeneity: τ^2=0.01; χ^2=12.48, df=6 (P=0.05); I^2=52%									
Test for overall effect: Z=1.50 (P=0.13)									
Total (95% CI)			1,562			1,457	100	−0.09 (−0.19, 0.01)	
Heterogeneity: τ^2=0.01; χ^2=13.73, df=8 (P=0.09); I^2=42%									
Test for overall effect: Z=1.79 (P=0.07)									
Test for subgroup differences: χ^2=0.05, df=1 (P=0.82); I^2=0%									

Favors (memantine)   Favors (control)

Figure 6 Forest plot of irritability/lability scores.
Abbreviations: 95% CI, 95% confidence interval; IV, inverse variance; SD, standard deviation; SMD, standardized mean difference.
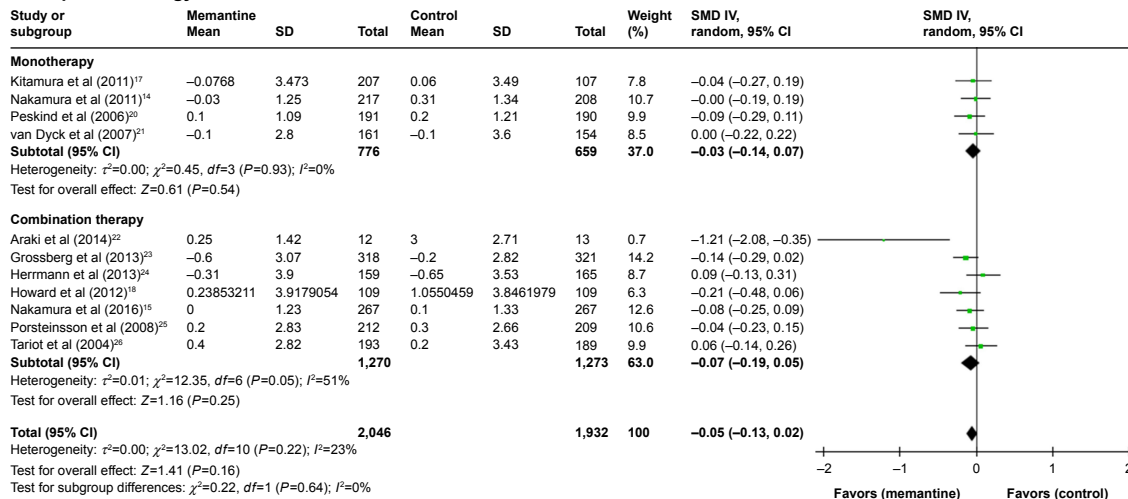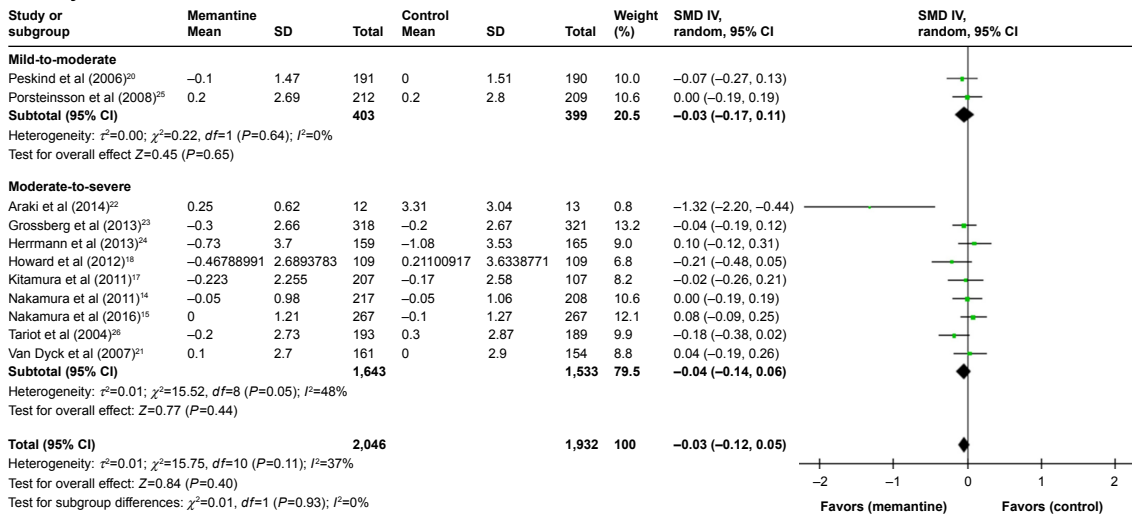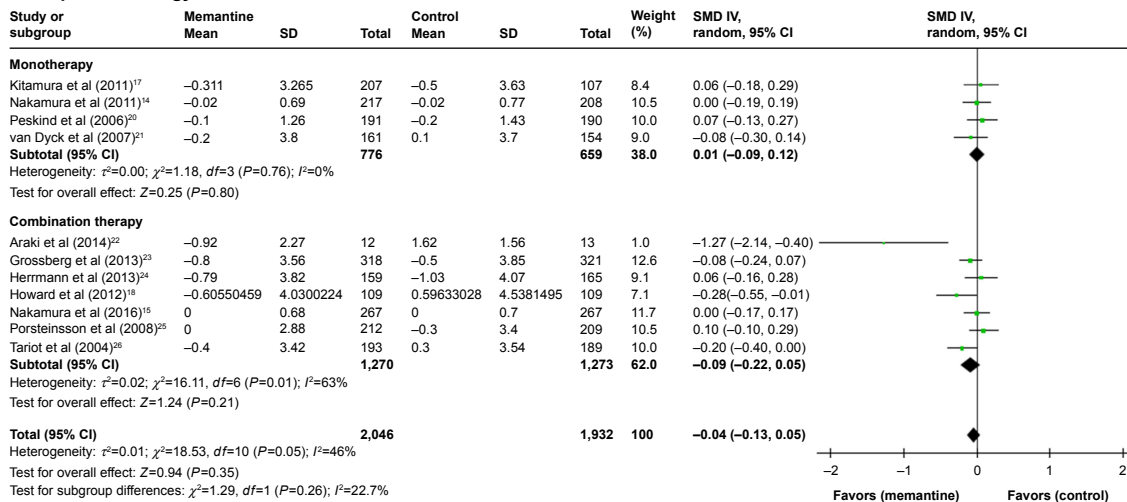
Therapeutic strategy

Study or subgroup	Memantine Mean	SD	Total	Control Mean	SD	Total	Weight (%)	SMD IV, random, 95% CI	SMD IV, random, 95% CI
Monotherapy									
Kitamura et al (2011)[17]	−0.0768	3.473	207	0.06	3.49	107	7.8	−0.04 (−0.27, 0.19)	
Nakamura et al (2011)[14]	−0.03	1.25	217	0.31	1.34	208	10.7	−0.00 (−0.19, 0.19)	
Peskind et al (2006)[20]	0.1	1.09	191	0.2	1.21	190	9.9	−0.09 (−0.29, 0.11)	
van Dyck et al (2007)[21]	−0.1	2.8	161	−0.1	3.6	154	8.5	0.00 (−0.22, 0.22)	
Subtotal (95% CI)			776			659	37.0	−0.03 (−0.14, 0.07)	
Heterogeneity: τ^2=0.00; χ^2=0.45, df=3 (P=0.93); I^2=0%									
Test for overall effect: Z=0.61 (P=0.54)									
Combination therapy									
Araki et al (2014)[22]	0.25	1.42	12	3	2.71	13	0.7	−1.21 (−2.08, −0.35)	
Grossberg et al (2013)[23]	−0.6	3.07	318	−0.2	2.82	321	14.2	−0.14 (−0.29, 0.02)	
Herrmann et al (2013)[24]	−0.31	3.9	159	−0.65	3.53	165	8.7	0.09 (−0.13, 0.31)	
Howard et al (2012)[18]	0.23853211	3.9179054	109	1.0550459	3.8461979	109	6.3	−0.21 (−0.48, 0.06)	
Nakamura et al (2016)[15]	0	1.23	267	0.1	1.33	267	12.6	−0.08 (−0.25, 0.09)	
Porsteinsson et al (2008)[25]	0.2	2.83	212	0.3	2.66	209	10.6	−0.04 (−0.23, 0.15)	
Tariot et al (2004)[26]	0.4	2.82	193	0.2	3.43	189	9.9	0.06 (−0.14, 0.26)	
Subtotal (95% CI)			1,270			1,273	63.0	−0.07 (−0.19, 0.05)	
Heterogeneity: τ^2=0.01; χ^2=12.35, df=6 (P=0.05); I^2=51%									
Test for overall effect: Z=1.16 (P=0.25)									
Total (95% CI)			2,046			1,932	100	−0.05 (−0.13, 0.02)	
Heterogeneity: τ^2=0.00; χ^2=13.02, df=10 (P=0.22); I^2=23%									
Test for overall effect: Z=1.41 (P=0.16)									
Test for subgroup differences: χ^2=0.22, df=1 (P=0.64); I^2=0%									

Favors (memantine)   Favors (control)

Figure 7 (*Continued*)

Severity of disease

Study or subgroup	Memantine Mean	SD	Total	Control Mean	SD	Total	Weight (%)	SMD IV, random, 95% CI
Mild-to-moderate								
Peskind et al (2006)[20]	0.1	1.09	191	0.2	1.21	190	9.9	−0.09 (−0.29, 0.11)
Porsteinsson et al (2008)[25]	0.2	2.83	212	0.3	2.66	209	10.6	−0.04 (−0.23, 0.15)
Subtotal (95% CI)			403			399	20.5	−0.06 (−0.20, 0.08)
Heterogeneity: $\tau^2=0.00$; $\chi^2=0.13$, $df=1$ ($P=0.72$); $I^2=0\%$								
Test for overall effect: $Z=0.85$ ($P=0.39$)								
Moderate-to-severe								
Araki et al (2014)[22]	0.25	1.42	12	3	2.71	13	0.7	−1.21 (−2.08, −0.35)
Grossberg et al (2013)[23]	−0.6	3.07	318	−0.2	2.82	321	14.2	−0.14 (−0.29, 0.02)
Herrmann et al (2013)[24]	−0.31	3.9	159	−0.65	3.53	165	8.7	0.09 (−0.13, 0.31)
Howard et al (2012)[18]	0.23853211	3.9179054	109	1.0550459	3.8461979	109	6.3	−0.21 (−0.48, 0.06)
Kitamura et al (2011)[17]	−0.0768	3.473	207	0.06	3.49	107	7.8	−0.04 (−0.27, 0.19)
Nakamura et al (2011)[14]	−0.03	1.25	217	0.31	134	208	10.7	−0.00 (−0.19, 0.19)
Nakamura et al (2016)[15]	0	1.23	267	0.1	1.33	267	12.6	−0.08 (−0.25, 0.09)
Tariot et al (2004)[26]	0.4	2.82	193	0.2	3.43	189	9.9	0.06 (−0.14, 0.26)
van Dyck et al (2007)[21]	−0.1	2.8	161	−0.1	3.6	154	8.5	0.00 (−0.22, 0.22)
Subtotal (95% CI)			1,643			1,533	79.5	−0.05 (−0.14, 0.04)
Heterogeneity: $\tau^2=0.01$; $\chi^2=12.87$, $df=8$ ($P=0.12$); $I^2=38\%$								
Test for overall effect: $Z=1.10$ ($P=0.27$)								
Total (95% CI)			2,046			1,932	100	−0.05 (−0.13, 0.02)
Heterogeneity: $\tau^2=0.00$; $\chi^2=13.02$, $df=10$ ($P=0.22$); $I^2=23\%$								
Test for overall effect: $Z=1.41$ ($P=0.16$)								
Test for subgroup differences: $\chi^2=0.01$, $df=1$ ($P=0.92$); $I^2=0\%$								

Favors (memantine)　Favors (control)

Figure 7 Forest plot of aberrant motor activity/activity disturbance scores.
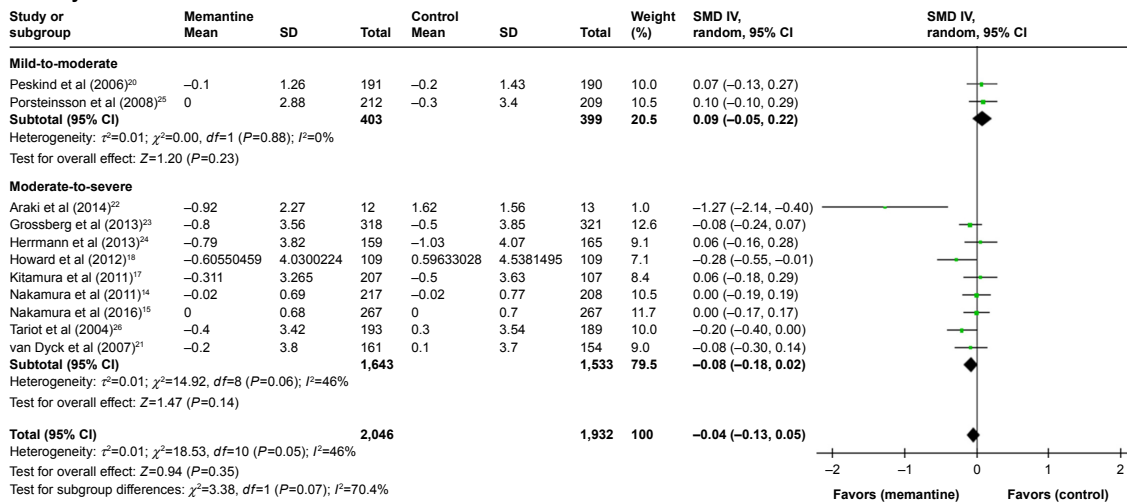**Abbreviations:** 95% CI, 95% confidence interval; IV, inverse variance; SD, standard deviation; SMD, standardized mean difference.
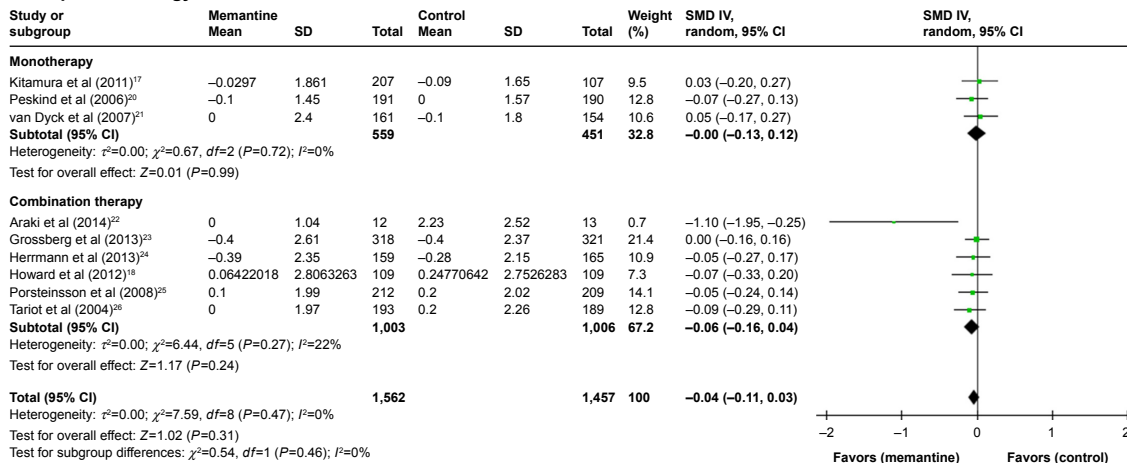
Therapeutic strategy

Study or subgroup	Memantine Mean	SD	Total	Control Mean	SD	Total	Weight (%)	SMD IV, random, 95% CI
Monotherapy								
Kitamura et al (2011)[17]	−0.223	2.255	207	−0.17	2.58	107	8.2	−0.02 (−0.26, 0.21)
Nakamura et al (2011)[14]	−0.05	0.98	217	−0.05	1.06	208	10.6	0.00 (−0.19, 0.19)
Peskind et al (2006)[20]	−0.1	1.47	191	0	1.51	190	10.0	−0.07 (−0.27, 0.13)
van Dyck et al (2007)[21]	0.1	2.7	161	0	2.9	154	8.8	0.04 (−0.19, 0.26)
Subtotal (95% CI)			776			659	37.6	−0.01 (−0.12, 0.09)
Heterogeneity: $\tau^2=0.00$; $\chi^2=0.49$, $df=3$ ($P=0.92$); $I^2=0\%$								
Test for overall effect: $Z=0.27$ ($P=0.78$)								
Combination therapy								
Araki et al (2014)[22]	0.25	0.62	12	3.31	3.04	13	0.8	−1.32 (−2.20, −0.44)
Grossberg et al (2013)[23]	−0.3	2.66	318	−0.2	2.67	321	13.2	−0.04 (−0.19, 0.12)
Herrmann et al (2013)[24]	−0.73	3.7	159	−1.08	3.53	165	9.0	0.10 (−0.12, 0.31)
Howard et al (2012)[18]	−0.46788991	2.6893783	109	0.21100917	3.6338771	109	6.8	−0.21 (−0.48, 0.05)
Nakamura et al (2016)[15]	0	1.21	267	−0.1	1.27	267	12.1	0.08 (−0.09, 0.25)
Porsteinsson et al (2008)[25]	0.2	2.69	212	0.2	2.8	209	10.6	0.00 (−0.19, 0.19)
Tariot et al (2004)[26]	−0.2	2.73	193	0.3	2.87	189	9.9	−0.18 (−0.38, 0.02)
Subtotal (95% CI)			1,270			1,273	62.4	−0.06 (−0.19, 0.07)
Heterogeneity: $\tau^2=0.02$; $\chi^2=15.17$, $df=6$ ($P=0.02$); $I^2=60\%$								
Test for overall effect: $Z=0.90$ ($P=0.37$)								
Total (95% CI)			2,046			1,932	100	−0.03 (−0.12, 0.05)
Heterogeneity: $\tau^2=0.01$; $\chi^2=15.75$, $df=10$ ($P=0.11$); $I^2=37\%$								
Test for overall effect: $Z=0.84$ ($P=0.40$)								
Test for subgroup differences: $\chi^2=0.29$, $df=1$ ($P=0.59$); $I^2=0\%$								

Favors (memantine)　Favors (control)

Severity of disease

Study or subgroup	Memantine Mean	SD	Total	Control Mean	SD	Total	Weight (%)	SMD IV, random, 95% CI
Mild-to-moderate								
Peskind et al (2006)[20]	−0.1	1.47	191	0	1.51	190	10.0	−0.07 (−0.27, 0.13)
Porsteinsson et al (2008)[25]	0.2	2.69	212	0.2	2.8	209	10.6	0.00 (−0.19, 0.19)
Subtotal (95% CI)			403			399	20.5	−0.03 (−0.17, 0.11)
Heterogeneity: $\tau^2=0.00$; $\chi^2=0.22$, $df=1$ ($P=0.64$); $I^2=0\%$								
Test for overall effect $Z=0.45$ ($P=0.65$)								
Moderate-to-severe								
Araki et al (2014)[22]	0.25	0.62	12	3.31	3.04	13	0.8	−1.32 (−2.20, −0.44)
Grossberg et al (2013)[23]	−0.3	2.66	318	−0.2	2.67	321	13.2	−0.04 (−0.19, 0.12)
Herrmann et al (2013)[24]	−0.73	3.7	159	−1.08	3.53	165	9.0	0.10 (−0.12, 0.31)
Howard et al (2012)[18]	−0.46788991	2.6893783	109	0.21100917	3.6338771	109	6.8	−0.21 (−0.48, 0.05)
Kitamura et al (2011)[17]	−0.223	2.255	207	−0.17	2.58	107	8.2	−0.02 (−0.26, 0.21)
Nakamura et al (2011)[14]	−0.05	0.98	217	−0.05	1.06	208	10.6	0.00 (−0.19, 0.19)
Nakamura et al (2016)[15]	0	1.21	267	−0.1	1.27	267	12.1	0.08 (−0.09, 0.25)
Tariot et al (2004)[26]	−0.2	2.73	193	0.3	2.87	189	9.9	−0.18 (−0.38, 0.02)
Van Dyck et al (2007)[21]	0.1	2.7	161	0	2.9	154	8.8	0.04 (−0.19, 0.26)
Subtotal (95% CI)			1,643			1,533	79.5	−0.04 (−0.14, 0.06)
Heterogeneity: $\tau^2=0.01$; $\chi^2=15.52$, $df=8$ ($P=0.05$); $I^2=48\%$								
Test for overall effect: $Z=0.77$ ($P=0.44$)								
Total (95% CI)			2,046			1,932	100	−0.03 (−0.12, 0.05)
Heterogeneity: $\tau^2=0.01$; $\chi^2=15.75$, $df=10$ ($P=0.11$); $I^2=37\%$								
Test for overall effect: $Z=0.84$ ($P=0.40$)								
Test for subgroup differences: $\chi^2=0.01$, $df=1$ ($P=0.93$); $I^2=0\%$								

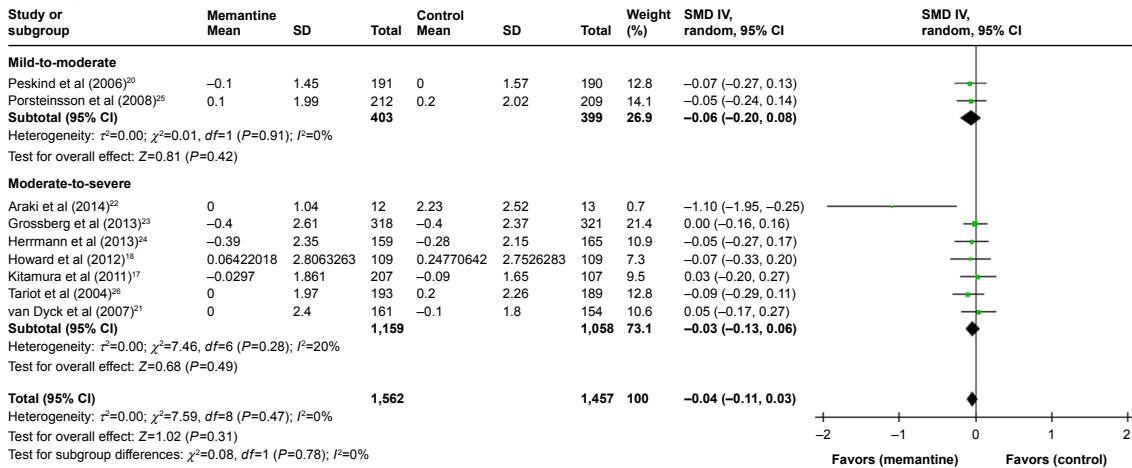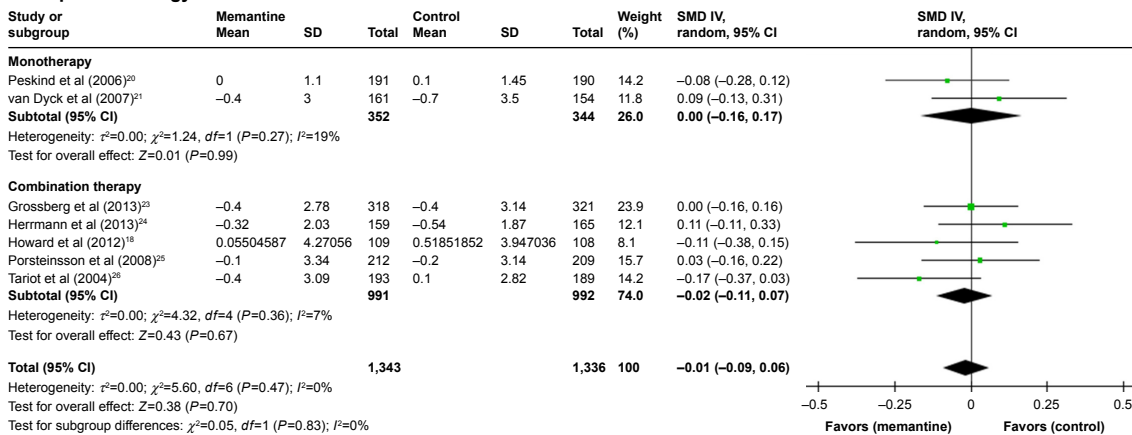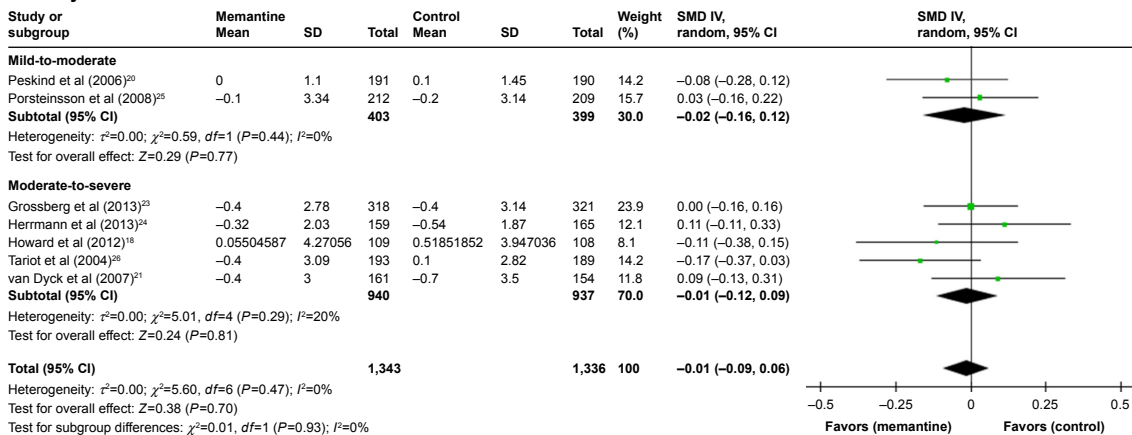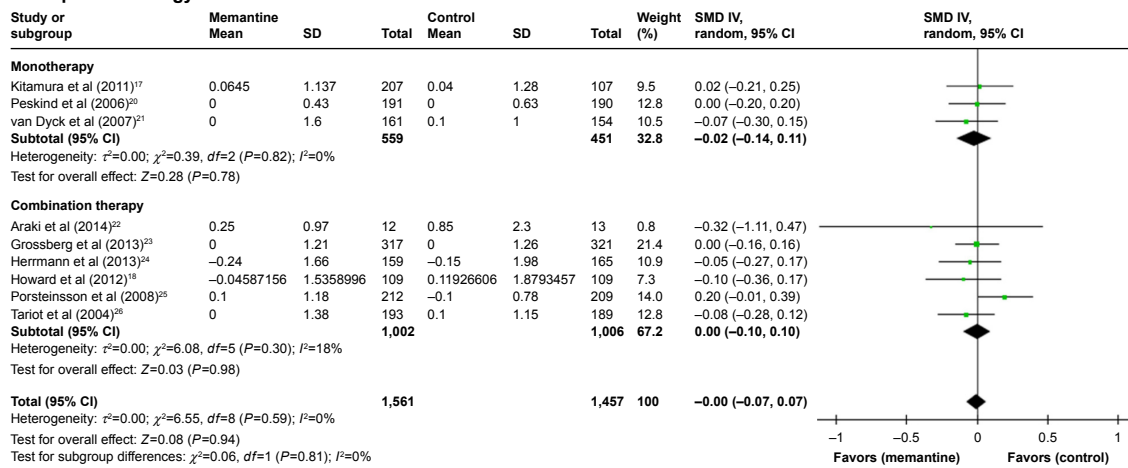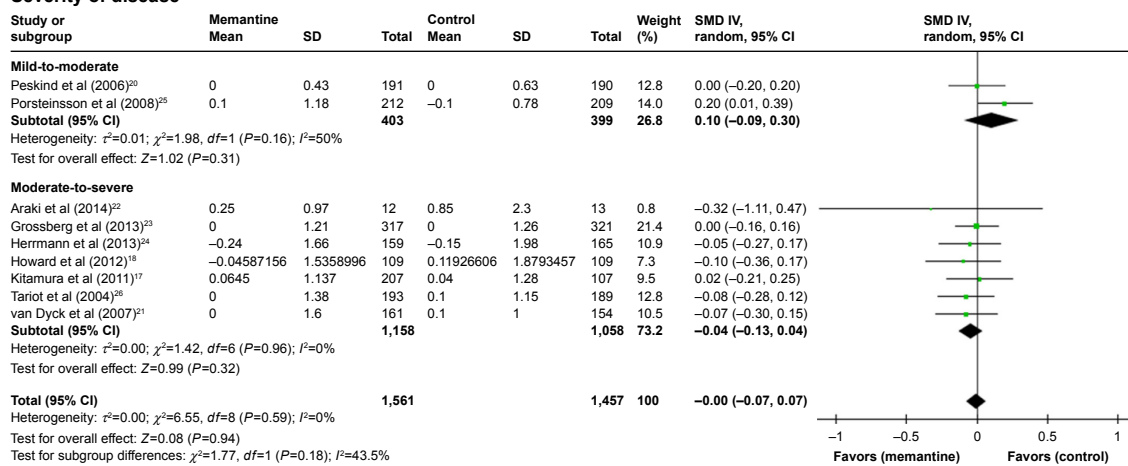Favors (memantine)　Favors (control)

Figure 8 Forest plot of anxiety/phobia scores.
Abbreviations: 95% CI, 95% confidence interval; IV, inverse variance; SD, standard deviation; SMD, standardized mean difference.

Therapeutic strategy

Study or subgroup	Memantine Mean	SD	Total	Control Mean	SD	Total	Weight (%)	SMD IV, random, 95% CI
Monotherapy								
Kitamura et al (2011)[17]	−0.311	3.265	207	−0.5	3.63	107	8.4	0.06 (−0.18, 0.29)
Nakamura et al (2011)[14]	−0.02	0.69	217	−0.02	0.77	208	10.5	0.00 (−0.19, 0.19)
Peskind et al (2006)[20]	−0.1	1.26	191	−0.2	1.43	190	10.0	0.07 (−0.13, 0.27)
van Dyck et al (2007)[21]	−0.2	3.8	161	0.1	3.7	154	9.0	−0.08 (−0.30, 0.14)
Subtotal (95% CI)			776			659	38.0	0.01 (−0.09, 0.12)
Heterogeneity: τ^2=0.00; χ^2=1.18, df=3 (P=0.76); I^2=0%								
Test for overall effect: Z=0.25 (P=0.80)								
Combination therapy								
Araki et al (2014)[22]	−0.92	2.27	12	1.62	1.56	13	1.0	−1.27 (−2.14, −0.40)
Grossberg et al (2013)[23]	−0.8	3.56	318	−0.5	3.85	321	12.6	−0.08 (−0.24, 0.07)
Herrmann et al (2013)[24]	−0.79	3.82	159	−1.03	4.07	165	9.1	0.06 (−0.16, 0.28)
Howard et al (2012)[18]	−0.60550459	4.0300224	109	0.59633028	4.5381495	109	7.1	−0.28 (−0.55, −0.01)
Nakamura et al (2016)[15]	0	0.68	267	0	0.7	267	11.7	0.00 (−0.17, 0.17)
Porsteinsson et al (2008)[25]	0	2.88	212	−0.3	3.4	209	10.5	0.10 (−0.10, 0.29)
Tariot et al (2004)[26]	−0.4	3.42	193	0.3	3.54	189	10.0	−0.20 (−0.40, 0.00)
Subtotal (95% CI)			1,270			1,273	62.0	−0.09 (−0.22, 0.05)
Heterogeneity: τ^2=0.02; χ^2=16.11, df=6 (P=0.01); I^2=63%								
Test for overall effect: Z=1.24 (P=0.21)								
Total (95% CI)			2,046			1,932	100	−0.04 (−0.13, 0.05)
Heterogeneity: τ^2=0.01; χ^2=18.53, df=10 (P=0.05); I^2=46%								
Test for overall effect: Z=0.94 (P=0.35)								
Test for subgroup differences: χ^2=1.29, df=1 (P=0.26); I^2=22.7%								

SMD IV, random, 95% CI — scale: −2 −1 0 1 2 — Favors (memantine) / Favors (control)

Severity of disease

Study or subgroup	Memantine Mean	SD	Total	Control Mean	SD	Total	Weight (%)	SMD IV, random, 95% CI
Mild-to-moderate								
Peskind et al (2006)[20]	−0.1	1.26	191	−0.2	1.43	190	10.0	0.07 (−0.13, 0.27)
Porsteinsson et al (2008)[25]	0	2.88	212	−0.3	3.4	209	10.5	0.10 (−0.10, 0.29)
Subtotal (95% CI)			403			399	20.5	0.09 (−0.05, 0.22)
Heterogeneity: τ^2=0.01; χ^2=0.00, df=1 (P=0.88); I^2=0%								
Test for overall effect: Z=1.20 (P=0.23)								
Moderate-to-severe								
Araki et al (2014)[22]	−0.92	2.27	12	1.62	1.56	13	1.0	−1.27 (−2.14, −0.40)
Grossberg et al (2013)[23]	−0.8	3.56	318	−0.5	3.85	321	12.6	−0.08 (−0.24, 0.07)
Herrmann et al (2013)[24]	−0.79	3.82	159	−1.03	4.07	165	9.1	0.06 (−0.16, 0.28)
Howard et al (2012)[18]	−0.60550459	4.0300224	109	0.59633028	4.5381495	109	7.1	−0.28 (−0.55, −0.01)
Kitamura et al (2011)[17]	−0.311	3.265	207	−0.5	3.63	107	8.4	0.06 (−0.18, 0.29)
Nakamura et al (2011)[14]	−0.02	0.69	217	−0.02	0.77	208	10.5	0.00 (−0.19, 0.19)
Nakamura et al (2016)[15]	0	0.68	267	0	0.7	267	11.7	0.00 (−0.17, 0.17)
Tariot et al (2004)[26]	−0.4	3.42	193	0.3	3.54	189	10.0	−0.20 (−0.40, 0.00)
van Dyck et al (2007)[21]	−0.2	3.8	161	0.1	3.7	154	9.0	−0.08 (−0.30, 0.14)
Subtotal (95% CI)			1,643			1,533	79.5	−0.08 (−0.18, 0.02)
Heterogeneity: τ^2=0.01; χ^2=14.92, df=8 (P=0.06); I^2=46%								
Test for overall effect: Z=1.47 (P=0.14)								
Total (95% CI)			2,046			1,932	100	−0.04 (−0.13, 0.05)
Heterogeneity: τ^2=0.01; χ^2=18.53, df=10 (P=0.05); I^2=46%								
Test for overall effect: Z=0.94 (P=0.35)								
Test for subgroup differences: χ^2=3.38, df=1 (P=0.07); I^2=70.4%								

SMD IV, random, 95% CI — scale: −2 −1 0 1 2 — Favors (memantine) / Favors (control)

Figure 9 Forest plot of apathy scores.
Abbreviations: 95% CI, 95% confidence interval; IV, inverse variance; SD, standard deviation; SMD, standardized mean difference.

Therapeutic strategy

Study or subgroup	Memantine Mean	SD	Total	Control Mean	SD	Total	Weight (%)	SMD IV, random, 95% CI
Monotherapy								
Kitamura et al (2011)[17]	−0.0297	1.861	207	−0.09	1.65	107	9.5	0.03 (−0.20, 0.27)
Peskind et al (2006)[20]	−0.1	1.45	191	0	1.57	190	12.8	−0.07 (−0.27, 0.13)
van Dyck et al (2007)[21]	0	2.4	161	−0.1	1.8	154	10.6	0.05 (−0.17, 0.27)
Subtotal (95% CI)			559			451	32.8	−0.00 (−0.13, 0.12)
Heterogeneity: τ^2=0.00; χ^2=0.67, df=2 (P=0.72); I^2=0%								
Test for overall effect: Z=0.01 (P=0.99)								
Combination therapy								
Araki et al (2014)[22]	0	1.04	12	2.23	2.52	13	0.7	−1.10 (−1.95, −0.25)
Grossberg et al (2013)[23]	−0.4	2.61	318	−0.4	2.37	321	21.4	0.00 (−0.16, 0.16)
Herrmann et al (2013)[24]	−0.39	2.35	159	−0.28	2.15	165	10.9	−0.05 (−0.27, 0.17)
Howard et al (2012)[18]	0.06422018	2.8063263	109	0.24770642	2.7526283	109	7.3	−0.07 (−0.33, 0.20)
Porsteinsson et al (2008)[25]	0.1	1.99	212	0.2	2.02	209	14.1	−0.05 (−0.24, 0.14)
Tariot et al (2004)[26]	0	1.97	193	0.2	2.26	189	12.8	−0.09 (−0.29, 0.11)
Subtotal (95% CI)			1,003			1,006	67.2	−0.06 (−0.16, 0.04)
Heterogeneity: τ^2=0.00; χ^2=6.44, df=5 (P=0.27); I^2=22%								
Test for overall effect: Z=1.17 (P=0.24)								
Total (95% CI)			1,562			1,457	100	−0.04 (−0.11, 0.03)
Heterogeneity: τ^2=0.00; χ^2=7.59, df=8 (P=0.47); I^2=0%								
Test for overall effect: Z=1.02 (P=0.31)								
Test for subgroup differences: χ^2=0.54, df=1 (P=0.46); I^2=0%								

SMD IV, random, 95% CI — scale: −2 −1 0 1 2 — Favors (memantine) / Favors (control)

Figure 10 (*Continued*)

Severity of disease

Study or subgroup	Memantine Mean	SD	Total	Control Mean	SD	Total	Weight (%)	SMD IV, random, 95% CI	SMD IV, random, 95% CI
Mild-to-moderate									
Peskind et al (2006)[20]	−0.1	1.45	191	0	1.57	190	12.8	−0.07 (−0.27, 0.13)	
Porsteinsson et al (2008)[25]	0.1	1.99	212	0.2	2.02	209	14.1	−0.05 (−0.24, 0.14)	
Subtotal (95% CI)			403			399	26.9	−0.06 (−0.20, 0.08)	
Heterogeneity: τ^2=0.00; χ^2=0.01, df=1 (P=0.91); I^2=0%									
Test for overall effect: Z=0.81 (P=0.42)									
Moderate-to-severe									
Araki et al (2014)[22]	0	1.04	12	2.23	2.52	13	0.7	−1.10 (−1.95, −0.25)	
Grossberg et al (2013)[23]	−0.4	2.61	318	−0.4	2.37	321	21.4	0.00 (−0.16, 0.16)	
Herrmann et al (2013)[24]	−0.39	2.35	159	−0.28	2.15	165	10.9	−0.05 (−0.27, 0.17)	
Howard et al (2012)[18]	0.06422018	2.8063263	109	0.24770642	2.7526283	109	7.3	−0.07 (−0.33, 0.20)	
Kitamura et al (2011)[17]	−0.0297	1.861	207	−0.09	1.65	107	9.5	0.03 (−0.20, 0.27)	
Tariot et al (2004)[26]	0	1.97	193	0.2	2.26	189	12.8	−0.09 (−0.29, 0.11)	
van Dyck et al (2007)[21]	0	2.4	161	−0.1	1.8	154	10.6	0.05 (−0.17, 0.27)	
Subtotal (95% CI)			1,159			1,058	73.1	−0.03 (−0.13, 0.06)	
Heterogeneity: τ^2=0.00; χ^2=7.46, df=6 (P=0.28); I^2=20%									
Test for overall effect: Z=0.68 (P=0.49)									
Total (95% CI)			1,562			1,457	100	−0.04 (−0.11, 0.03)	
Heterogeneity: τ^2=0.00; χ^2=7.59, df=8 (P=0.47); I^2=0%									
Test for overall effect: Z=1.02 (P=0.31)									
Test for subgroup differences: χ^2=0.08, df=1 (P=0.78); I^2=0%									

Favors (memantine)   Favors (control)

Figure 10 Forest plot of dysphoria scores.
Abbreviations: 95% CI, 95% confidence interval; IV, inverse variance; SD, standard deviation; SMD, standardized mean difference.

Therapeutic strategy

Study or subgroup	Memantine Mean	SD	Total	Control Mean	SD	Total	Weight (%)	SMD IV, random, 95% CI	SMD IV, random, 95% CI
Monotherapy									
Peskind et al (2006)[20]	0	1.1	191	0.1	1.45	190	14.2	−0.08 (−0.28, 0.12)	
van Dyck et al (2007)[21]	−0.4	3	161	−0.7	3.5	154	11.8	0.09 (−0.13, 0.31)	
Subtotal (95% CI)			352			344	26.0	0.00 (−0.16, 0.17)	
Heterogeneity: τ^2=0.00; χ^2=1.24, df=1 (P=0.27); I^2=19%									
Test for overall effect: Z=0.01 (P=0.99)									
Combination therapy									
Grossberg et al (2013)[23]	−0.4	2.78	318	−0.4	3.14	321	23.9	0.00 (−0.16, 0.16)	
Herrmann et al (2013)[24]	−0.32	2.03	159	−0.54	1.87	165	12.1	0.11 (−0.11, 0.33)	
Howard et al (2012)[18]	0.05504587	4.27056	109	0.51851852	3.947036	108	8.1	−0.11 (−0.38, 0.15)	
Porsteinsson et al (2008)[25]	−0.1	3.34	212	−0.2	3.14	209	15.7	0.03 (−0.16, 0.22)	
Tariot et al (2004)[26]	−0.4	3.09	193	0.1	2.82	189	14.2	−0.17 (−0.37, 0.03)	
Subtotal (95% CI)			991			992	74.0	−0.02 (−0.11, 0.07)	
Heterogeneity: τ^2=0.00; χ^2=4.32, df=4 (P=0.36); I^2=7%									
Test for overall effect: Z=0.43 (P=0.67)									
Total (95% CI)			1,343			1,336	100	−0.01 (−0.09, 0.06)	
Heterogeneity: τ^2=0.00; χ^2=5.60, df=6 (P=0.47); I^2=0%									
Test for overall effect: Z=0.38 (P=0.70)									
Test for subgroup differences: χ^2=0.05, df=1 (P=0.83); I^2=0%									

Favors (memantine)   Favors (control)

Severity of disease

Study or subgroup	Memantine Mean	SD	Total	Control Mean	SD	Total	Weight (%)	SMD IV, random, 95% CI	SMD IV, random, 95% CI
Mild-to-moderate									
Peskind et al (2006)[20]	0	1.1	191	0.1	1.45	190	14.2	−0.08 (−0.28, 0.12)	
Porsteinsson et al (2008)[25]	−0.1	3.34	212	−0.2	3.14	209	15.7	0.03 (−0.16, 0.22)	
Subtotal (95% CI)			403			399	30.0	−0.02 (−0.16, 0.12)	
Heterogeneity: τ^2=0.00; χ^2=0.59, df=1 (P=0.44); I^2=0%									
Test for overall effect: Z=0.29 (P=0.77)									
Moderate-to-severe									
Grossberg et al (2013)[23]	−0.4	2.78	318	−0.4	3.14	321	23.9	0.00 (−0.16, 0.16)	
Herrmann et al (2013)[24]	−0.32	2.03	159	−0.54	1.87	165	12.1	0.11 (−0.11, 0.33)	
Howard et al (2012)[18]	0.05504587	4.27056	109	0.51851852	3.947036	108	8.1	−0.11 (−0.38, 0.15)	
Tariot et al (2004)[26]	−0.4	3.09	193	0.1	2.82	189	14.2	−0.17 (−0.37, 0.03)	
van Dyck et al (2007)[21]	−0.4	3	161	−0.7	3.5	154	11.8	0.09 (−0.13, 0.31)	
Subtotal (95% CI)			940			937	70.0	−0.01 (−0.12, 0.09)	
Heterogeneity: τ^2=0.00; χ^2=5.01, df=4 (P=0.29); I^2=20%									
Test for overall effect: Z=0.24 (P=0.81)									
Total (95% CI)			1,343			1,336	100	−0.01 (−0.09, 0.06)	
Heterogeneity: τ^2=0.00; χ^2=5.60, df=6 (P=0.47); I^2=0%									
Test for overall effect: Z=0.38 (P=0.70)									
Test for subgroup differences: χ^2=0.01, df=1 (P=0.93); I^2=0%									

Favors (memantine)   Favors (control)

Figure 11 Forest plot of eating disturbance scores.
Abbreviations: 95% CI, 95% confidence interval; IV, inverse variance; SD, standard deviation; SMD, standardized mean difference.

Therapeutic strategy

Study or subgroup	Memantine Mean	SD	Total	Control Mean	SD	Total	Weight (%)	SMD IV, random, 95% CI
Monotherapy								
Kitamura et al (2011)[17]	0.0645	1.137	207	0.04	1.28	107	9.5	0.02 (−0.21, 0.25)
Peskind et al (2006)[20]	0	0.43	191	0	0.63	190	12.8	0.00 (−0.20, 0.20)
van Dyck et al (2007)[21]	0	1.6	161	0.1	1	154	10.5	−0.07 (−0.30, 0.15)
Subtotal (95% CI)			**559**			**451**	**32.8**	**−0.02 (−0.14, 0.11)**
Heterogeneity: τ²=0.00; χ²=0.39, df=2 (P=0.82); I²=0%								
Test for overall effect: Z=0.28 (P=0.78)								
Combination therapy								
Araki et al (2014)[22]	0.25	0.97	12	0.85	2.3	13	0.8	−0.32 (−1.11, 0.47)
Grossberg et al (2013)[23]	0	1.21	317	0	1.26	321	21.4	0.00 (−0.16, 0.16)
Herrmann et al (2013)[24]	−0.24	1.66	159	−0.15	1.98	165	10.9	−0.05 (−0.27, 0.17)
Howard et al (2012)[18]	−0.04587156	1.5358996	109	0.11926606	1.8793457	109	7.3	−0.10 (−0.36, 0.17)
Porsteinsson et al (2008)[25]	0.1	1.18	212	−0.1	0.78	209	14.0	0.20 (−0.01, 0.39)
Tariot et al (2004)[26]	0	1.38	193	0.1	1.15	189	12.8	−0.08 (−0.28, 0.12)
Subtotal (95% CI)			**1,002**			**1,006**	**67.2**	**0.00 (−0.10, 0.10)**
Heterogeneity: τ²=0.00; χ²=6.08, df=5 (P=0.30); I²=18%								
Test for overall effect: Z=0.03 (P=0.98)								
Total (95% CI)			**1,561**			**1,457**	**100**	**−0.00 (−0.07, 0.07)**
Heterogeneity: τ²=0.00; χ²=6.55, df=8 (P=0.59); I²=0%								
Test for overall effect: Z=0.08 (P=0.94)								
Test for subgroup differences: χ²=0.06, df=1 (P=0.81); I²=0%								

SMD IV, random, 95% CI
−1   −0.5   0   0.5   1
Favors (memantine)   Favors (control)

Severity of disease

Study or subgroup	Memantine Mean	SD	Total	Control Mean	SD	Total	Weight (%)	SMD IV, random, 95% CI
Mild-to-moderate								
Peskind et al (2006)[20]	0	0.43	191	0	0.63	190	12.8	0.00 (−0.20, 0.20)
Porsteinsson et al (2008)[25]	0.1	1.18	212	−0.1	0.78	209	14.0	0.20 (0.01, 0.39)
Subtotal (95% CI)			**403**			**399**	**26.8**	**0.10 (−0.09, 0.30)**
Heterogeneity: τ²=0.01; χ²=1.98, df=1 (P=0.16); I²=50%								
Test for overall effect: Z=1.02 (P=0.31)								
Moderate-to-severe								
Araki et al (2014)[22]	0.25	0.97	12	0.85	2.3	13	0.8	−0.32 (−1.11, 0.47)
Grossberg et al (2013)[23]	0	1.21	317	0	1.26	321	21.4	0.00 (−0.16, 0.16)
Herrmann et al (2013)[24]	−0.24	1.66	159	−0.15	1.98	165	10.9	−0.05 (−0.27, 0.17)
Howard et al (2012)[18]	−0.04587156	1.5358996	109	0.11926606	1.8793457	109	7.3	−0.10 (−0.36, 0.17)
Kitamura et al (2011)[17]	0.0645	1.137	207	0.04	1.28	107	9.5	0.02 (−0.21, 0.25)
Tariot et al (2004)[26]	0	1.38	193	0.1	1.15	189	12.8	−0.08 (−0.28, 0.12)
van Dyck et al (2007)[21]	0	1.6	161	0.1	1	154	10.5	−0.07 (−0.30, 0.15)
Subtotal (95% CI)			**1,158**			**1,058**	**73.2**	**−0.04 (−0.13, 0.04)**
Heterogeneity: τ²=0.00; χ²=1.42, df=6 (P=0.96); I²=0%								
Test for overall effect: Z=0.99 (P=0.32)								
Total (95% CI)			**1,561**			**1,457**	**100**	**−0.00 (−0.07, 0.07)**
Heterogeneity: τ²=0.00; χ²=6.55, df=8 (P=0.59); I²=0%								
Test for overall effect: Z=0.08 (P=0.94)								
Test for subgroup differences: χ²=1.77, df=1 (P=0.18); I²=43.5%								

SMD IV, random, 95% CI
−1   −0.5   0   0.5   1
Favors (memantine)   Favors (control)

Figure 12 Forest plot of euphoria scores.
Abbreviations: 95% CI, 95% confidence interval; IV, inverse variance; SD, standard deviation; SMD, standardized mean difference.

respect to apathy (P=0.07), this subgroup analysis showed that memantine was similar to control in moderate-to-severe Alzheimer's disease patients, as well as mild-to-moderate Alzheimer's disease patients (Figure 9).

Agitation/aggression, delusion, disinhibition, and nighttime disturbance/diurnal rhythm disturbances were outcomes, where memantine was superior to control in the moderate-to-severe Alzheimer's disease patients' subgroup, but not in the mild-to-moderate Alzheimer's disease patients' subgroup (Figures 1–4).

Discussion

This meta-analysis showed that memantine showed significant efficacy compared to controls in improving delusion, agitation/aggression, disinhibition, and nighttime disturbance/ diurnal rhythm disturbances in patients with Alzheimer's disease. Moreover, memantine seems to benefit the treatment of hallucination and irritability/lability. These symptoms are classified as positive symptoms.[10] Memantine was similar to controls for negative symptoms, such as dysphoria, anxiety/ phobia, euphoria, apathy, aberrant motor activity/activity disturbances, and eating disturbances. Memantine improves cognitive functions,[8,9] and anti-dementia drugs may prevent brain atrophy in patients with Alzheimer's disease.[27] Therefore, we considered that the evidence that memantine did not deteriorate negative symptoms, such as behavioral disturbances in patients with Alzheimer's disease, was very important for the clinicians and the patients. If the patients receiving memantine have negative symptoms, the evidence suggests that the patients do not need to stop taking memantine.

Although we did not detect any considerable heterogeneity in all of the meta-analysis, we performed two subgroup analysis (severity of disease and therapeutic strategy) to detect confounding factors. We did not find significant subgroup differences. Subgroup analysis could provide the following evidence, although we did not address multiple comparisons: 1) memantine has benefits for the treatment of delusion in patients with not only combination therapy but also memantine monotherapy; 2) patients with combination therapy may have more benefits for the treatment of agitation/aggression, and disinhibition than patients with memantine monotherapy; and 3) patients with moderate-severe Alzheimer's disease may have more benefit for the treatment of agitation/aggression, delusion, disinhibition and nighttime disturbance/diurnal rhythm disturbances than patients with mild-moderate Alzheimer's disease.

There were several limitations in this study which need to be addressed. First, patient characteristics differed between the studies examined including: symptom severity, inclusion criteria, race, ethnicity, and study duration. These differences could generate heterogeneity, when combining data for systematic review and meta-analysis. Second, most studies included in this study were industry-sponsored studies. Therefore, there remains a possibility for sponsorship bias in our results. Third, most of all studies included in the study did not report sufficient information about concomitant drugs such as psychotropic drugs (Table 1). Therefore, we did not examine whether concomitant drugs influence on the results of the meta-analysis. Fourth, because mean patients' age among the studies included in the meta-analysis were very similar (Table 1), we did not perform the meta-regression analysis to examine whether the effect size of memantine was associated with patient age. Fifth, our study focused on memantine treatment for Alzheimer's disease. We considered that it needed to conduct a network meta-analysis of anti-dementia drugs for Alzheimer's disease on efficacy and safety because network meta-analysis can combine direct and indirect evidence to address the frequent absence of randomized trials that directly compare all the interventions of interest. This should offer suggestion on which pharmacological interventions for the Alzheimer's disease is best.

Conclusion

The meta-analysis suggest that memantine has benefits for the treatment of most of the behavioral disturbances in patients with Alzheimer's disease. Memantine does not deteriorate negative symptoms as behavioral disturbances in patients with Alzheimer's disease.

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Supplementary materials

Potentially relevant studies (N=2,239)

Duplicate studies (N=1,498)

Unique studies identified and screened (N=741)

Studies excluded at abstract level (N=693)

Full text studies retrieved for detailed evaluation (N=48)

Studies excluded because of not meeting inclusion criteria (N=28)
- Review articles (N=22)
- Single arm study (N=4)
- Data based on same sample (N=2)

Hand searches (N=10)
- We found 7 RCTs (Nakamura et al 2011,[1] JapicCTI-050079,[2] Nakamura et al 2016,[3] NCT00097916,[4] NCT00476008,[5] NCT00768261,[6] NCT0093360[7]) from clinical trial registries
- We found 2 RCTs (Kitamura et al 2011,[8] Zheng 2011[9]) from review articles (Matsunaga et al 2015,[10] Tsoi et al 2016[11])
- We found 1 RCT (IE2201[12]) from the drug package insert[13]

RCTs included in the study (N=30)
- RCTs included in the meta-analysis (N=11)

Figure S1 PRISMA flow diagram.
Abbreviations: PRISMA, Preferred Reporting Items for Systematic Reviews and Meta-Analysis; RCT, randomized controlled trial; N, number of randomized controlled trials.
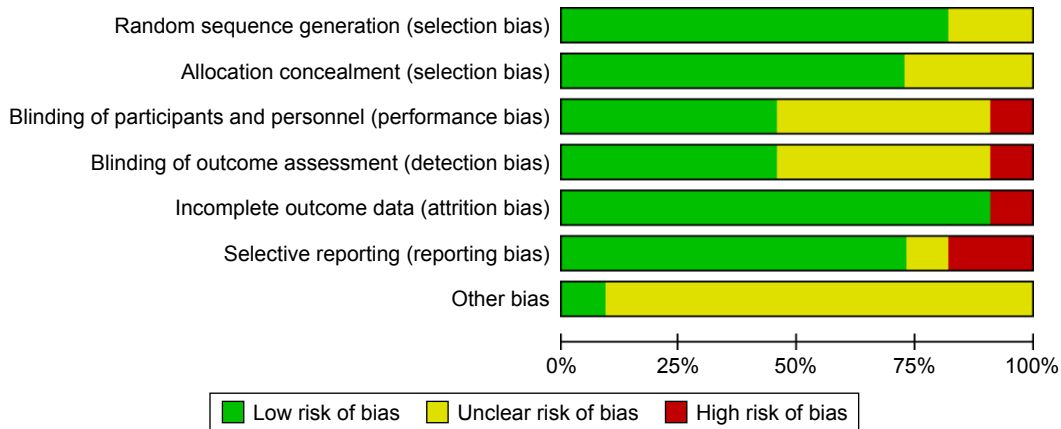
Figure S2 Risk of bias graph.

Figure S3 Risk of bias summary.

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