Early Career Psychiatrists

It is illegal to post this copyrighted PDF on any website. Depression, Cognitive Functions, and Impaired Functioning in Middle-Aged Adults From the CONSTANCES Cohort

Hélène Vulser, MD, PhD^{a,b,*}; Emmanuel Wiernik, PhD^c; Nicolas Hoertel, MD, PhD^{a,b,d}; Maria Melchior, PhD^{e,f}; Mura Thibault, MD, PhD^{c,g,h}; Romain Olekhnovitch, PhD^c; Philippe Fossati, MD, PhD^{i,j}; Frédéric Limosin, MD, PhD^{a,b,d}; Marcel Goldberg, MD, PhD^{a,c}; Marie Zins, MD, PhD^{a,c}; and Cédric Lemogne, MD, PhD^{a,b,d}

ABSTRACT

Objective: This large-scale population-based prospective study examined the association between depressive symptoms and cognitive performance at baseline with later functioning in middle-aged adults.

Methods: The Center for Epidemiologic Studies Depression Scale, the Digit Symbol Substitution Test (DSST), the Trail Making Test B (TMT-B), and the Semantic Verbal Fluency test (SVF) were completed at baseline by 7,426 participants aged \geq 45 years from February 2012 to December 2013. Role limitations and social functioning were later assessed with the second version of the 12-Item Short Form Health Survey. The association between depressive symptoms and cognitive performance at baseline with functioning at follow-up was examined using general linear models and mediation analyses including sex, age, education, alcohol intake, and cannabis use as covariates.

Results: Altered functioning at follow-up was predicted by depressive symptoms (β per standard deviation [95% confidence intervals]: -1.10 [-1.16 to -1.03] and -1.02 [-1.08, -0.96] for role limitations and social functioning, respectively) and DSST, TMT-B, and SVF performance (for role limitations: 0.11 [0.09 to 0.14], -0.11 [-0.13 to -0.08], and 0.03 [0.01 to 0.06], respectively; for social functioning: 0.10 [0.07 to 0.12], -0.08 [-0.11 to -0.06], and 0.04 [0.01 to 0.05], respectively) at baseline. Depressive symptoms were associated with poorer cognitive performance at baseline (-0.19 [-0.25 to -0.13], 0.15 [0.08 to 0.21], and -0.11 [-0.17 to -0.04], respectively). Cognitive performance accounted for only 0.3%-1.4% of the relationship between depressive symptoms and functioning. In contrast, depressive symptoms accounted for 19.5%-43.7% of the association between cognitive performance and functioning.

Conclusions: In middle-aged adults from the general population, cognitive impairment is unlikely to substantially explain the association between depressive symptoms and later role limitations and social functioning.

J Clin Psychiatry 2018;79(6):17m12003

To cite: Vulser H, Wiernik E, Hoertel N, et al. Depression, cognitive functions, and impaired functioning in middle-aged adults from the CONSTANCES cohort. *J Clin Psychiatry*. 2018;79(6):17m12003.

To share: https://doi.org/10.4088/JCP.17m12003 © Copyright 2018 Physicians Postgraduate Press, Inc.

^aParis Descartes University, Sorbonne Paris Cité, Faculty of Medicine, Paris, France ^bAP-HP, West Paris University Hospitals, Department of Psychiatry, Paris, France ^cINSERM, Population-based Epidemiological Cohorts Unit, UMS 011, Villejuif, France ^dINSERM U894, Psychiatry and Neurosciences Center, Paris, France

^eINSERM, UMR_S 1136, Pierre Louis Institute of Epidemiology and Public Health, Department of Social Epidemiology, Paris, France

^fSorbonne Universités, UPMC Univ Paris 06, UMR-S 1136, Pierre Louis Institute of Epidemiology and Public Health, Department of Social Epidemiology, F-75013 Paris, France

⁹INSERM, U1061, Neuropsychiatry: Epidemiological and Clinical Research, Montpellier, France

^hMontpellier University Hospital, Montpellier University, Montpellier, France ⁱAP-HP, Pitié-Salpêtrière Hospital, Department of Psychiatry, Paris, France ^jINSERM U 1127, CNRS UMR 7225, Sorbonne Universités, UPMC Univ Paris 06, UMR S 1127, Institut du Cerveau et de la Moelle, ICM, Social and Affective Neuroscience (SAN) Laboratory, Paris, France

*Corresponding author: Hélène Vulser, MD, PhD, Unité de Psychologie et Psychiatrie de liaison et d'urgence, Hôpital Européen Georges Pompidou, 20 rue Leblanc, 75908 Paris Cedex 15, France (helene.vulser@aphp.fr).

epression affects 350 million people worldwide and is a major contributor to the overall global burden of disease.¹ Depression is strongly associated with functional impairment,² which has been found to be comparable to or worse than that associated with several major chronic medical conditions (eg, diabetes, arthritis, angina).³ Impaired subjective performance has been reported in widespread domains of functioning, such as household; work; relationships with partners, family members, and friends; and leisure activities.4,5 These impairments in functioning are strongly associated with the severity of depressive symptoms^{6,7} and reduced quality of life.⁸ In addition, depression may contribute to objective social damage, such as work absenteeism, presenteeism, and unemployment.^{7,9} Several studies have reported a significant positive effect of antidepressant treatments on quality of life and on social and work functioning,¹⁰ although residual functional impairment is frequent in patients who achieved remission.¹¹ However, the mechanisms underlying the association between depression and functioning remain largely unknown. In recent decades, it has been suggested that cognitive dysfunctions associated with depression might mediate its impact on functioning.^{12,13} Indeed, a broad range of cognitive domains may be affected in depressed patients, such as information processing speed, verbal fluency, working memory, attentional control, and cognitive inhibition.^{14,15} Such cognitive impairments, which all relate to some extent to executive function, have even been found in patients during their first depressive episode.^{16,17} While cognitive function may improve after pharmacologic treatment of depression,^{18,19} deficits can still be detected in euthymic patients,^{20,21} which might explain persistent functional impairment in remission.

Several studies have reported an association between objectively assessed cognitive dysfunction and functional impairment in patients with depression.^{12,13} For example, Jaeger et al²² reported that several cognitive measures were associated with disability 6 months following

Vulser et al It is illegal to post this copyrighted PDF on any website.

 Depression is associated with both cognitive and functional impairment.

nical Points

- In middle-aged adults from the general population with depressive symptoms, cognitive impairment is unlikely to substantially explain altered functioning.
- Interventions aimed at reducing the functional impairment associated with depression should primarily target depressive symptoms themselves; such interventions are likely to improve cognitive functioning at the same time.

hospitalization for a major depressive episode in 48 patients; in addition, 6-month cognitive performance was strongly associated with self-reported functioning after adjusting for residual depression. In 21 adults treated for depression, Naismith et al²³ found a moderate relationship between objectively measured psychomotor speed and physical disability, even after adjusting for depression severity. In 52 inpatients with depression, Withall et al²⁴ found that poor event-based prospective memory and more perseverative errors on the shortened Wisconsin Card Sorting Test at admission predicted worse social and occupational outcomes at a 3-month follow-up. Considering occupational status, Baune et al²⁵ reported that, among 70 patients with depression, those who were unemployed had poorer results on neuropsychological tests than those who were employed. Finally, in 483 currently non-depressed patients with major depressive disorder receiving selective serotonin reuptake inhibitors, improvements in cognitive performance were found to predict improvements in functioning.²⁶

Overall, these studies are consistent with the hypothesis that cognitive impairment may account for a substantial part of functional limitation in patients with depression. However, in contrast with more severe mental disorders such as schizophrenia or bipolar disorder,²⁷ the evidence supporting this hypothesis remains weak. Most of the aforementioned studies were based on relatively small, highly selected samples; few had a longitudinal design; and none performed formal mediation analyses.^{12,28,29} In a cross-sectional survey including 21,425 adults from 6 European countries, Buist-Bouwman et al³⁰ reported that more than 25% of the association of depression with role functioning was directly attributable to self-reported cognitive complaints (ie, concentration, attention, and memory problems). However, that large-scale study did not use objective measures of cognitive function or have a longitudinal design. Besides, due to its associations with both impaired cognition and altered functioning, depression is a plausible confounding factor that may partly explain the association between cognitive and functional impairment.

In this study, we used data from CONSTANCES, a French large-scale population-based study, to investigate the prospective associations between depressive symptoms and cognitive functions with later functioning in middleaged adults.

Participants

All participants were recruited from the CONSTANCES cohort (www.constances.fr). This project aims at providing a general prospective cohort of a large sample of the French population aged 18-69 years.^{31,32} Participants were recruited since 2012 among people affiliated to the main national health insurance provider, which covers more than 85% of the French population. The cohort was designed to be representative of the target population according to age, sex, employment status, and occupational class. A random sample from the target population was invited by mail to join the cohort. Those who agreed had to fill out self-administrated questionnaires dealing with lifestyle, health, physical limitations, and social and personal characteristics. They were invited to go to one of the 21 participating Health Screening Centers throughout France to benefit from an extensive health examination (medical and paraclinical examinations, blood tests). In addition, cognitive tests were performed for those aged 45 years or older. A follow-up self-administered questionnaire was then completed annually by the participants at home, using either paper or web-based questionnaires.

In the present study, we used data from the participants aged 45 years or older included from February 21, 2012, to December 31, 2013. Eligibility criteria were being able to fill out the study questionnaires, ability to speak French, and having no missing data for selected variables, including assessment of functioning at follow-up in 2014 (Supplementary Figure 1).

All confidentiality, safety, and security procedures were approved by the French legal authorities. In accordance with French regulations, the CONSTANCES cohort project obtained the authorization of the National Data Protection Authority (Commission Nationale de l'Informatique et des Libertés). Written informed consent was obtained from all participants.

Assessment of Depressive Symptoms at Baseline

Depressive symptoms were measured at baseline with the French version of the Center for Epidemiologic Studies Depression Scale (CES-D).^{33,34} The total score ranged from 0 (no depressive symptom) to 60. We used a cutoff score of 19 (CES-D score \geq 19 versus < 19) to define depression status³⁴ (Appendix 1).

Assessment of Cognitive Functions at Baseline

Cognitive functions were assessed at baseline using objective neuropsychological tests for which impaired performance has been previously reported in patients with major depression¹⁵: Digit Symbol Substitution Test (DSST),³⁵ Trail Making Test part B (TMT-B),³⁶ and Semantic Verbal Fluency test (SVF)³⁷ (Appendix 1). TMT-B score was log transformed to achieve a close-to-normal distribution. All of these tests engage executive function to some extent.

It is illegal to post this copyrighted

in Multivariate Models $(N = 7,426)^a$

Follow-Up

at

Baseline With Role Limitations and Social Functioning

able 1. Associations of Depression Status and Cognitive Performance at

The second version of the Short-Form-12 Health Survey $(SF-12v2)^{38-40}$ was part of the annual follow-up questionnaire sent in 2014 to all participants (Appendix 1). Mean (SD) duration between baseline assessments and reception of the 2014 follow-up questionnaire was 497 (157) days. Because we were interested in functional impairment associated with depression specifically, we decided a priori to use 2 subscales as primary outcomes, as Spijker et al⁴¹ did: role limitations due to emotional problems, henceforth referred to as "role limitations" (score range, 2–10), and social functioning (score range, 1–5). For both scales, a higher score corresponds with better functioning.

Other Covariates

Other covariates included age at baseline, sex, year of inclusion, education level (no diploma, lower secondary education, professional education, upper secondary education, bachelor, fourth-year university level, master's degree or higher, other), alcohol intake frequency (never, once or less than once in a month, 2 or 3 times in a month, once or more in a week), and lifetime cannabis use. Education level, alcohol intake frequency, and lifetime cannabis use were assessed at baseline.

Statistical Analysis

Cognitive and functioning scores were z-transformed. The relationships between variables were assessed within the framework of generalized linear models (GLMs) using R software (http:// cran.r-project.org, version 3.3.1). First, analyses were conducted including role limitations and social functioning at follow-up as the dependent variables and depression status at baseline as the independent variable. Sex, age, year of inclusion, education level, alcohol intake frequency, and lifetime cannabis use at baseline were entered as covariates. The association between depression status (as a binary variable) and each cognitive score at baseline was also assessed. Then, each of the 3 cognitive scores (DSST, TMT-B, SVF) was entered as the independent variable instead of depression status in 3 separate models. Finally, both depression status and cognitive scores were entered in the 3 separate models. GLM coefficients were presented per standard deviation of the SF-12v2 subscale.

To examine whether changes in regression coefficients across the aforementioned models were statistically significant, formal mediation analyses were conducted with functioning scores as the dependent variable based on algorithms devised by Imai et al.⁴² Sex, age, year of inclusion, education level, alcohol intake frequency, and lifetime cannabis use at baseline were entered as covariates.

У	r	0	Jh	t	e		P.				C		n	ē			У	V	V	e	D		5	t	e.	
	%Clf				Ref		10 21 +0 11 00						Ref			12.09 to 58.12	d 3c include	alcohol intake							formed)	
VF	%Med ^g				Ref		C7 61	7/174					Ref			29.39	i 3a, 3b, an	tion level,							loa-transf	5
S	Cld				0.01 to 0.06		0.01 +0.0.01						0.01 to 0.05			0.01 to 0.05	Model 2c. Models	inclusion, educa							Test B (total time	
	β ^c				0.03		NCOO	70.0					0.04			0.03	nd SVF in I	ge, year of							il Making	2
	%Clf				Ref		11.17 to 28.33					Ref			14.11 to 35.09		-B in Model 2b, ai	djusted for sex, aç			2v2).				core). TMT-B=Tra	
T-B	%Med ^g				Ref		19.53					Ret			23.45		lel 2a, TMT-	vere also ac			urvey (SF-1				est (total so	
TM	Cld				-0.13 to -0.08		–0.11 to –0.06					-0.11 to -0.06			–0.09 to –0.04		el 1, DSST in Moc	l 3c). All models v		:	ort Form Health S				- Verhal Fluency t	
	β ^c				-0.11		-0.09					-0.08			-0.06		tus in Mod	VF (Mode		1	2-item Sho				= Semantic	
	%Clf			Ref		19.47 to 36.93					Ket			20.19 to 40.17			vs: depression sta	(Model 3b), and S			d version of the 1		as mediator.		s as mediator.	
DSST	%Med ^g			Ref		26.77					Ket			29.11			e as follow	3a), TMT-B		ļ	the secon		ve scores à		sion status t_Ref=refe	
	Cld			0.09 to 0.14		0.06 to 1.05					0.0/ to 0.12			0.04 to 0.09			lent variables ai	r DSST (Model			the subscale of		es using cogniti		es using depres = not significan	
	β			0.11		0.08				0	0.10			0.07			depend	usted fc		ore ≥ 1	tion of		analyse		analys re) NS	
	%Clf	nt Variable	Ref			0.93 to 2.21	0.64 to 1.82		dent Variable	Ref				0.85 to 1.97	0.49 to 1.46	0.07 to 0.71	when noted. In	le and were adji		Scale (CES-D) sc	r standard devia	parameter.	on in mediation	on proportion.	on in mediation	
n Status ^b	%Med ^e	Depende	Ref			1.44	1.19 0.15 ^{NS}		the Depen	Ref				1.31	0.92	0.33	11), except	lent variab	Ise.	epression	ficient) per	estimated	ria mediati	e mediatic	via mediati Substitutio	
Depressio	Cld	Limitations as the	-1.16 to -1.03			-1.14 to -1.02	-1.14 to -1.02		al Functioning as l	–1.08 to –0.96				–1.07 to –0.94	–1.07 to –0.95	–1.08 to –0.95	significant (P<.00	is as the independ	ifetime cannabis u	niologic Studies D	ameter (GLM coef	ice interval of the	on of total effect v	ence interval of th	on of total effect v	
	β ^c	Jsing Role	-1.10			-1.08	-1.08		Jsing Socia	-1.02				-1.01	-1.01	-1.02	istics were	ssion statu	ency, and li	for Epiden	mated par	% contider	= proportic	5% confid∈	= proportic	
	lodel	Aodel L		e.	a v	a	م ہ	: - 	lodel (e .	q	U,	a	q	U	All stat	depre	freque	Center	β=esti	CL=95	%Med	%CI = 9.	%Med	

You are prohibited from making this PDF publicly available

Vulser et al It is illegal to post this First, depression-status at baseline was considered opyrighted PDF

as the independent variable, and cognitive scores at baseline (DSST, TMT-B, and SVF separately) as the "mediator" variables between depression status at baseline and functioning at follow-up. Then, each cognitive score was entered as the independent variable with depression status at baseline as the "mediator" variable between each cognitive score at baseline and functioning at follow-up. These mediation models were fit with GLM, and output objects were bootstrapped 500 times with replacement using a parametric mediational analysis. In mediation analysis, a significant mediating effect is defined by a 95% confidence interval (CI) of the regression coefficient that does not include zero.42

To examine the robustness of our findings, we also carried out sensitivity analyses. We performed similar analyses (1) using CES-D as a continuous score, taking the interval between the 25th and the 75th percentile as the unit to provide clinically meaningful regression coefficients; and (2) using a more restricted definition for depression requiring both CES-D score \geq 19 and self-reported limitation at inclusion. Self-reported limitation at inclusion was defined as having answered "yes" to "Have you been limited, for at least 6 months, in your routine activities by a health problem?" and then having answered "depressive state" to "If yes, for what reasons?"

RESULTS

The final study population consisted of 7,426 participants (3,551 men, 47.82%) with a mean (SD) age of 57.79 (7.20) years. Study population selection is described in Supplementary Figure 1. The mean (SD) CES-D score was 9.88 (8.35) (range, 0-53); 13.24% of participants (n = 983) were depressed at baseline (CES-D score \geq 19) (Supplementary Table 1). Characteristics of participants lost to follow-up and comparisons with the study population are displayed in Supplementary Table 2.

First, after adjustment for covariates, depression status was significantly associated with each of the 3 cognitive scores at baseline as expected (DSST: $\beta = -0.19$; 95% CI, -0.25 to -0.13; P < .001; TMT-B: $\beta = 0.15$; 95% CI, 0.08 to 0.21; P<.001; SVF: $\beta = -0.11$; 95% CI, -0.17 to -0.04; P = .001) and with both functioning scores (role limitations and social functioning) at follow-up (Table 1). Second, each of the 3 cognitive scores at baseline was significantly associated with the 2 functioning scores at follow-up as expected (Table 1). Third, after further adjustment for each cognitive score, the relationship between depression status and functioning remained virtually unchanged, and

Figure 1. Graphic Representation of Mediation Analyses^a

A. Cognitive Scores as Mediator



^aCognitive scores served as a mediator of the relationship between depression status and role limitations/social functioning at follow-up (A). Depression status served as a mediator of the relationship between cognitive scores and role limitations/social functioning at follow-up (B). Covariates are sex, age, year of inclusion, education level, alcohol intake frequency, and lifetime cannabis use.

^bProportions of mediated effect are listed as follows: top row: models using DSST/ TMT-B/SVF total scores, with role limitations as dependent variable; bottom row: models using DSST/TMT-B/SVF total scores, with social functioning as dependent variable. ***P<.001

Abbreviations: DSST = Digit Symbol Substitution Test (total score), NS = not significant (P > .05), SVF = Semantic Verbal Fluency (total score), TMT-B = Trail Making Test B (total time).

mediation analyses showed that cognitive scores at baseline accounted for only 0.3%-1.4% of the relationship between depression status at baseline and functioning at follow-up (Table 1, Figure 1). In contrast, depression status at baseline accounted for 19.5%-43.7% of the relationship between cognitive at baseline and functioning scores at follow-up (Table 1, Figure 1).

In sensitivity analyses based on continuous CES-D scores, depressive symptoms were also associated with each of the 3 cognitive scores (DSST: $\beta = -0.10$; 95% CI, -0.12 to -0.07; P < .001; TMT-B: $\beta = .09$; 95% CI, 0.07 to 0.12; P < .001; SVF: $\beta = -0.06$; 95% CI, -0.08 to -0.03; P<.001) and with both functioning scores (Table 2). The association between depressive symptoms and the 2 functioning scores remained virtually unchanged after further adjustment for each cognitive score, which accounted for only 0.2%-1.1% of this relationship (P<.001 considering DSST and TMT-B, not significant for SVF) (Table 2). In contrast, continuous CES-D scores accounted for 42.5%-85.3% of the relationship between cognitive and functioning scores (all *P* < .001) (Table 2).

When a more restricted definition of depression status is used, combining both CES-D score \geq 19 and self-reported limitation (n = 205, ie, 2.76% of the total sample), the association of depression status with each of the cognitive scores (DSST: $\beta = -0.34$; 95% CI, -0.47 to -0.22; P < .001; TMT-B: $\beta = .29$; 95% CI, 0.16 to 0.42; P < .001; SVF: $\beta = -0.25$; 95% CI, -0.38 to -0.11; P < .001) and with both functioning scores (Table 3) strengthened. However, cognitive scores accounted for only

Depression, Cognition, and Functional Impairment of the relationship betw

scores and functioning, whereas depression status still explained 13.3%-33.8% of the relationship between cognitive and functioning scores (all *P*<.001) (Table 3).

DISCUSSION

This prospective large-scale populationbased study aimed to investigate the association of depressive symptoms and cognitive performance at baseline with both role limitations and social functioning at follow-up in adults aged 45 years or older. We found that the association between depressive symptoms and later functioning was not substantially explained by cognitive performance, regardless of the definition of depression (ie, binary or continuous), the cognitive test (ie, DSST, TMT-B or SVF), or the functioning variable (ie, role limitations or social functioning). In contrast, depression explained a substantial proportion of the association between cognition at baseline and functioning at follow-up.

Strengths of the study are the large size, the population-based sample, the prospective design, and the use of objective measures of cognitive functions. Thanks to a sufficient statistical power and standardized neuropsychological tests, our results were consistent with decades of literature linking depressive symptoms with both cognitive^{14,15} and functional^{2-6,8-11} impairment. However, it is noteworthy that this literature did not examine the contribution of cognitive function or functional impairment associated with depression. To our knowledge, our study might indeed be the first to explore this issue in a large prospective sample.

Some limitations should also be acknowledged. First, the population study is not representative of the general population and was confined to participants aged 45 to 69 years. Thus, our results cannot be generalized to younger or older adults. Second, the duration of the follow-up was short. However, the majority of the studies on this topic have been crosssectional. Third, the diagnosis of depression was based on a self-report scale rather than on a standardized interview. However, sensitivity analyses using CES-D as a continuous score or a more restricted definition of depression status (including both CES-D score \geq 19 and selfreported limitation at baseline) yielded similar results. Fourth, as in other studies on this topic,⁴⁰ functioning was measured with a selfadministered questionnaire (ie, the SF-12v2).

at Eollow-Up in Multivariate Social Functioning Role Limitations at Raceline ICES-D Sco Accortations

Fr TMT-B SVF SUF SUF <th></th>	
$ \frac{100}{100} \frac{1}{100} \frac$	
$ \frac{19}{3} \frac{\% C f}{6} \frac{\beta'}{6} \frac{C l^d}{6} \frac{\% M e d^9}{6} \frac{\% C l^d}{6} \frac{\beta'}{6} \frac{C l^d}{6} \frac{\% M e d^9}{6} \frac{\% C l^d}{6} \% $	SST
Ref -0.11 -0.13to -0.08 Ref Ref 0.03 0.01to 0.05 Ref Ref 3 35.80 to 57.00 -0.06 -0.08to -0.04 42.51 33.01 to 54.26 0.03 0.01 to 0.03 Ref Ref -0.06 -0.08to -0.04 42.51 33.01 to 54.26 0.00 ^{NS} -0.01 to 0.03 85.28 50.00 to 566.7 Ref -0.06 -0.08to -0.04 42.51 33.01 to 54.26 0.001 0.001 666.7 8 -0.06 -0.0104 42.51 33.01 to 54.26 0.001 0.001 666.7 8 -0.06 -0.0104 42.51 33.01 to 54.26 0.004 Ref Ref 9 -0.056 Ref 0.04 0.02 to 0.006 Ref Ref 9 -0.066 to -0.02 53.35 38.32 to 91.36 0.02 to 0.004 55.69 36.78 to 109.99	6Med ^g %Cl ^f β^c 6
$\frac{\mathrm{Ref}}{3 35.80 \mathrm{to} 57.00} -0.11 -0.13 \mathrm{to} -0.08 \\ -0.06 -0.08 \mathrm{to} -0.04 \\ -0.06 -0.08 \mathrm{to} -0.04 \\ -0.01 \mathrm{to} -0.03 \\ -0.04 \mathrm{to} -0.01 \mathrm{to} -0.03 \\ -0.01 \mathrm{to} -0.04 \\ -0.00 \mathrm{to} -0.04 \\ -0.04 \mathrm{to} -0.04 \$	
$ \begin{array}{c ccccccccccccccccccccccccccccccccccc$	
$3 35.80 \text{ to } 57.00 \\ -0.06 -0.08 \text{ to } -0.04 \\ 42.51 \\ 33.01 \text{ to } 54.26 \\ 0.00^{\text{NS}} \\ -0.01 \text{ to } 0.03 \\ 8.2.8 \\ 5.00 \text{ to } 566.79 \\ 8.2.00 \text{ to } 566.79 \\ $	Ref Ref –0.11 –0.13
$ \begin{array}{cccccccccccccccccccccccccccccccccccc$	
-0.06 -0.08 to -0.04 42.51 33.01 to 54.26 0.00 ^{NS} -0.01 to 0.03 85.28 50.00 to 566.79 Ref -0.08 -0.11 to -0.06 Ref Ref 0.04 0.02 to 0.06 Ref Ref 0.04 -0.02 to 0.06 36.78 to 109.99	5.53 35.80 to 57.00
Ref -0.08 -0.11 to -0.06 Ref 0.04 0.02 to 0.06 Ref Ref 5 36.38 to 65.42 -0.04 -0.06 to -0.02 53.95 38.32 to 91.36 0.02 0.00 to 0.04 55.69 36.78 to 109.99	-0.06 -0.08
Ref -0.08 -0.11 to -0.06 Ref Ref 5 36.38 to 65.42 -0.04 -0.06 to -0.02 53.95 38.32 to 91.36 0.02 0.00 to 0.04 55.69 36.78 to 109.99	
Ref -0.08 -0.11 to -0.06 Ref 0.04 0.02 to 0.06 Ref Ref 5 36.38 to 65.42 -0.04 -0.06 to -0.02 53.95 38.32 to 91.36 0.02 0.00 to 0.04 55.69 36.78 to 109.99	
–0.08 –0.11 to –0.06 Ref Ref 0.04 0.02 to 0.06 Ref Ref 36.38 to 65.42 –0.04 –0.06 to -0.02 53.95 38.32 to 91.36 0.02 0.00 to 0.04 55.69 36.78 to 109.99	ef Ref
5 36.38 to 65.42 -0.04 -0.06 to -0.02 53.95 38.32 to 91.36 0.02 0.00 to 0.04 55.69 36.78 to 109.99	-0.08 -0.11
–0.04 –0.06 to –0.02 53.95 38.32 to 91.36 0.02 0.00 to 0.04 55.69 36.78 to 109.99	1.05 36.38 to 65.42
0.02 0.00 to 0.04 55.69 36.78 to 109.99	-0.04 -0.06
ויט (וווטטבו טט), מווט טייו (וווטטבו טכן. הזו וווטטבו אבוב מוטט מטןמזבט וטו זכה, מטב, זכמו טו ווורוטטיטו, בטטכמיטו ובעבן, מרטוטו וווגמאב	een the 25th and the 75th percentile (9
re (model bd), and by (model bd). An inddeb were also adjusted for sex, age, year of inclusion, education revel, arcorof intake en the 25th and the 75th percentile (9.56) as unit.	
ten the 25th and the 75th percentile (9.56) as unit.	ores as mediator.
ten the 25th and the 75th percentile (9.56) as unit.	
ten the 25th and the 12-item Short Form Health Survey (5F-12v2). cond version of the 12-item Short Form Health Survey (5F-12v2). es as mediator.	status as mediator. = reference value, SVF = Semantic Verba

Vulser	et	al											
.lt i	S		le	aal	to	p	09	st t	his	copyric	ahte	ec	d PDF on any website.
				9	22.62				8.39	a a			In particular, one may argue that the items
		6Cl ^f		Ref	10 J			Ref	to 4	inta			that were selected a priori from the SF-12V2
lels		0`			5.70				0.63	c inc	ned)		symptoms so that there might be little room
100					16				,	, alc	sforr		for a mediating effect of objective cognitive
te N		edg		ų.				f	38	ib, al evel	trans		functioning. However, it should be noted that
aria	F	WW%		Å	33.			Re	21.	3a, 3 ion l	log-1		not only depressive symptoms but also cognitive
tiva	S			ſ) 4			~	9	dels ucati	me,		functions were associated with the 2 SF-12v2
Mul				0.0	0.0			0.0	0.0	Mod , edu	al tii		subscales, suggesting that these measures were
. 드		Ū		01 tc	01 tc			02 tc	01 tc	l 2c. sion	tot		sensitive enough to capture relevant effects.
ļ				0.0	-0.0			0.0	0.0	lode	est B		Furthermore, depressive symptoms did account
Ň					2					in N rofi	ng T		for a substantial part of the association between
R		й		03	.02 ^N			.04	.04	SVF yea	Aaki		cognitive functions and these 2 subscales.
at				C	0			0	0	and age,	rail N		Therefore, the lack of mediation effect by
ing					9.84				2.47	2b, a sex, a	3= Tr		cognitive functions is unlikely to be explained
ion		%Cl ^f		Ref	to 1			Ref	to 2	for s	MT-E		by the subjective versus objective nature of the
D L					7.63				9.16	n Mc sted 2).	e), T		for intelligence quotient (IO). Future studies
ЦЦ									0.	T-B i adju -12v	scor		would benefit in reproducing our results while
ocia		Aed ^g		Ref	3.32			lef	5.07	, TM Ilso a Iso a	otal		additionally adjusting for IO, particularly for
s/S	8-1	% ∧			<u>(1)</u>			ш	1	el 2a ere <i>i</i> irvey	st (t		cognitive tests that are closely related to IQ. For
ion	Ľ⊻			08	07			06	05	Aode Is w h Su	cy te		example, in older adults, TMT score has been
itat		ъ		-0-	.0-			-0-	-0-	in N node Healt	nen		found to be more strongly associated with IQ
Ē		Ū		3 tc	2 tc			1 tc)9 tc	All m tm H	al Fl		than education level. ⁴³ However, IQ may also be
ole				-0.1	-0.1			-0.1	-0.0	il 1, [3c). , t Fo	Verb		considered as the composite of neurobehavioral
h R				-	6			œ		lode Shor	ntic		abilities assessed in neurocognitive tests. ⁴⁴ In
Wit		ğ		-0.1	-0.0			-0.0	-0.0	in Mo (Mo	ema		this study, analyses were adjusted for education
ne					_			·		atus SVF 12-it	= = S(level, which has been found to be positively
seli					23.21				53.3	and and the	, SVF		associated with both IQ and neuropsychological
t Ba		%Cl ^f		Ref	to		2	кег	to	essic 3b), 3b), n of ator.	liato alue		study ⁴⁶ reported a positive association between
e ai					0.92				1.05	lepre odel ersio	mec ce v		education level and cognitive association between
and					-				-	ws: c 8 (Mc tion tion as m	is as eren		on more cognitively complex tests such as the
L E		1ed ⁹		lef	.69		4	eI	.89	ollov MT-E mita ecor ores	statu = ref		TMT-B or the DSST, in contrast with more
erfo	SSC	% ∧		œ	16		6	r	16	e as f a), Tl a), Tl a)	ion s Ref		simple tests such as the TMT-A. Thus, schooling
e P				4	5		ć	7	-	s are del 3. borte porte i of t	ressi ant,		may foster the development of cognitive
Ē		p		0 0.1	0 0.1		Ċ	 0	0 0.	able Moc If-rej scale cog	dep Inific		processes that underpin performance on IQ.
ogr		0		.09 t	.07 t		5	1/0.	.06 t	vari SST (SST (Sub: sub: sing	sing ot sig		This study confirms results from previous
0 P				0	0		¢	^o	0	fent or DS 9 an the es us	es u: = no		ones ^{2-6,8-11} of a strong association between
s an		ğ		.1	60.		, ,	01.	.08	oenc ed fo ed fo s≥1 n of n of	alys , NS		depression and later altered functioning. We
atu:				0	0		Ċ	0	0	ndel Jjust score iatio	n ar core)		limitations were significantly altered up to
St			d)		3.04 2.82 0.75	ble			3.08 2.36 1.22	ed. l re ac -D) dev er. iatio	ion. liatic tal se		24 months after depression assessment. The
sior		%Cl ^f	iable	Iau	4 to 2 to	/aria	Ref		2 to 0 to	I we (CES dard nete	port med t (to		strength of these associations was similar to
res	1s ^b		t Var		1.2 -0.0	ent /			0.2	/hen : anc cale stanc aarar	n in Tes		the figures obtained by Spiker et al ⁴¹ with the
Dep	statu		den			ende				pt w able on So per s ed p ation	ation		36-item Short Form Health Survey in individuals
ed [on	Med [®]	hen	ē	07 78 31 ^{NS}	Dep	Ref		.99 .61	exce vari essic essic imat imat	iedia nedi stitu		with major depression. Furthermore, we found
ict	ressi	₩	e De	-	0 7	the	-		0	01), dent use. Depr fficie e esti via n	ne m via n Sub		a significant negative association between
est	Dep		is th	.	6464	g as	24		22 25	 <.0 ><.0 abis abis ies [ies [the f the fect 	of tl fect nbol		CES-D scores as a continuous measure and
of B	ted	σ	ons a			onin	-1-			int (F anna Stud GLM al efi	erval al ef t Syn		later functioning, suggesting that, as previously
su	stric	D	tatic	22 10	56 tc 56 tc 59 tc	nctio	51 to		48 to 49 to 50 to	ifica the c ne c ogic ; ter (ter (tot	inte f tot Digii		reported, ^o these impairments in functioning
atio	Re		Limi	- - 		I Fu			$\frac{1}{1}$ $\frac{1}{1}$ $\frac{1}{1}$	sign s as fetir iolc ame ame ce ii	ence on of 6T=1		were associated with the severity of depressive
oci			Role	0	щщы	Socia	80		2 9 1	vere statu ind l iden pari fider fider	nfide vortie : DS		This study also confirms and avtands in
Ass (6) ^a		g	ing F	<u>.</u>	-1.5 -1.5 -1.5	ng S	- 1.3		-1.3 -1.3	ics v ion s cy, a cy, a ir Ep ated conf	6 col orop ions:		a general population sample the previously
e 3. 7,42		-	I Us			I Usi				atist ress luen er fo stim: d=p	= 95% ed = 1 viati		well-described associations between cognitive
able		lode	lode	а-О,	, n D n	lode		יי בע מו	u D a	All st dep freq Cent S = es Cl = 9 Me	6Cl = %Me bbre		impairment (as measured by 3 cognitive tests)
IF 5	I .	\geq	2 ,	- v <u>v</u> v	๚๛๛๛	2	- c	йыл	m m m	က် မိမ်မိုက်	Ъ́ Я́ A	I .	

It is illegal to post this copp and both depression⁴⁻¹⁷ and altered functioning^{2,13,22} (as measured by 2 functioning scores). However, despite these findings, this large population-based study did not support the hypothesis that cognitive dysfunction could substantially explain the association between depression and functioning. This negative result could be explained by the population-based design of our study, which excluded severely depressed participants. However, we found similar results with a more stringent definition of depression. Thus, these analyses did not provide additional evidence for a mediation effect of cognitive impairment. To our knowledge, only 1 large-scale study³⁰ found support for this hypothesis by formally testing the mediation. However, that study relied on subjective cognitive complaints that are poorly correlated with objective cognitive functions such as those measured by standardized neuropsychological tests.⁴⁷ Some studies^{12,13} have reported an association between objectively assessed cognitive function and functional impairment in individuals with depression, thus providing preliminary evidence for a mediating role of cognitive function, but such mediation effect was not reported. Furthermore, since depression is associated with both cognitive and functional impairment, the association between objectively assessed cognitive function and functional impairment could have been confounded by depression itself.

Consistent with this alternative hypothesis, and contrasting with the lack of evidence for a mediating role of cognitive function, the mediation analyses suggested that depression could explain up to 44% of the relationship between cognitive deficits and altered functioning. Our results suggest that depression and cognitive impairment are strongly interrelated and both negatively impact functioning. However, depression without cognitive impairment may alter functioning to a greater extent than cognitive impairment without depression in a general population sample. Another plausible interpretation of this result is that depression might cause both cognitive and functional impairment. For instance, depression may result in both cognitive and functional impairment through altered motivation or

ghted PDF on any website, decreased self-efficacy. Depression may also simultaneously affect cognitive and functional impairment, but by different mechanisms. For instance, ruminative thoughts associated with depression may reduce cognitive resources during performance of externally oriented cognitive tasks,⁴⁸ whereas poor self-esteem and embarrassment may have detrimental impact on social functioning.³⁰ However, these findings are controversial. For example, in 117 remitted patients with major depressive disorder, no association was found between residual symptoms such as self-blaming, feeling worthless, or hopeless and impaired cognition.⁴⁹ As mediation and confounding are identical statistically, they can be distinguished only on conceptual grounds, even in longitudinal studies.⁵⁰ Therefore, strictly speaking, our results are also consistent with the hypothesis that depression could mediate, rather than confound, the association between cognitive and functional impairment. For instance, the DSST, TMT-B, and SVF outcomes might result from impaired cognitive control, which is also involved in poor emotion regulation and thus vulnerability to depression.⁴⁸ Although these 2 hypotheses (ie, confusion versus mediation) are not mutually exclusive, they both imply that the impact of depression on functional impairment is independent of cognitive impairment.

In adults aged 45 years or older from the general population, the association between depression at baseline and role limitations and social functioning at follow-up could not be explained by lower scores on cognitive tests. Although the management of cognitive impairment associated with depression is central to the treatment of depression, it may not be sufficient to improve the functioning beyond what is expected from the improvement of depression per se at the general population level. Further studies based on more ecological cognitive tests (eg, tests involving social cognition or integration of cognitive tests in social context), but also using objective measures of functioning (eg, absenteeism), are needed to further refine our understanding of the mechanisms explaining why depression is one of the most disabling conditions worldwide.⁵¹

Submitted: November 6, 2017; accepted May 14, 2018.

Published online: November 13, 2018.

Potential conflicts of interest: Pr Fossati received consulting and speaker honoraria from Janssen and Lundbeck and grant support from Servier. Pr Limosin received consulting and speaker honoraria from AstraZeneca, Euthérapie/Servier, Janssen, Lundbeck, Otsuka, and Roche. Pr Lemogne reports grants, personal fees, and non-financial support from Lundbeck; personal fees from Daiichi-Sankyo, Janssen, and Servier; and non-financial support from Otsuka, outside of this work. Drs Vulser, Wiernik, Hoertel, Melchior, Thibault, Olekhnovitch, Goldberg, and Zins have no conflict of interest to disclose.

Funding/support: The CONSTANCES cohort study was supported and funded by the Caisse nationale d'assurance maladie des travailleurs salariés (CNAMTS). The CONSTANCES cohort study is an "Infrastructure nationale en Biologie et Santé" and benefits from a grant from Agence nationale de la recherche (ANR-11-INBS-0002). CONSTANCES is also partly funded by Merck Sharpe & Dohme, AstraZeneca, and Lundbeck.

Role of the sponsor: Funding sources had no role in the conduct or publication of the study.

Acknowledgments: The authors thank the CNAMTS and the "Centres d'examens de santé" of the French Social Security, which collected a large part of the data, as well as the "Caisse nationale d'assurance vieillesse," ClinSearch, Asqualab, and Eurocell, which are in charge of the data quality control.

Supplementary material: Available at PSYCHIATRIST.COM.

REFERENCES

- World Health Organization. Depression. WHO website. http://www.who.int/mediacentre/ factsheets/fs369/en/ Accessed March 3, 2017.
- Broadhead WE, Blazer DG, George LK, et al. Depression, disability days, and days lost from work in a prospective epidemiologic survey. JAMA. 1990;264(19):2524–2528.

- Wells KB, Stewart A, Hays RD, et al. The functioning and well-being of depressed patients: results from the Medical Outcomes Study. JAMA. 1989;262(7):914–919.
- Adler DA, McLaughlin TJ, Rogers WH, et al. Job performance deficits due to depression. Am J Psychiatry. 2006;163(9):1569–1576.
- Weinstock LM, Keitner GI, Ryan CE, et al. Family functioning and mood disorders: a comparison between patients with major depressive disorder and bipolar I disorder. J Consult Clin Psychol. 2006;74(6):1192–1202.
- Reed C, Monz BU, Perahia DGS, et al. Quality of life outcomes among patients with depression after 6 months of starting treatment: results from FINDER. J Affect Disord. 2009;113(3):296–302.
- Kim JM, Chalem Y, di Nicola S, et al. A crosssectional study of functional disabilities and perceived cognitive dysfunction in patients with major depressive disorder in South Korea: the PERFORM-K study. *Psychiatry Res.* 2016;239:353–361.
- 8. Strine TW, Kroenke K, Dhingra S, et al. The

For reprints or permissions, contact permissions@psychiatrist.com.
© 2018 Copyright Physicians Postgraduate Press, Inc.
J Clin Psychiatry 79:6, November/December 2018
PSYCHIATRIST.COM
PSychiatry 79:6, November/December 2018

Vulser et al **It is illegal to post this copyrighted PDF on average state** associations between depression, healthrelated quality of life social support life

related quality of life, social support, life satisfaction, and disability in communitydwelling US adults. *J Nerv Ment Dis.* 2009;197(1):61–64.

- 9. Rizvi SJ, Cyriac A, Grima E, et al. Depression and employment status in primary and tertiary care settings. *Can J Psychiatry*. 2015;60(1):14–22.
- Soares CN, Kornstein SG, Thase ME, et al. Assessing the efficacy of desvenlafaxine for improving functioning and well-being outcome measures in patients with major depressive disorder: a pooled analysis of 9 double-blind, placebo-controlled, 8-week clinical trials. J Clin Psychiatry. 2009;70(10):1365–1371.
- Kennedy N, Foy K, Sherazi R, et al. Long-term social functioning after depression treated by psychiatrists: a review. *Bipolar Disord*. 2007;9(1–2):25–37.
- Evans VC, Iverson GL, Yatham LN, et al. The relationship between neurocognitive and psychosocial functioning in major depressive disorder: a systematic review. J Clin Psychiatry. 2014;75(12):1359–1370.
- Lam RW, Kennedy SH, McIntyre RS, et al. Cognitive dysfunction in major depressive disorder: effects on psychosocial functioning and implications for treatment. *Can J Psychiatry*. 2014;59(12):649–654.
- McDermott LM, Ebmeier KP. A meta-analysis of depression severity and cognitive function. J Affect Disord. 2009;119(1–3):1–8.
- Snyder HR. Major depressive disorder is associated with broad impairments on neuropsychological measures of executive function: a meta-analysis and review. *Psychol Bull*. 2013;139(1):81–132.
- Lee RSC, Hermens DF, Porter MA, et al. A metaanalysis of cognitive deficits in first-episode major depressive disorder. J Affect Disord. 2012;140(2):113–124.
- 17. Ahern E, Semkovska M. Cognitive functioning in the first-episode of major depressive disorder: a systematic review and metaanalysis. *Neuropsychology*. 2017;31(1):52–72.
- Mahableshwarkar AR, Zajecka J, Jacobson W, et al. A randomized, placebo-controlled, activereference, double-blind, flexible-dose study of the efficacy of vortioxetine on cognitive function in major depressive disorder. *Neuropsychopharmacology*. 2015;40(8):2025–2037.
- Rosenblat JD, Kakar R, McIntyre RS. The cognitive effects of antidepressants in major depressive disorder: a systematic review and meta-analysis of randomized clinical trials. Int J Neuropsychopharmacol. 2015;19(2):pyv0852.
- Bora E, Harrison BJ, Yücel M, et al. Cognitive impairment in euthymic major depressive disorder: a meta-analysis. *Psychol Med.* 2013;43(10):2017–2026.
- Rock PL, Roiser JP, Riedel WJ, et al. Cognitive impairment in depression: a systematic review and meta-analysis. *Psychol Med*. 2014;44(10):2029–2040.
- Jaeger J, Berns S, Uzelac S, et al. Neurocognitive deficits and disability in major depressive disorder. *Psychiatry Res.* 2006;145(1):39–48.

Natsmith SL, Longley WA, Scott EM, et al. Disability in major depression related to selfrated and objectively-measured cognitive deficits: a preliminary study. *BMC Psychiatry*. 2007;7(1):32.

- 24. Withall A, Harris LM, Cumming SR. The relationship between cognitive function and clinical and functional outcomes in major depressive disorder. *Psychol Med*. 2009;39(3):393–402.
- Baune BT, Miller R, McAfoose J, et al. The role of cognitive impairment in general functioning in major depression. *Psychiatry Res.* 2010;176(2–3):183–189.
- Rothschild AJ, Raskin J, Wang CN, et al. The relationship between change in apathy and changes in cognition and functional outcomes in currently non-depressed SSRI-treated patients with major depressive disorder. *Compr Psychiatry*. 2014;55(1):1–10.
- Bowie CR, Depp C, McGrath JA, et al. Prediction of real-world functional disability in chronic mental disorders: a comparison of schizophrenia and bipolar disorder. *Am J Psychiatry*. 2010;167(9):1116–1124.
- McIntyre RS, Cha DS, Soczynska JK, et al. Cognitive deficits and functional outcomes in major depressive disorder: determinants, substrates, and treatment interventions. *Depress Anxiety*. 2013;30(6):515–527.
- Woo YS, Rosenblat JD, Kakar R, et al. Cognitive deficits as a mediator of poor occupational function in remitted major depressive disorder patients. *Clin Psychopharmacol Neurosci*. 2016;14(1):1–16.
- Buist-Bouwman MA, Ormel J, de Graaf R, et al; ESEMeD/MHEDEA 2000 investigators. Mediators of the association between depression and role functioning. *Acta Psychiatr Scand*. 2008;118(6):451–458.
- Zins M, Bonenfant S, Carton M, et al. The CONSTANCES cohort: an open epidemiological laboratory. *BMC Public Health*. 2010;10(1):479.
- Zins M, Goldberg M; CONSTANCES team. The French CONSTANCES population-based cohort: design, inclusion and follow-up. *Eur J Epidemiol.* 2015;30(12):1317–1328.
- Radloff LS. The CES-D Scale: a self-report depression scale for research in the general population. *Appl Psychol Meas*. 1977;1(3):385–401.
- Morin AJS, Moullec G, Maïano C, et al. Psychometric properties of the Center for Epidemiologic Studies Depression Scale (CES-D) in French clinical and nonclinical adults. Rev Epidemiol Sante Publique. 2011;59(5):327–340.
- Wechsler D. WAIS-R, Wechsler Adult Intelligence Scale- Revised, Manual. New York, NY: Psychological Corporation; 1981.
- Gaudino EA, Geisler MW, Squires NK. Construct validity in the Trail Making Test: what makes Part B harder? J Clin Exp Neuropsychol. 1995;17(4):529–535.
- Borkowski JG, Benton AL, Spreen O. Word fluency and brain damage. *Neuropsychologia*. 1967;5(2):135–140.
- Ware J Jr, Kosinski M, Keller SDA. A 12-Item Short-Form Health Survey: construction of scales and preliminary tests of reliability and

- Ware JE; QualityMetric Incorporated, New England Medical Center Hospital; Health Assessment Lab. How to Score Version 2 of the SF-12 Health Survey (With a Supplement Documenting Version 1). Lincoln, RI: Health Assessment Lab; Boston, MA: QualityMetric Inc; 2005.
- Cheak-Zamora NC, Wyrwich KW, McBride TD. Reliability and validity of the SF-12v2 in the medical expenditure panel survey. *Qual Life Res.* 2009;18(6):727–735.
- Spijker J, Graaf R, Bijl RV, et al. Functional disability and depression in the general population: results from the Netherlands Mental Health Survey and Incidence Study (NEMESIS). Acta Psychiatr Scand. 2004;110(3):208–214.
- Imai K, Keele L, Tingley D. A general approach to causal mediation analysis. *Psychol Methods*. 2010;15(4):309–334.
- Steinberg BA, Bieliauskas LA, Smith GE, et al. Mayo's Older Americans Normative Studies: Age- and IQ-Adjusted Norms for the Trail-Making Test, the Stroop Test, and MAE Controlled Oral Word Association Test. Clin Neuropsychol. 2005;19(3–4):329–377.
- Larrabee GJ. Association between IQ and neuropsychological test performance: commentary on Tremont, Hoffman, Scott, and Adams (1998). *Clin Neuropsychol.* 2000;14(1):139–145.
- Ceci SJ. How much does schooling influence general intelligence and its cognitive components? a reassessment of the evidence. *Dev Psychol.* 1991;27(5):703–722.
- Guerra-Carrillo B, Katovich K, Bunge SA. Does higher education hone cognitive functioning and learning efficacy? findings from a large and diverse sample. *PLoS One*. 2017;12(8):e0182276.
- Burmester B, Leathem J, Merrick P. Subjective cognitive complaints and objective cognitive function in aging: a systematic review and meta-analysis of recent cross-sectional findings. *Neuropsychol Rev.* 2016;26(4):376–393.
- Nejad AB, Fossati P, Lemogne C. Self-referential processing, rumination, and cortical midline structures in major depression. Front Hum Neurosci. 2013;7:666.
- Pedrelli P, Baer L, Losifescu DV, et al. Relationship between residual symptoms of depression and self-reported cognitive impairment. CNS Spectr. 2010;15(1):46–51.
- MacKinnon DP, Krull JL, Lockwood CM. Equivalence of the mediation, confounding and suppression effect. *Prev Sci.* 2000;1(4):173–181.
- Baldessarini RJ, Forte A, Selle V, et al. Morbidity in depressive disorders. *Psychother Psychosom*. 2017;86(2):65–72.

Editor's Note: We encourage authors to submit papers for consideration as a part of our Early Career Psychiatrists section. Please contact Erika F. H. Saunders, MD, at esaunders@psychiatrist.com.

See supplementary material for this article at PSYCHIATRIST.COM.



THE OFFICIAL JOURNAL OF THE AMERICAN SOCIETY OF CLINICAL PSYCHOPHARMACOLOGY

Supplementary Material

- Article Title: Depression, Cognitive Functions, and Impaired Functioning in Middle-Aged Adults From the CONSTANCES Cohort
- Author(s): Hélène Vulser, MD, PhD; Emmanuel Wiernik, PhD; Nicolas Hoertel, MD; Maria Melchior, PhD; Mura Thibault, MD, PhD; Romain Olekhnovitch, PhD; Philippe Fossati, MD, PhD; Frédéric Limosin, MD, PhD; Marcel Goldberg, MD, PhD; Marie Zins, MD, PhD; and Cédric Lemogne, MD, PhD
- DOI Number: https://doi.org/10.4088/JCP.17m12003

List of Supplementary Material for the article

- 1. Appendix 1
- 2. Figure 1 Flow chart of the study population selection
- 3. <u>Table 1</u> Characteristics of participants according to depression status (N=7426)
- 4. <u>Table 2</u> Characteristics of lost to follow-up participants and comparisons with the study population

Disclaimer

This Supplementary Material has been provided by the author(s) as an enhancement to the published article. It has been approved by peer review; however, it has undergone neither editing nor formatting by in-house editorial staff. The material is presented in the manner supplied by the author.

© Copyright 2018 Physicians Postgraduate Press, Inc.

Appendix 1

Assessment of depressive symptoms

The CES-D consists of 20 items that are designed to measure self-reported depressive symptoms during the week prior to the test ¹ with adequate internal consistency (Cronbach's alpha: 0.88 in the current sample). The total score ranged from 0 (no depressive symptom) to 60. We used a cut-off score of 19 (CES-D score \geq 19 versus <19) to define depression status, according to the validation study of the French version of the CES-D (sensitivity/specificity for the diagnosis of major depression: 0.853/0.859)².

Assessment of cognitive functions

The DSST is a subtest of the Wechsler Adult Intelligence Scale-Revised, a timed paper- and pencil- task that measures psychomotor speed, sustained attention and logical reasoning ³. It consists of matching symbols with their corresponding numerical digit as fast as possible. The DSST score represents the number of correctly matched symbols in 120 seconds. TMT-B requires to draw lines sequentially connecting alternatively encircled numbers and letters (e.g., 1, A, 2, B, 3, C, etc.) distributed on a sheet of paper ⁴. The TMT-B score represents the amount of time required to complete the task. SVF requires participants to say as many words as possible from the "Animal" category in 60 seconds ⁵.

Assessment of functioning

The SF-12v2 is a widely used measure of health-related quality of life, with adequate reliability and validity ⁶⁻⁸. It measures eight health aspects, namely general health, physical functioning, role limitations due to physical health problems, bodily pain, vitality, social functioning, role limitations due to emotional problems, and mental health. Mental health and

vitality subscales were not taken into account because of their obvious overlap with CES-D items ("Did you have a lot of energy?", "Have you felt downhearted and depressed?"). Because we were interested in functional impairment associated with depression specifically, we a priori decided to use two subscales as primary outcomes ⁹: role limitations due to emotional problems and social functioning. These three items were rated from 1 ("All of the time") to 5 ("None of the time"), leading to a score from 2 to 10 for role limitations and from 1 to 5 for social functioning ⁹. For both scales, a higher score corresponds with a better functioning. Role limitations due to emotional problems, henceforth referred to as "role limitations", was assessed with two items: "During the past 4 weeks, how much of the time have you had any of the following problems with your work or other regular daily activities as a result of any emotional problems (such as feeling depressed or anxious)? 1) Accomplished less than you would like, 2) Did work or other activities less carefully than usual". Social functioning was assessed with one item: "During the past 4 weeks, how much of the time has your physical health or emotional problems interfered with your social activities (like visiting friends, relatives, etc.)?"). These three items were rated from 1 ("All of the time") to 5 ("None of the time"), leading to a score from 2 to 10 for role limitations and from 1 to 5 for social functioning⁹. For both scales, a higher score corresponds with a better functioning.

References

1. Radloff LS. The CES-D Scale: A self-report depression scale for research in the general population. *Applied Psychological Measurement*. 1977;1(3):385-340.

2. Morin AJS, Moullec G, Maïano C, et al. Psychometric properties of the Center for Epidemiologic Studies Depression Scale (CES-D) in French clinical and nonclinical adults. *Rev Epidemiol Sante Publique*. 2011;59(5):327-340.

3. Wechsler D. *WAIS-R, Wechsler Adult Intelligence Scale- Revised, Manual.* New York, NY:Psychological Corporation; 1981.

4. Gaudino EA, Geisler MW, Squires NK. Construct validity in the Trail Making Test: what makes Part B harder? *J Clin Exp Neuropsychol*. 1995;17(4):529-535.

5. Borkowski JG, Benton AL, Spreen O. Word fluency and brain damage. *Neuropsychologia*. 1967;5(2):135-140.

6. Ware J, Kosinski M, Keller SD. A 12-Item Short-Form Health Survey: construction of scales and preliminary tests of reliability and validity. *Med Care*. 1996;34(3):220-233.

7. Ware J, QualityMetric Incorporated, New England Medical Center Hospital, Health Assessment Lab. *How to score version 2 of the SF-12 health survey (with a supplement documenting version 1)*. Lincoln, R.I.; Boston, Mass.: QualityMetric Inc. ; Health Assessment Lab; 2005.

8. Cheak-Zamora NC, Wyrwich KW, McBride TD. Reliability and validity of the SF-12v2 in the medical expenditure panel survey. Qual Life Res Int J Qual Life Asp Treat Care Rehabil. 2009;18(6):727-735.

9. Spijker J, Graaf R, Bijl RV, et al. Functional disability and depression in the general population. Results from the Netherlands Mental Health Survey and Incidence Study (NEMESIS). *Acta Psychiatr Scand*. 2004;110(3):208-214.

Supplementary Figure 1. Flow chart of the study population selection



Supplementary Table 1: Characteristics of participants according to depression status (N=7426)

	(CES-D) score	(CES-	-D score		
	<1	9)	≥	:19)		
	N=6	443	N=	=983		
Continuous variables	mean	sd	mean	sd	t	р
Age	57.91	7.24	57.03	6.88	-3.70	< 0.001
N of days between inclusion and FU	497.47	158.52	497.47	148.74	0.03	>0.99
CES-D score	7.34	4.91	26.56	6.95	83.53	< 0.001
DSST score	67.55	14.99	65.48	15.62	-3.89	< 0.001
TMT-B score	66.79	31.39	70.78	32.69	3.59	< 0.001
SVF score	23.92	5.78	23.18	5.75	-3.78	< 0.001
Role limitations	8.73	1.59	6.70	1.93	-31.56	< 0.001
Social functioning	4.26	0.83	3.29	0.95	-30.37	< 0.001
Discrete variables	Ν	%	Ν	%	χ²	р
Sex					115.17	< 0.001
Men	3238	50.26	313	31.84		
Women	3205	49.74	670	68.16		
Date of inclusion					1.11	0.29
2012	1046	16.23	146	14.85		
2013	5397	83.77	837	85.15		
Education level					30.77	< 0.001
No diploma	103	1.60	27	2.75		

Lower secondary education	513	7.96	99	10.07		
Professional education	1207	18.73	203	20.65		
Upper secondary education	1104	17.13	197	20.04		
Bachelor	1477	22.92	211	21.46		
Fourth year university level	658	10.21	88	8.95		
Master degree or higher	1174	18.22	134	13.63		
Other	207	3.21	24	2.44		
Alcohol intake					53.38	< 0.001
Never	189	2.93	45	4.58		
≤ 1 glass/month	714	11.08	179	18.21		
2-3glasses/month	1224	19.00	187	19.02		
≥1glass/week	4316	66.99	572	58.19		
Life cannabis use					2.10	0.35
Yes	1140	17.69	192	19.53		
No	5263	81.69	786	79.96		
No intent to answer	40	0.62	5	0.51		

CES-D: Center for Epidemiologic Studies Depression Scale; DSST: Total score for Digit Symbol Substitution Test; TMT-B: Total time for Trail Making Test B, SVF: Semantic Verbal Fluency; FU: follow-up; sd: standard deviation; χ^2 : chi-square value; t: t value.

Supplementary Table 2: Characteristics of lost to follow-up participants and

comparisons with the study population

	Include stu	ed in the 1dy	Lost to	follow-up		
	N=7	7426	N=	1474		
Continuous variables	mean	sd	mean	sd	t	р
Age	57.80	7.20	57.19	7.23	-2.93	0.003
CES-D score	9.88	8.35	12.28	10.05	8.60	< 0.001
DSST score	67.27	15.09	62.88	15.31	-10.08	< 0.001
TMT-B score	67.32	31.60	75.38	38.20	7.60	< 0.001
SVF score	23.82	5.78	22.51	5.76	-7.99	< 0.001
Discrete variables	Ν	%	Ν	%	χ^2	р
Sex						
Men	3551	47.82	743	50.41	3.20	0.07
Women	3875	52.18	731	49.59		
Date of inclusion						
2012	1192	16.05	251	17.03	0.79	0.37
2013	6234	83.95	1223	82.97		
Education level						
No diploma	130	1.75	81	5.50	143.23	< 0.001
Lower secondary education	612	8.24	195	13.23		
Professional education	1410	18.99	317	21.51		
Upper secondary education	1301	17.52	229	15.54		

Bachelor	1688	22.73	290	19.67		
Fourth year university level	746	10.05	102	6.92		
Master degree or higher	1308	17.61	204	13.84		
Other	231	3.11	56	3.80		
Alcohol intake					0.90	0.82
Never	234	3.15	44	2.99		
≤ 1 glass/month	893	12.03	179	12.14		
2-3glasses/month	1411	19.00	266	18.05		
≥1glass/week	4888	65.82	985	66.82		
Life cannabis use					4.61	0.10
Yes	1332	17.94	274	18.59		
No	6049	81.46	1184	80.33		
No intent to answer	45	0.61	16	1.09		

CES-D: Center for Epidemiologic Studies Depression Scale; DSST: Total score for Digit Symbol Substitution Test; TMT-B: Total time for Trail Making Test B, SVF: Semantic Verbal Fluency; FU: follow-up; sd: standard deviation; χ^2 : chi-square value; t: t value.