

Data supplement

Appendix DS1

Assessment of risk of bias

Each study was assessed (by two independent raters) on the following 13 items.

- 1 The source population is adequately described for key characteristics (choose one of the following possibilities)
 1. A selected sample from the general population
 2. Patients with heart disease
 3. Patients with cancer
 4. Patients with another somatic disorder
 5. Other clearly defined sample
 6. The source population is not clearly described
- 2 The sampling frame and recruitment are adequately described:
 - 2(a) The participants are recruited through
 1. the general population (with a clear description of the method)
 2. a medical setting, number of hospitals/institutes
 3. another clearly described method
 4. a method not clearly described
 - 2(b) The period of inclusion is clearly defined (at least the years are indicated)
 1. True
 2. Not true
 - 2(c) The geographical location of recruitment is clearly indicated (e.g. the name and city of the hospital, the name of the area)
 1. True
 2. Not true
- 3 Inclusion and exclusion criteria are clearly described
 1. True
 2. Not true
- 4 Is the study sample an adequate representation of the target population?
 1. Yes
 2. No
- 5 The baseline study sample (individuals entering the study) is adequately described for key characteristics
 1. Yes
 2. No
- 6 Mortality data at follow-up are available for at least 90% of the baseline sample
 1. True
 2. Not true
 3. Unclear
- 7 Attempts to collect information on participants who dropped out of the study are described
 1. Yes
 2. No drop-out
 3. No
 4. Unclear
- 8 Reasons for drop-out from baseline to follow-up are provided
 1. Yes
 2. No drop-out
 3. No
 4. Unclear
- 9 Participants who dropped out are adequately described in terms of key characteristics (including at least the number with depression)
 1. Yes
 2. No drop-out
 3. No
 4. Unclear
- 10 Mortality
 - 10(a) It is clearly reported what was done to establish the mortality status of participants
 1. True
 2. Not true
 - 10(b) The follow-up period for which mortality is measured is clearly described
 1. True
 2. Not true

- 11 Have the following confounders been measured?
 - 11(a) Demographic variables
 1. Yes
 2. No
 - 11(b) One or more lifestyle variables (smoking, body mass index, exercise)
 1. Yes
 2. No
 - 11(c) One or more illness-related variables (severity of the illness, somatic comorbidity, characteristics of the illness, etc.)
 1. Yes
 2. No
- 12 Have analyses been conducted to examine the influence of the confounders described in item 11 on the association between depression and mortality (usually through multivariable analyses)?
 1. Yes, all three groups of relevant confounders have been examined in multivariable analyses
 2. One or two groups of confounders have been examined in multi-variable analyses
 3. No confounder was included in the analyses
 4. No confounder was reported in question 11
- 13 The analyses have been conducted adequately. There are two possibilities:

This is a prospective study in a population. In these studies survival analyses are conducted.

This is a case–control study. In these studies logistic regression analyses have been conducted.

Have these analyses been conducted?

 1. Yes
 2. No

Scoring

After the rating the studies were scored on the main five main areas using the following rules.

Study participation (items 1–5)

Does the study sample represent the population of interest on key characteristics, sufficient to limit potential bias to the results?

Yes (5 items are positive)

Partly (3 or 4 items are positive)

No (0–2 items are positive)

Item 1 is positive if one of the answers 1–5 is given (6 is negative)

Item 2 is positive if 2(a) to 2(c) are all positive

Item 3 is positive when 1 is chosen

Item 4 is positive when 1 is chosen

Item 5 is positive when 1 is chosen

Study attrition (items 6–9)

Loss to follow-up (from sample to study population) is not associated with key characteristics (i.e. the study data adequately represent the sample), sufficient to limit potential bias

Yes (4 items are positive)

Partly (2 or 3 items are positive)

No (0 or 1 item is positive)

Unclear

Item 6 is positive when 1 is chosen

Item 7 is positive when 1 or 2 is chosen

Item 8 is positive when 1 or 2 is chosen

Item 9 is positive when 1 or 2 is chosen

Outcome measurement (item 10)

The outcome of interest is adequately measured in study participants to sufficiently limit potential bias

Yes (2 sub-items are positive)

Partly (1 sub-item is positive)

No (0 sub-item is positive)

Unclear

Item 10(a) is positive when 1 is chosen

Item 10(b) is positive when 1 is chosen

Confounding measurement and account (items 11 and 12)
 Important potential confounders are appropriately accounted for, limiting potential bias with respect to the prognostic factor of interest
 Yes (all three groups of confounders have been measured and accounted for; items 11(a)–(c) are positive and 1 is selected for item 12)
 Partly (one or two groups of confounders have been measured and accounted for; at least one of items 11(a)–(c) is positive and 2 is selected for item 12)
 No (all other ratings)
 Item 11(a) is positive when 1 is chosen

Item 11(b) is positive when 1 is chosen
 Item 11(c) is positive when 1 is chosen

Analysis (item 13)

The statistical analysis is appropriate for the design of the study, limiting potential for presentation of invalid results
 Yes (adequate analyses)
 No
 Unsure
 Item 13 is positive when 1 is chosen

Table DS1 Quality scores for the included studies ^a						
	Study participation	Study attrition	Outcome measurement	Confounding measurement	Analysis	Total quality score
Bush, 2001 ³⁶	+	+	+		-	3.5
Frasure-Smith, 1995 ³⁷	+	+	+	+	-	4.0
Fredman, 1989 ³⁸	+	+	+		-	3.5
Gallo, 1997 ³⁹	+	-			+	3.0
Janzing, 1999 ⁴⁰		-			-	1.5
Jiang, 2001 ⁴¹	+	+	+	-	+	4.0
Jorm, 1991 ⁴²			+	-	-	2.0
McCusker, 2006 ⁴³	+	+	+		+	4.0
Morris, 1993a ⁴⁴	+	-	+	+	-	3.0
Morris, 1993b ⁴⁵	+	-	+	-	-	2.0
Nakaya, 2006 ⁴⁶	+	-		+	+	3.5
Parakh, 2008 ⁴⁷	+	+	+	+	+	5.0
Penninx, 1999 ³⁵	+	-	+	+	+	4.0
Prieto, 2005 ⁴⁸	+	+		+	+	4.5
Rovner, 1991 ⁴⁹	+	+			+	4.0
Ryan, 2008 ⁵⁰	+			+	+	4.0
Schleifer, 1989 ⁵¹	+		+	-	-	2.0
Sharma, 1998 ⁵²	-			-	-	1.0
Sullivan, 2003 ⁵³	+	-	+		+	3.5
Von Ammon, 2001 ⁵⁵		+			-	2.5
Tilvis, 1998 ⁵⁴	-	-	-		+	1.5
Winkley, 2007 ⁵⁶		+	+	+	+	4.5

a. Key: +, positive (score 1); -, negative (score 0); , partly positive (score 0.5).

Table Ds2 Selected characteristics of studies examining excess mortality in major depressive disorder and subthreshold depression

Study	Patient group	Recruitment	Women %	Depressive disorder	Subthreshold depression	n	Follow-up period years	Country
Bush, 2001 ³⁶	Myocardial infarction patients	Consecutive patients admitted to hospital	41.7	MDD (SCID, DSM-III-R)	BDI \geq 10	271	0.33	USA
Frasure-Smith, 1995 ³⁷	Hospitalised MI patients	Patients admitted to hospital	21.6	MDD (DIS)	BDI \geq 10	222	1.5	Canada
Fredman, 1999 ³⁸	Adults	Community sample	65.0	MDD (DIS, DSM-III)	Minor depression (DIS)	1606	2	USA
Gallo, 1997 ³⁹	Adults (\geq 50 years)	Community sample (ECA Baltimore)	62.6	MDD (DIS, DSM-III)	Depression with sadness	1612	13	USA
Janzing, 1999 ⁴⁰	Dementia patients	Inhabitants of residential homes for the elderly	87.7	Depressive disorder (GMS/AGECAT)	Subthreshold depression (GMS/AGECAT)	73	1	The Netherlands
Jiang, 2001 ⁴¹	Congestive heart failure patients	Patients admitted to cardiology service at hospital	37.5	MDD (DIS)	BDI \geq 10	374	1	USA
Jorm, 1991 ⁴²	Older adults (\geq 70 years)	Community sample	NR	MDD (GMS/DSM-III)	Dysphoric mood (GMS)	228	5	Australia
McCusker, 2006 ⁴³	Older medical in-patients (\geq 65 years)	Patients admitted to intensive care or cardiac units of two hospitals	63.0	MDD (DIS, DSM-IV)	Minor depression (DIS, DSM-IV)	715	2.8	Canada
Morris, 1993a ⁴⁴	Stroke patients	Consecutive patients	40.7	MDD (PSE/DSM-III)	Minor depression (PSE, DSM-III)	91	10	USA
Morris, 1993b ⁴⁵	Stroke patients	Consecutive patients undergoing rehabilitation	52.9	MDD (CID, DSM-III)	Minor depression (CID, DSM-III)	82	1.25	Australia
Nakaya, 2006 ⁴⁶	Lung cancer patients	Patients with postoperative cancer	39.7	MDD (DSM-III-R, SCID)	POMS-d \geq 7	229	5.75	Japan
Parakh, 2008 ⁴⁷	Hospitalised MI patients	Patients with acute MI admitted to cardiology service at hospital	43.0	MDD and dysthymia (SCID, DSM-IV)	BDI \geq 10	208	8	USA
Peminx, 1999 ³⁵	Older adults (55-85 years)	Community sample (LASA)	51.7	MDD (DIS, DSM-III)	CEES-D \geq 16	3056	4.2	The Netherlands
Prieto, 2005 ⁴⁸	Haematological cancer patients	Patients recruited through hospital	41.7	MDD and minor depression (clinical interview; modified DSM-IV)	Minor depression (clinical interview)	199	5	Spain
Rovner, 1991 ⁴⁹	Nursing home residents	Consecutive admissions	77.3	Depressive disorder (M-PSE, DSM-III-R)	Depressive symptoms (M-PSE)	454	1	USA
Ryan, 2008 ⁵⁰	Older adults (\geq 65 years)	Community sample	60.8	MDD (DSM-IV, MINI)	CEES-D \geq 16	7363	4	France
Schleifer, 1989 ⁵¹	Patients with MI	Consecutive patients with MI admitted to one hospital	36.0	MDD (RDC criteria)	Minor depression (RDC)	283	0.25	USA
Sharma, 1998 ⁵²	Patients with depression compared with subthreshold and people without depression	Selection from community sample	70.6	Depressive disorder (GMS/AGECAT)	Subthreshold depression (GMS/AGECAT)	245	5	UK
Sullivan, 2003 ⁵³	Patients with stable coronary disease	HMO patients	17.2	MDD (DIS, DSM-IV)	Minor depression (DIS, DSM-IV)	199	5	USA
Tilvis, 1998 ⁵⁴	Older adults (65, 75, 80, 85 years)	Community sample	NR	Depressive disorder (interview according to DSM-III)	SDS \geq 40	1330	5	Finland
Von Ammon Cavanaugh, 2001 ⁵⁵	Medical in-patients	Consecutive admissions to one hospital	34.9	MDD (SADS/DSM-IV)	Minor depression (SADS/DSM-IV)	151	NR	Brazil
Winkley, 2007 ⁵⁶	Patients with first diabetic foot ulcer	Community chiropody and hospital foot clinics	36.4	MDD (SCAN, DSM-IV)	Minor depression (SCAN, DSM-IV)	253	1.5	UK

AGECAT, Automated Geriatric Examination for Computer Assisted Taxonomy; BDI, Beck Depression Inventory; CES-D, Center for Epidemiological Studies Depression scale; CID, Composite International Diagnostic Interview; DIS, Diagnostic Interview Schedule; ECA, Epidemiologic Catchment Area; GMS, Geriatric Mental State; HMO, health maintenance organisation; LASA, Longitudinal Aging Study Amsterdam; M-PSE, Modified Present State Examination; MDD, major depressive disorder; MI, myocardial infarction; MINI, Mini International Neuropsychiatric Interview; NR, not reported; POMS-d, Profile of Mood States Depression scale; PSE, Present State Examination; RDC, Research Diagnostic Criteria; SADS, Schedule for Affective Disorders and Schizophrenia; SCAN, Schedules for Clinical Assessment in Neuropsychiatry; SCID, Structured Clinical Interview for DSM; SDS, Self-Rating Depression Scale.