

ORIGINAL RESEARCH

Impact of Cognitive Symptoms on Health-Related Quality of Life and Work Productivity in Chinese Patients with Major Depressive Disorder: Results from the PROACT Study

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Patients and Methods: This was an epidemiological, non-interventional, prospective cohort study in adult outpatients with moderate-to-severe MDD initiating antidepressant monotherapy (first or second line). Crude and adjusted analyses of covariance were performed to assess the association of perceived cognitive symptoms (20-item Perceived Deficits Questionnaire-Depression [PDQ-D] total score) or observed cognitive performance (Digit Symbol Substitution Test [DSST] score) with HRQoL (EuroQoL 5-Dimensions Questionnaire index) and work productivity (Work Productivity and Activity Impairment [WPAI] or Sheehan Disability Scale [SDS] absenteeism and presenteeism scores). Adjusted analyses included depression severity, age, sex, residential area (urban/rural), and educational level.

Results: Of 1008 patients enrolled in the PROACT study, 986 were included in this analysis. Severity of perceived cognitive symptoms (ie, higher PDQ-D total score) was significantly associated with worse HRQoL (P<0.001) and higher levels of absenteeism (P=0.020 for the WPAI and P=0.002 for the SDS) and presenteeism (P<0.001 for both scales). The association of perceived cognitive symptoms with HRQoL and presenteeism was independent of depression severity. The association between observed cognitive performance (DSST score) and HRQoL was less robust. No association was seen between observed cognitive performance and levels of absenteeism or presenteeism assessed by either scale.

Conclusion: Results of this real-world study illustrate the impact of cognitive symptoms on HRQoL and work productivity in Chinese patients with MDD, and highlight the importance of assessing and targeting cognitive symptoms in order to improve functional outcomes when treating patients with MDD.

Keywords: major depressive disorder, cognitive symptoms, health-related quality of life, work productivity, presenteeism, absenteeism

Introduction

Major depressive disorder (MDD) is a prevalent and disabling condition, accounting for 8.2% of years lived with disability worldwide. In China, weighted lifetime prevalence rates for depressive disorders and MDD were 6.8% and 3.4%, respectively, in 2013. MDD is the second leading cause of years lived with disability in

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Tel +45 3083 2880 Email leem@lundbeck.com China;³ however, available data suggest that this condition is underdiagnosed and undertreated in Chinese populations.^{4–6} In a recent cross-sectional study, only 8.3% of respondents meeting diagnostic criteria for MDD had been clinically diagnosed, and of these, only 51.5% were receiving antidepressant medication.⁶ Of note, Chinese adults with MDD reported significantly worse health-related quality of life (HRQoL) and greater productivity loss than those without depression.⁶

It is well documented that MDD imposes significant impairments in HRQoL and work productivity. ^{7,8} MDD is associated both with having to take time off work (absenteeism) and with reduced productivity when at work (presenteeism), ⁹ and its impact on the ability to work and work productivity appears greater than that of other common mental and physical health disorders. ^{10–12} Indeed, reduced work productivity accounts for the majority of the financial costs attributed to MDD. ^{10,13}

Cognitive symptoms are common in patients with MDD, with deficits observed across multiple domains (eg, executive function, memory, and attention). 14-16 Such symptoms have been shown to negatively impact both clinical and functional outcomes in patients with MDD, including HRQoL and work productivity. 17-28 However, data are lacking concerning the impact of cognitive symptoms on HRQoL and work productivity in Chinese patients with MDD. The Prospective Research Observation to Assess Cognition in Treated patients with MDD (PROACT) study was undertaken to evaluate the prevalence and course of cognitive symptoms in Chinese patients initiating new antidepressant treatment in a real-world setting.²⁹ We present results of a post hoc analysis undertaken to further explore the association of cognitive symptoms with HRQoL and work productivity in the PROACT study population at the time of treatment initiation.

Methods

Study Design

PROACT was an epidemiological, non-interventional, prospective, cohort study conducted at 15 sites in China between March 2016 and July 2017. The study design has been reported in detail previously. Eligible participants were outpatients aged 18–65 years with a diagnosis of MDD (International Statistical Classification of Diseases and Related Health Problems [10th revision] criteria) and a total 17-item Hamilton Depression Rating Scale (HAM-D₁₇) score \geq 17 (indicating moderate-to-

severe depression), and were initiating antidepressant monotherapy (first- or second-line therapy). Key exclusion criteria included: comorbidities, including schizophrenia or other psychotic disorders, bipolar disorder, and alcohol or substance dependence; use of combination therapy (ie, receiving more than one antidepressant, adjunctive antipsychotics, or mood stabilizers); acute suicidality; and pregnancy/breastfeeding.

Ethical Approval and Consent to Participate

The study was conducted in accordance with the ethical principles described in the Declaration of Helsinki and the approved protocol, as well as all other applicable laws and regulations. Ethical approval was obtained from the necessary committees for each study site and all patients provided written informed consent for participation.

Study Assessments

Depression severity was evaluated by clinicians using the HAM-D₁₇ and by patients using the 9-item Patient Health Questionnaire (PHQ-9).^{30,31} The HAM-D₁₇ assesses the severity of psychological and somatic depressive symptoms over the past week through a semi-structured interview.³⁰ Each item is rated on a scale from 0 (absent) to 2 or 4 (maximum severity); individual item scores are then combined to yield a total score ranging from 0 to 52. The PHQ-9 assesses the severity of depressive symptoms over the previous 2 weeks on a scale from 0 to 27.³¹ On both scales, higher scores indicate more severe depression.

Perceived cognitive symptoms were assessed by patients using the 20-item Perceived Deficits Questionnaire-Depression (PDQ-D). The PDQ-D assesses self-perceived difficulties over the previous 7 days across four domains of cognitive function: attention/concentration, prospective memory, retrospective memory, and planning/organization. Each of the 20 questions (5 per domain) is rated from 0 to 4, yielding a possible total score of 0 to 80; higher scores indicate greater perceived cognitive symptoms. The Chinese version of the PDQ-D has been shown to be psychometrically valid for the evaluation of subjective cognitive symptoms in patients with MDD. 33

Cognitive performance was evaluated using the Digit Symbol Substitution Test (DSST), a neuropsychological coding test that requires the patient to substitute simple symbols for digits.³⁵ The number of correct symbols substituted for digits during a 90-second period provides

Dovepress Wang et al

a score ranging from 0 to 133, with higher scores reflecting better cognitive performance. The DSST assesses cognitive performance across a number of domains, including those known to be impaired in patients with MDD, such as executive function, processing speed and attention. The DSST has shown sensitivity to change in MDD populations, ^{17,36,37} and age-related norms have been defined for US adults. ³⁵

HRQoL was assessed with the widely used self-reported EuroQoL 5-Dimensions Questionnaire (EQ-5D). This instrument encompasses five health dimensions (mobility, self-care, usual activities, pain/discomfort, and anxiety/depression). Responses were recorded using three levels of severity (no problem, some problems, or extreme problems). The EQ-5D assesses HRQoL at the time of completion; there is no recall period. The EQ-5D utility index score was computed, based on coefficients derived from regression results in a Chinese population. An EQ-5D utility index score of 1 indicates perfect health, 0 indicates a state equivalent to being dead, and <0 indicates a state considered worse than being dead.

Work productivity was assessed by the Work Productivity and Activity Impairment - Specific Health Problems Questionnaire (WPAI-SHP)⁴⁰ and the Sheehan Disability Scale (SDS). 41,42 The WPAI questionnaire is a validated self-reported instrument for the assessment of work-related impairment due to a health problem such as MDD over the past 7 days. 40 The WPAI measures work hours missed due to the health problem, work hours missed due to other reasons, hours worked, the degree to which the health problem affected productivity while working, and the degree to which the health problem affected non-work daily activities. Based on validated algorithms, the WPAI yields four subscores: absenteeism (percent work time missed), presenteeism (percent time impaired while working), work productivity loss (percent overall work impairment), and percent overall activity impairment. Higher scores indicate greater impairment and lower work productivity.

The SDS is a brief self-administered tool that evaluates functional impairment during the previous 7 days across three domains: work/school, social life/leisure activities, and family life/home duties. The severity of impairment in each domain is rated by the patient on a scale from 0 (no impairment) to 10 (extreme impairment). The SDS also assesses the number of work/study days lost (absenteeism) and the number of underproductive work/study days (presenteeism) over the past week.

Statistical Analysis

The population for this analysis comprised all enrolled patients who complied with the study inclusion and exclusion criteria. Crude and adjusted analyses of covariance were performed to assess the association of perceived cognitive symptoms (PDQ-D total score) or observed cognitive performance (DSST total score) with HROoL (EO-5D utility index) and work productivity (WPAI or SDS absenteeism and presenteeism scores). Patients were categorized into groups based on quartiles of PDQ-D total score at study entry: (a) 0-20; (b) 21-32; (c) 33-44; and (d) 45-80. For DSST, patients were categorized into four mutually exclusive categories based on DSST total score and age at baseline: (a) within norm; (b) 1/3-2/3 standard deviation (SD) below norm; (c) ²/₃-1 SD below norm; and (d) ≥1 SD below norm. As DSST norms were developed using data from US adults,³⁵ patients were also categorized into groups based on DSST total score quartiles at baseline: (a) 0-37; (b) 38-51; (c) 52-62; and (d) 63-133. For this analysis, lost work days and underproductive work days were disregarded if patients indicated that they had not worked or studied at all during the past week for reasons unrelated to MDD. In the adjusted analyses, depression severity (PHQ-9 or HAM-D₁₇ score) was adjusted as a continuous variable, together with age, sex, residential area (urban/rural), and educational level, as these were identified as possible confounders based on clinical experience and in line with the primary analysis of data from this study.²⁹ All analyses were performed using R version 3.5.1.43

Results

Of 1008 patients enrolled in the PROACT study, 986 were included in this analysis (97.8%). Reasons for exclusion from the study, and hence this analysis population, were: not initiating antidepressant monotherapy at baseline (n=14); use of antipsychotics or mood stabilizers at baseline (n=7); pregnant or \leq 6 months postpartum (n=12); and unable to read and/or understand the information sheet and informed consent form (n=2). Patients could have had more than one reason for exclusion.

Patient demographics and clinical characteristics are shown in Table 1. Mean (SD) age was 35.9 (12.0) years, and patients were mostly female (68.5%), lived in urban areas (82.2%), and had received formal education up to at least high school level (76.7%). Just over half of the study participants were employed (57.0%). Patients were

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Table I Baseline Patient Demographics and Clinical Characteristics

Characteristic	Patients with MDD (n=986)
Age (years), mean (SD)	35.9 (12.0)
Female sex, % (n)	68.5 (675)
Region in China, % (n)	
North	56.9 (561)
South	6.8 (67)
East	17.5 (173)
West	18.8 (185)
Residential area, % (n)	
Urban	82.2 (810)
Rural	17.8 (176)
Employed, % (n)	57.0 (562)
Educational level, % (n)	
Middle school or lower	23.3 (230)
High school or junior college	31.7 (313)
University or postgraduate	44.9 (443)
Clinical history, % (n)	
Previous episode	39.2 (387)
Switching antidepressant at baseline	20.6 (203)
Baseline assessment scores,	
mean (SD)	
HAM-D ₁₇ total score	23.2 (4.4)
PHQ-9 total score	16.7 (5.8)
DSST	50.7 (16.8)
PDQ-D total score	33.6 (16.3)
EQ-5D utility index	0.74 (0.14)
SDS absenteeism (work days lost)	2.0 (2.5)
SDS presenteeism (work days	3.7 (2.5)
underproductive)	
WPAI absenteeism (%)	26.3 (35.7)
WPAI presenteeism (%)	54.5 (26.8)
PDQ-D quartile, % (n)	
0–20	24.0 (231)
21–32	25.7 (247)
33–44	24.9 (239)
45–80	25.4 (244)
Missing (n)	25
DSST subgroup, % (n)	45.4.(445)
Within norm	45.4 (445)
1/3-2/3 SD below norm	13.3 (130)
² / ₃ −I SD below norm	10.9 (107)
≥I SD below norm	30.5 (299)
Missing (n)	5

(Continued)

Table I (Continued).

Characteristic	Patients with MDD (n=986)
DSST quartile, % (n)	
0–37	23.8 (233)
38–51	25.0 (245)
52–62	25.9 (254)
63–133	25.4 (249)
Missing (n)	5

Abbreviations: DSST, Digit Symbol Substitution Test; EQ-5D, EuroQol 5-Dimensions Questionnaire; HAM-D₁₇, 17-item Hamilton Depression Rating Scale; MDD, major depressive disorder; PDQ-D, 20-item Perceived Deficits Questionnaire-Depression; PHQ-9, 9-item Patient Health Questionnaire; SD, standard deviation; SDS, Sheehan Disability Scale; WPAI, Work Productivity and Activity Impairment.

predominantly initiating first-line antidepressant treatment (79.4%). Mean (SD) HAM-D₁₇ total score was 23.2 (4.4) and mean (SD) PHQ-9 total score was 16.7 (5.8). Over half of all patients (54.6%) had DSST score $\geq \frac{1}{3}$ SD below norm. Mean (SD) DSST score was 50.7 (16.8) and mean (SD) PDQ-D total score was 33.6 (16.3). With regard to impairment in HRQoL and work productivity, mean (SD) EQ-5D utility index score at study entry was 0.74 (0.14), mean (SD) WPAI absenteeism and presenteeism scores were 26.3% (35.7%) and 54.5% (26.8%), respectively, and the mean (SD) number of working days during the past week that were lost or unproductive, as assessed using the SDS, were 2.0 (2.5) and 3.7 (2.5), respectively.

Results of the analyses performed to assess the association of perceived cognitive symptoms (PDQ-D score) with HRQoL (EQ-5D utility index) and work productivity (WPAI or SDS absenteeism and presenteeism scores) are shown in Table 2. Higher PDQ-D total score (ie, greater severity of perceived cognitive symptoms) was found to be associated with both lower HRQoL (P<0.001) and higher levels of absenteeism (P=0.020 for the WPAI and P=0.002 for the SDS) and presenteeism (P < 0.001 for both scales). The association of PDQ-D total score with HRQoL and presenteeism was shown to be independent of depression severity, as it remained statistically significant when the analysis was adjusted for PHQ-9 or HAM-D₁₇ scores. However, the association between PDQ-D total score and absenteeism did not persist when adjusted for PHQ-9 or HAM-D₁₇ scores, suggesting that this was largely driven by depression severity.

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Table 2 Association of PDQ-D Total Scores with HRQoL and Work Productivity Scores in Crude and Adjusted^a Analyses

Assessment	Model	PDQ-D Quartiles at Baseline	t Baseline			df	Error df	ш	Global P-value	Partial η ²
		0-20	21–32	33–44	45–80					
HRQoL	Crude PHQ-9 adjusted HAM-D ₁₇ adjusted	0.80 (0.78, 0.81) 0.75 (0.74, 0.80) 0.78 (0.77, 0.80)	0.74 (0.73, 0.76) 0.73 (0.72, 0.75) 0.73 (0.72, 0.75)	0.73 (0.72, 0.75) 0.73 (0.72, 0.75) 0.74 (0.72, 0.75)	0.67 (0.66, 0.69) 0.71 (0.69, 0.72) 0.68 (0.66, 0.70)	ж ж ж	948 938 942	33.78 4.42 25.89	<0.001 0.004 <0.001	0.097 0.014 0.076
SDS absenteeism	Crude PHQ-9 adjusted HAM-D ₁₇ adjusted	1.52 (1.15, 1.89) 2.01 (1.55, 2.46) 1.83 (1.41, 2.25)	1.85 (1.49, 2.21) 2.10 (1.70, 2.50) 2.13 (1.73, 2.53)	2.33 (1.96, 2.70) 2.51 (2.09, 2.93) 2.47 (2.05, 2.89)	2.40 (2.05, 2.76) 2.32 (1.88, 2.76) 2.46 (2.05, 2.8)		724 714 718	5.03 1.21 2.45	0.002 0.307 0.063	0.020 0.005 0.010
SDS presenteeism	Crude PHQ-9 adjusted HAM-D ₁₇ adjusted	2.60 (2.25, 2.96) 2.87 (2.45, 3.29) 2.44 (2.04, 2.84)	3.55 (3.21, 3.89) 3.44 (3.06, 3.82) 3.36 (2.97, 3.74)	4.06 (3.70, 4.41) 3.79 (3.40, 4.19) 3.82 (3.42, 4.23)	4.37 (4.03, 4.70) 3.75 (3.34, 4.16) 4.13 (3.74, 4.52)	m m m	718 708 712	19.04 4.38 15.69	<0.001 0.005 <0.001	0.074 0.018 0.062
WPAI absenteeism	Crude PHQ-9 adjusted HAM-D ₁₇ adjusted	22.81 (16.27, 29.36) 29.12 (21.06, 37.19) 26.53 (19.05, 34.01)	20.12 (13.85, 39.76) 21.86 (14.76, 28.95) 22.34 (15.30, 29.38)	30.36 (23.95, 36.76) 32.77 (25.20, 40.34) 30.66 (23.15, 38.18)	32.05 (26.06, 38.04) 30.15 (22.65, 37.65) 32.46 (25.38, 39.53)	8 8 8	489 480 483	3.31 2.09 1.93	0.020 0.101 0.124	0.020 0.013 0.012
WPAI presenteeism	Crude PHQ-9 adjusted HAM-D ₁₇ adjusted	40.20 (35.46, 44.94) 46.80 (41.35, 52.25) 39.65 (34.30, 45.00)	51.26 (46.72, 55.80) 51.75 (46.94, 56.56) 50.58 (45.51, 55.65)	53.37 (48.60, 58.13) 51.60 (46.34, 56.86) 50.86 (45.32, 56.41)	70.66 (66.3, 74.99) 62.11 (58.89, 67.33) 67.55 (62.29, 72.81)		432 423 426	30.35 6.65 24.43	<0.001 <0.001 <0.001	0.174 0.045 0.147

Notes: Data are presented as means with 95% confidence intervals. F statistic, global P-value and partial η² were obtained via Type III sum of squares. ^aAge, sex, residential area and educational level also included as adjustment variables in the adjusted analyses.

Abbreviations: df, degrees of freedom; HAM-D₁₇, 17-item Hamilton Depression Rating Scale; HRQoL, health-related quality of life; PDQ-D, 20-item Perceived Deficits Questionnaire-Depression; PHQ-9, 9-item Patient Health Questionnaire; SDS, Sheehan Disability Scale; WPAI, Work Productivity and Activity Impairment.

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Table 3 Association of DSST Scores (by Subgroup) with HRQoL and Work Productivity Scores in Crude and Adjusted^a Analyses

HROol.		Door Jangroup at Basenne						_	Global	Partial
		Within Norm	½ to ½ SD Below Norm	% to I SD Below Norm	I SD or More Below Norm		p		P-value	Ju,
	Crude PHQ-9 adjusted HAM-D ₁₇ adjusted	0.75 (0.74, 0.77) 0.74 (0.73, 0.76) 0.75 (0.73, 0.77)	0.73 (0.70, 0.75) 0.72 (0.70, 0.74) 0.73 (0.70, 0.75)	0.74 (0.72, 0.77) 0.74 (0.71, 0.76) 0.73 (0.71, 0.76)	0.71 (0.70, 0.73) 0.72 (0.71, 0.74) 0.72 (0.71, 0.74)	m m m	965 954 959	5.42	0.001 0.162 0.090	0.017 0.005 0.007
SDS absenteeism Crude PHQ-9 HAM-D	Crude PHQ-9 adjusted HAM-D ₁₇ adjusted	1.97 (1.71, 2.23) 2.22 (1.86, 2.58) 2.22 (1.86, 2.57)	1.89 (1.40, 2.38) 2.08 (1.56, 2.61) 2.10 (1.58, 2.63)	1.99 (1.39, 2.58) 2.21 (1.59, 2.83) 2.25 (1.63, 2.87)	2.20 (1.86, 2.54) 2.34 (1.96, 2.72) 2.31 (1.94, 2.69)		737 726 731	0.49 0.24 0.16	0.687 0.870 0.921	0.002 0.001 0.0007
SDS presenteeism Crude PHQ-9 HAM-D	Crude PHQ-9 adjusted HAM-D ₁₇ adjusted	3.75 (3.50, 4.01) 3.55 (3.21, 3.89) 3.45 (3.10, 3.80)	3.60 (3.11, 4.08) 3.45 (2.96, 3.95) 3.43 (2.92, 3.94)	3.29 (2.71, 3.87) 3.18 (2.59, 3.76) 3.12 (2.51, 3.72)	3.65 (3.32, 3.98) 3.54 (3.19, 3.89) 3.59 (3.23, 3.95)		732 721 726	0.73 0.52 0.65	0.533 0.671 0.583	0.003 0.002 0.003
WPAI absenteeism Crude PHQ-9 HAM-D	Crude PHQ-9 adjusted HAM-D ₁₇ adjusted	26.14 (21.64, 30.64) 29.86 (23.41, 36.31) 29.30 (22.96, 35.64)	24.39 (15.79, 32.99) 25.20 (16.10, 34.30) 26.98 (17.95, 36.00)	26.36 (16.60, 36.12) 28.36 (17.96, 38.76) 29.04 (18.74, 39.35)	27.76 (21.77, 33.75) 27.38 (20.62, 34.15) 26.24 (19.55, 32.93)	3 3 3	498 489 492	0.14 0.32 0.22	0.936 0.814 0.880	0.0008 0.002 0.001
WPAI presenteeism Crude PHQ-9 HQ-9	Crude PHQ-9 adjusted HAM-D ₁₇ adjusted	54.93 (51.37, 58.49) 53.82 (49.20, 58.44) 50.74 (45.68, 55.80)	48.95 (41.97, 55.92) 47.80 (41.27, 54.32) 48.14 (40.91, 55.38)	57.92 (50.32, 65.52) 56.60 (49.43, 63.76) 55.40 (47.48, 63.32)	55.17 (50.36, 59.97) 53.90 (49.14, 58.66) 53.66 (48.41, 58.90)	е е е	440 431 434	1.12	0.341 0.239 0.409	0.008 0.01 0.007

Notes: Data are presented as means with 95% confidence intervals. F statistic, global P-value and partial n? were obtained via Type III sum of squares. Age, sex, residential area and educational level also included as adjustment variables in the adjusted analyses.

Abbreviations: df, degrees of freedom; DSST, Digit Symbol Substitution Test; HAM-D₁₇, 17-item Hamilton Depression Rating Scale; HRQoL, health-related quality of life; PHQ-9, 9-item Patient Health Questionnaire; SD, standard deviation; SDS, Sheehan Disability Scale; WPAI, Work Productivity and Activity Impairment.

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Table 4 Association of DSST Scores (by Quartile) with HRQoL and Work Productivity Scores in Crude and Adjusted^a Analyses

Assessment	Model	DSST Quartile at Baseline	kaseline			₽	Error df	ш	Global P-value	Partial η^2
		0–37	38–51	52–62	63-133					
HRQoL	Crude PHQ-9 adjusted HAM-D ₁₇ adjusted	0.71 (0.69, 0.73) 0.72 (0.70, 0.74) 0.72 (0.70, 0.74)	0.73 (0.72, 0.75) 0.73 (0.71, 0.75) 0.73 (0.71, 0.75)	0.75 (0.73, 0.77) 0.74 (0.73, 0.76) 0.75 (0.73, 0.77)	0.75 (0.73, 0.77) 0.74 (0.72, 0.76) 0.74 (0.72, 0.77)	3 3 3	965 954 959	4.82 1.40 1.43	0.002 0.242 0.233	0.015 0.004 0.004
SDS absenteeism	Crude PHQ-9 adjusted HAM-D ₁₇ adjusted	2.05 (1.62, 2.47) 2.27 (1.79, 2.75) 2.22 (1.74, 2.70)	2.05 (1.70, 2.40) 2.28 (1.88, 2.67) 2.29 (1.90, 2.68)	2.14 (1.80, 2.49) 2.32 (1.91, 2.73) 2.31 (1.90, 2.72)	1.88 (1.54, 2.22) 2.08 (1.63, 2.54) 2.11 (1.66, 2.56)		737 726 731	0.38 0.32 0.25	0.767 0.811 0.860	0.002 0.001 0.001
SDS presenteeism	Crude PHQ-9 adjusted HAM-D ₁₇ adjusted	3.55 (3.14, 3.97) 3.59 (3.14, 4.04) 3.63 (3.16, 4.10)	3.37 (3.03, 3.71) 3.26 (2.89, 3.64) 3.26 (2.87, 3.64)	3.72 (3.38, 4.06) 3.52 (3.14, 3.91) 3.46 (3.06, 3.87)	3.95 (3.61, 4.28) 3.69 (3.26, 4.11) 3.59 (3.15, 4.03)		732 721 726	2.01 1.10 0.87	0.112 0.348 0.456	0.008 0.005 0.004
WPAI absenteeism	Crude PHQ-9 adjusted HAM-D ₁₇ adjusted	27.81 (20.43, 35.18) 26.58 (18.34, 34.82) 24.27 (16.05, 32.48)	27.73 (21.61, 33.86) 28.45 (21.42, 35.48) 29.35 (22.42, 36.27)	25.55 (19.71, 31.40) 28.13 (21.00, 35.27) 27.97 (20.94, 35.01)	24.95 (18.88, 31.03) 29.28 (21.20, 37.37) 29.86 (21.87, 37.85)	3 3 3	498 489 492	0.21 0.08 0.43	0.891 0.969 0.731	0.001 0.0005 0.003
WPAI presenteeism	Crude PHQ-9 adjusted HAM-D ₁₇ adjusted	53.82 (47.76, 59.88) 54.27 (48.32, 60.21) 53.56 (46.96, 60.16)	53.68 (48.79, 58.56) 53.05 (48.09, 58.01) 52.41 (46.97, 57.85)	54.48 (49.76, 59.20) 52.75 (47.56, 57.95) 50.76 (45.05, 56.47)	55.87 (51.17, 60.58) 52.68 (47.12, 58.24) 50.64 (44.51, 56.77)		440 431 434	0.16	0.922 0.979 0.895	0.001

Notes: Data are presented as means with 95% confidence intervals. F statistic, global P-value and partial η² were obtained via Type III sum of squares. ^aAge, sex, residential area and educational level also included as adjustment variables

in the adjusted analyses.

Abbreviations: df, degrees of freedom; DSST, Digit Symbol Substitution Test; HAM-D₁,, 17-item Hamilton Depression Rating Scale; HRQoL, health-related quality of life; PHQ-9, 9-item Patient Health Questionnaire; SDS, Sheehan Disability Scale; WPAI, Work Productivity and Activity Impairment.

Wang et al Dovepress

Results of the analyses performed to assess the association of cognitive performance (DSST score) with HRQoL (EQ-5D utility index) and work productivity (WPAI or SDS absenteeism and presenteeism scores) are shown in Tables 3 and 4. Lower DSST total score (ie, lower cognitive performance) was associated with lower HRQoL (*P*=0.001 by DSST subgroup and *P*=0.002 by DSST quartile). However, the association between DSST total score and HRQoL did not persist when adjusted for PHQ-9 or HAM-D₁₇ scores. No association was seen between DSST total score and levels of absenteeism or presenteeism assessed by either scale.

Discussion

To our knowledge, this is the first study to describe the impact of cognitive symptoms on HRQoL and work productivity in Chinese patients with MDD. Patients were required to have moderate-to-severe depression to be eligible for study entry and most were initiating first-line antidepressant treatment. Most patients reported cognitive symptoms and over half were shown to have impaired cognitive performance by DSST. Mean DSST score at the time of treatment of initiation in this study (50.7) was similar to that reported in Japanese patients initiating new antidepressant therapy in another recent study (54.3),⁴⁴ but higher than those reported in comparable US and European populations (42.0 in FOCUS and 43.1 in CONNECT). 36,45 This may at least partly reflect cultural differences between the different geographic populations. Levels of impairment of HRQoL and work productivity were consistent with those reported in studies in similar MDD populations in other countries. 18,20,21,23-25 Our findings also suggest that HRQoL may be more impaired by MDD than by other common chronic diseases in Chinese patients. The reported mean EQ-5D utility index score in this study was lower than reported for Chinese patients with other chronic health conditions, such as diabetes, hypertension, and epilepsy, 46 and similar to those reported for Chinese patients with a range of cancer types.47

Patients with more severe perceived cognitive symptoms were shown to experience greater impairments in HRQoL and work productivity (absenteeism and presenteeism), even after adjustment for depression severity. In contrast, less robust correlations were seen between observed cognitive performance (assessed using the DSST) and HRQoL, with no associations seen between DSST and work productivity scores. This finding is not unexpected,

as several other studies have also found discrepancies between self-reported measures and performance tests of cognitive symptoms in patients with MDD. ^{25,26,48-51} For example, in a study in the United States involving 602 patients with MDD reporting cognitive symptoms, 64% met criteria for objectively defined cognitive impairment assessed using a battery of neuropsychological tests (including the DSST) and 48% met criteria for subjectively defined cognitive impairment assessed using the Cognitive and Physical Functioning Questionnaire (CPFQ). ⁵⁰ Overall, 80% of patients met criteria for either subjective or objective cognitive impairment; however, only 31% met both criteria.

The observed dissonance between objective and subjective cognitive performance in patients with MDD may be due to differences in the cognitive subdomains assessed by the various questionnaires and tests. There is also evidence to suggest differential effects of potential moderating factors. For example, data from the Cognitive Dysfunction in Asians with Depression (CogDAD) study show depression severity to be predictive of subjective cognitive performance, but not objective measures of cognition.⁴⁹ From a clinical perspective, subjective and objective methods for assessing cognition complement each other and both methods are valuable. While the potential for bias cannot be excluded with self-assessment of cognitive symptoms, it is possible that subjective assessment of cognitive symptoms may more closely reflect the degree of impairment experienced by patients in daily life than objective assessment in a highly standardized test setting.⁵²

As in other recent studies in patients with depression, ^{20,23,25,26,53,54} presenteeism was found to be more prevalent than absenteeism in the study population. Patients reported missing an average of 2 working days during the previous week and over half of their time spent at work was deemed unproductive (3.7 days). Mean WPAI absenteeism and presenteeism scores in this study (26.3% and 54.5%) were considerably higher than those reported in a US study of patients with depression initiating anti-depressant treatment (8.2% and 35.2%). MDD-related work impairment has been shown to be associated with substantial economic costs. ^{8,53,56}

Coupled with the results from the primary analysis of the PROACT study, in which structural equation modelling showed severity of cognitive symptoms to consistently predict both clinical and functional outcomes over a 6-month period independent of depressive symptom severity,²⁹ our findings suggest that cognitive symptoms Dovepress Wang et al

should be considered an important and independent treatment target in Chinese patients with MDD. Results of a recent meta-analysis suggest differential effects of currently available antidepressants in terms of improving cognitive symptoms in patients with MDD.⁵⁷

A major strength of the PROACT study is that it was performed in a large and well-defined population of Chinese patients with MDD in a real-world setting; hence, findings should be generalizable to routine practice. In addition, patient-reported outcome measures were used to assess disease severity and impact on HRQoL and work productivity from the patient's own perspective; this is known to be particularly valid in patients with mental health disorders such as MDD.⁵⁸ A potential limitation is that the study population mainly comprised well-educated, urban residents initiating first- or second-line antidepressant monotherapy, which may limit generalization of the study findings to other groups of patients. Furthermore, as norms are not available for Chinese patients, age-related norms for US adults were used in the DSST subgroup analyses described in this paper. However, to address this potential limitation, DSST analyses were also performed according to DSST total-score quartile at baseline.

In summary, results of this real-world study illustrate the impact of cognitive symptoms on HRQoL and work productivity in Chinese patients with MDD, and highlight the importance of assessing and targeting cognitive symptoms in order to improve functional outcomes when treating patients with MDD.

Abbreviations

DSST, Digit Symbol Substitution Test; EQ-5D, EuroQoL 5-Dimensions Questionnaire; HAM-D₁₇, 17-item Hamilton Depression Rating Scale; HRQoL, health-related quality of life; MDD, major depressive disorder; PDQ-D, Perceived Deficits Questionnaire-Depression; PHQ-9, 9-item Patient Health Questionnaire; PROACT, Prospective Research Observation to Assess Cognition in Treated patients with MDD; SD, standard deviation; SDS, Sheehan Disability Scale; WPAI-SHP, Work Productivity and Activity Impairment – Specific Health Problems Questionnaire.

Data Sharing Statement

The authors declare that the data relating to the reported analyses are available within the article. H. Lundbeck A/S may be contacted for further data sharing, including individual-level data.

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Author Contributions

GW, HR and LH-H were involved in conceptualization and design of the study, GW was involved in the acquisition of data, TKHX was responsible for data analysis, and all authors contributed to data interpretation. The authors are entirely responsible for the scientific content of this paper and all authors contributed to drafting or revising the article, gave final approval of the version to be published, and agree to be accountable for all aspects of the work.

Disclosure

GW has received honoraria for being an advisor to or providing educational talks for Lundbeck, Pfizer, Sumitomo, Johnson & Johnson, and Eli Lilly and Company. TKHX was a full-time employee of Lundbeck Singapore Pte Ltd when the study and analyses were performed. HR and LH-H are full-time employees of H. Lundbeck A/S. The authors report no other conflicts of interest in this work.

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