

# Correlations Between Working Memory Impairment and Neurometabolites of the Prefrontal Cortex in Drug-Naive Obsessive-Compulsive Disorder

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**Purpose:** This study aimed to investigate the mechanism of working memory (WM) impairment in drug-naive obsessive-compulsive disorder (OCD) by using neuropsychological tests and proton magnetic resonance spectroscopy (<sup>1</sup>H-MRS).

**Patients and Methods:** A total of 55 patients with drug-naive OCD and 55 healthy controls (HCs) were recruited for this study. The working memory (WM) was evaluated using the digit span test (DST), visual space memory test (VSMT), and the 2-back task and stroop color word test (SCWT). The bilateral metabolite levels of the prefrontal cortex (PFC) were evaluated by <sup>1</sup>H-MRS, then determined the ratios of N-acetyl aspartate (NAA), choline-containing compounds (Cho), and myo-inositol (MI) to creatine (Cr). The independent sample *t*-test was used to analyse the differences in WM performance and neurometabolite ratios. Multivariate linear regression analysis was performed to screen the influential factors of WM, with an introduction level of 0.05 and a rejection level of 0.10.

**Results:** 1) Patients with OCD performed significantly worse on DST (score), VSMT (score), 2-back task (accuracy rate), SCWT (execution time) when compared with HCs. 2) NAA/Cr and Cho/Cr in the left PFC (IPFC) and MI/Cr ratios in the bilateral PFC of OCD patients were significantly lower when compared to HCs. 3) For OCD patients, the NAA/Cr ratio in the IPFC was negatively correlated with the score of DST (forwards), the Cho/Cr ratio in the IPFC was positively correlated with the accuracy rate of 2-back task, and the MI/Cr ratio in the right PFC (rPFC) was positively correlated with the score of DST (forwards) and the accuracy rate of VSMT. We also found that the compulsive symptoms showed a positive correlation with MI/Cr ratio of the rPFC.

**Conclusion:** Drug-naive OCD patients have demonstrated WM impairments, including phonological loop, visual-spatial sketchpad and central executive system, and the WM impairments might be associated with hypometabolism in the PFC, especially the IPFC.

**Keywords:** obsessive compulsive disorder, prefrontal cortex, working memory, neurometabolites, proton magnetic resonance spectroscopy

## Introduction

Obsessive compulsive disorder (OCD) is a severe, chronic, and complex neuropsychiatric disorder,<sup>1,2</sup> with a lifetime prevalence of 1% to 3%.<sup>3,4</sup> The incidence of OCD has recently shown an upward trend, but its diagnosis and treatment are often delayed. In most cases of OCD, it usually takes approximately 8 years from symptom onset to medical treatment,<sup>5,6</sup> and the effective rate of treatment remains only between 40% - 60%.<sup>7</sup>

Most studies in literature have confirmed that OCD involves cognitive impairment, which is considered as one of the neurophenotypic markers of OCD.<sup>8–12</sup> Working memory (WM) is considered a key cognitive function<sup>13</sup> and is closely related to OCD.<sup>14–16</sup> WM is a psychological process that involves storing and processing of information when performing tasks and activities, which include verbal WM, visuospatial WM and central executive system (including updating, shifting and inhibitory functions). WM also supports other high-level cognitive abilities (including but not limited to general fluid intelligence, learning, problem solving and decision-making),<sup>17,18</sup> as well as low-level psychological operations that require cognitive control.<sup>19,20</sup> Recurrent intrusive thoughts and/or compulsions in OCD are often repetitive behaviors or mental acts, followed by checking, rumination, and disinhibition,<sup>1</sup> and these are associated with poorer memory in OCD. For example, memory deficits might explain the compulsive checking behavior.<sup>21</sup> Impaired response inhibition accounts for unstoppable repetitive rituals in OCD patients<sup>22</sup>. There exists a large body of working memory literature on OCD, and yet results from this literature are highly mixed, such as visuospatial WM and verbal memory.<sup>15,23,24</sup> The inconsistent pattern of results described is possibly attributed to several factors, which include the highly heterogeneous nature of OCD, the high rate of comorbidity, and incomplete or inconsistent measurement tools. It is also possible that the mechanism of WM impairment in OCD remains unclear. Therefore, we used a more comprehensive assessment tools to study WM, including Digit Span Test (DST), Visual Reproduction Subtest (VSMT), 2-back task and Stroop Color Word Test (SCWT). DST mainly reflects the ability of spatial WM capacity as well as verbal WM, which is included in the phonological loop and episodic buffer subsystem. VSMT is used to measure the visuospatial WM, which is assumed to be responsible for processing and short-term storage of visuospatial information and is the part of visual-spatial sketchpad and episodic buffer.<sup>25,26</sup> The 2-back task is one of the verbal n-back tasks, which is a commonly used paradigm to investigate verbal WM and updating function of the central executive system. SCWT is used to evaluate the updating, shifting and inhibitory functions of the central executive system.

A growing body of evidence in imaging studies also revealed neurobiological abnormalities in OCD, which is speculated to be a potential neuroimaging endophenotype.<sup>27–30</sup> It has been particularly convincing

in suggesting that prefrontal–basal ganglia–thalamic–prefrontal circuits are responsible for the mediation of OCD symptoms.<sup>29,30</sup> Complementary approaches examining regional neurochemistry now assure additional insights into the neurobiology of OCD. Some studies have found that the imbalances in neuronal metabolite and neurotransmitter within CSTC circuit have been shown as the leading reasons of the OCD onset.<sup>31</sup> In particular, the proton magnetic resonance spectroscopy (<sup>1</sup>H-MRS) permits in vivo quantification of specific neurochemicals in various regions of the brain. The most commonly reported resonances of <sup>1</sup>H-MRS spectrum are N-acetyl aspartate (NAA), choline-containing (Cho) compounds, myo-inositol (MI), and creatine (Cr). The pathophysiology of OCD has been studied by <sup>1</sup>H-MRS, but the results are still inconclusive.<sup>32,33</sup> Like Pauls et al comments, there is little agreement between these investigations,<sup>28</sup> and this is probably because of several factors, such as the sample size, in which most of the studies included low patient samples, with an average of 13 subjects per study. Secondly, the heterogeneity of the disease, wherein the samples are often included into heterogeneous groups (comorbidities, drugs/treatment, untreated, etc.). Thirdly, with regard to the inspection equipment of <sup>1</sup>H-MRS technique, only 4 (14%) of 28 studies have used field strength  $\geq 3$ -Tesla (3T).<sup>28,34</sup> In order to reduce the above interference in the current study, we examined biochemical metabolites in the prefrontal cortex (PFC) of treatment-naïve, non-late-life OCD patients without comorbidity by using multi-voxel 3.0T <sup>1</sup>H-MRS. The multi-voxel technique used in our study possesses the advantage of simultaneously obtaining multiple <sup>1</sup>H-MRS spectra and allowing for examination of a larger volume of tissue than that by single voxel spectroscopy. In addition, the spectra of the normal human brains at 3.0T demonstrated a 20% improvement of the signal to noise ratios compared to 1.5T at a short TE.

The prefrontal cortex (PFC) is a key area of cortico-striato-thalamo-cortical (CSTC) circuit,<sup>30</sup> and dysfunction in the PFC might be constituted as a neurophenotype of OCD. However, there is very little research till now that has been conducted in OCD patients on their WM and neurometabolites of the PFC. A combination of metabolic brain alterations with neuropsychological tests might be an effective way to explore the correlation between brain function change and OCD.

Therefore, we examined WM and biochemical metabolites in PFC of patients with drug-naïve OCD in this

study. We hypothesized that the deficits of WM and abnormal biochemical metabolism could be found in patients with drug-naive OCD. We were also interested in the relationship between WM and abnormal biochemical metabolism in drug-naive OCD. During this approach, characteristic changes of WM and metabolites in OCD patients were explored, and to find out the potentially unknown associations between neural mechanism and WM impairment in OCD.

## Materials and Methods

### Participants

A total of 55 patients affected by OCD (Diagnostic and Statistical Manual of Mental Disorder, 5th Edition - criteria, DSM-5) (APA, 2013) (29 males, 26 females; mean  $\pm$  SD: 26.70 $\pm$ 7.43 years of age), and 55 healthy controls (HCs) (21 males, 34 females; mean  $\pm$  SD: 25.33 $\pm$ 5.28 years of age) were studied. All subjects were right-handed and belonged to the Han ethnic group. All patients and HCs voluntarily participated in this study and signed an informed consent.

### OCD Group

Between March 2014 and July 2015, 55 adults with OCD were recruited from the psychiatric department of the First Affiliated Hospital of Jinan University (Guangzhou, China). All patients were assessed by psychiatrists involved in the study. Written informed consent was obtained from all participants before any study procedure was performed.

The inclusion criteria of OCD group were as follows: diagnosed with OCD according to the DSM-5 criteria, aged between 18 and 50 years, educational level above junior high school, with a total score of the Yale-Brown Obsessive-Compulsive Scale (Y-BOCS) of  $\geq 16$  points (if only one of the compulsive behaviors or compulsive ideas exist, the evaluation standard then becomes to the total number of Y-BOCS scores of  $\geq 8$  points), psychiatric drug naive and treatment naive.

Exclusion criteria of OCD group were as follows: any other psychiatric axis-I or axis-II disorders, pregnant or lactating females, any other neurological disorders, brain organic mental disorders or somatic complaints, a history of alcohol or drug abuse, abnormal brain structure or magnetic resonance contraindications, achromatopsia or hemochromatosis, those who received electric convulsive treatment (ECT) recently or had long-term anti-psychotic drugs.

### Healthy Control (HC) Group

A sample of 55 healthy volunteers was recruited by advertising. The inclusion criteria of control group were as follows: healthy adults, aged between 18 and 50, educational level above junior high school.

Exclusion criteria of control group were as follows: a history of psychiatric illness or a family history of mental disorders, pregnant or lactating females, brain organic mental disorders or somatic complaints, a history of alcohol or drug abuse, abnormal brain structure or magnetic resonance contraindications, and achromatopsia or hemochromatosis.

### Clinical Assessments

A general information questionnaire was utilized to investigate the subject's gender, age, nationality, marital status, profession, education, history of somatic diseases, history of drug allergy, status of smoking and alcohol addiction.

Y-BOCS: OCD symptom characteristics and severity were assessed using Y-BOCS. Those with a total score of 25 or more (simple obsessive thinking or compulsive behavior, only 15 points or more) were considered severe; 16–25 points (mere obsessive thinking or compulsive behavior, only 10–14 points) were considered moderately severe; 10–15 points were considered mildly severe; and 10 points were considered extremely low.

### WM Assessments

DST (Wechsler, 2000) was used to measure the WM capacity and verbal WM, which consists of digit span forward (DSF) and digit span backward (DSB). Correct responses on the DSF and DSB were calculated. DST mainly reflects the ability of spatial WM capacity as well as verbal WM, which is included in the phonological loop and episodic buffer subsystem.

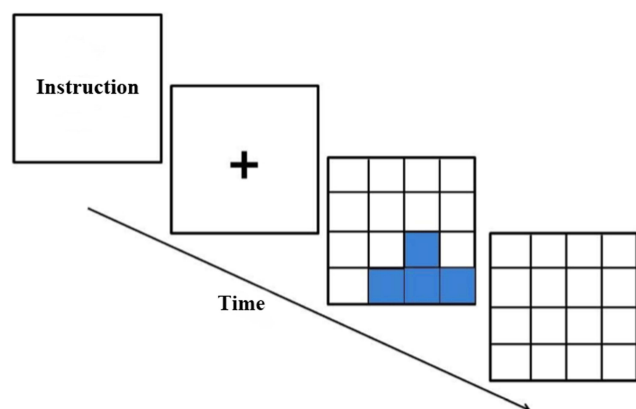
2-back task is one of verbal n-back task, which is a commonly used paradigm to investigate verbal WM and updating function of the central executive system. The 2-back task was programmed with E-prime 2.0 software. During the test, any number from 1 to 9 is randomly displayed in the center of the computer screen, each number is displayed for 1000 milliseconds, and the number is separated by 1000 milliseconds. If the currently displayed number is consistent with the previously displayed number, press "F" key; if the currently displayed number is inconsistent with the previous number, press "J" key. Reaction time and accuracy rate on the 2-back were calculated.

VSMT was also programmed with E-prime 2.0 software, which was used to measure the visuospatial WM (see Figure 1). During the test, a 4\*4 black-and-white grid matrix appears in the center of the computer screen, in which blue patches are randomly distributed. After 1000 milliseconds, the blue color block disappeared, and then the participant marked the position of the blue color block in the blank square matrix. Subjects were requested again to recall the spatial location as quickly and as accurately as possible. Reaction time and accuracy rate on the VSMT were calculated.

SCWT-CW was used to evaluate the updating, shifting and inhibitory functions of the central executive system. The stimulus card contains a total of 100 characters. Participants were required to name the color of each Chinese character in the displayed symbol card in turn (for example, the “green” character displayed in yellow font is named “yellow”). The reaction time on the SCWT-CW was calculated.

## Image Acquisition

Magnetic resonance imaging (MRI) and <sup>1</sup>H-MRS were performed using the Discovery MR 750 3.0 T MRI scanner (General Electric, Milwaukee, WI, USA) with a conventional gradient system. The gradient field strength used was 44mT/m and the gradient switching rate was 200T/m/s. An 8-channel head coil was used for radio-frequency transmission and reception of MR signal. The subjects' heads were fixed by a sponge pad, in which the center line was in the middle, and symmetrical on both sides in order to avoid motion artifacts.

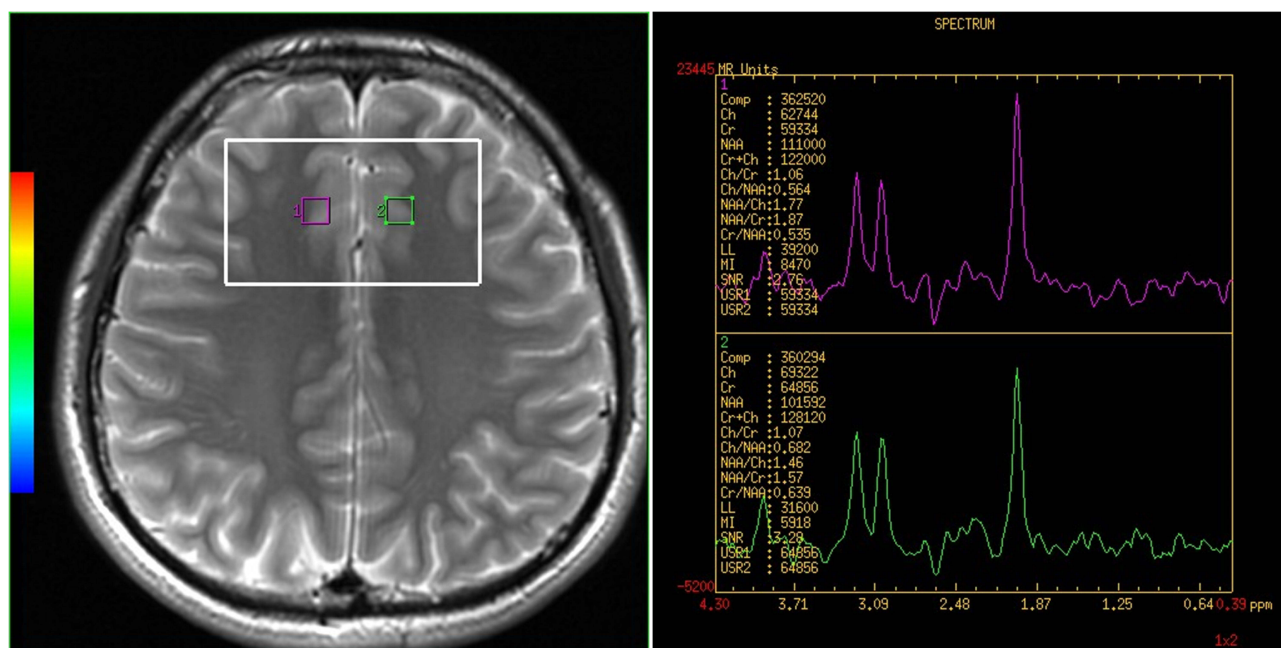


**Figure 1** During the test, a 4\*4 black-and-white grid matrix appears in the center of the computer screen, in which blue patches are randomly distributed. After 1000 milliseconds, the blue color block disappeared, and then the participant marked the position of the blue color block in the blank square matrix. Subjects were requested again to recall the spatial location as quickly and as accurately as possible.

Routine axial T1-weighted fluid attenuation inversion recovery (T1Flair) [repetition time (TR) = 1750 ms, echo-time (TE) = 24 ms] and fast spin echo T2-weighted MR images (TR = 8400ms, TE = 145ms) were obtained to confirm the absence of any structural and signal abnormalities of the brain.

The magnetic resonance protocol was conducted in the axial plane with a T2-weighted fast spin-echo (FSE) sequence (TR = 3500 ms, TE = 102 ms, number of excitation (NEX) = 2, slice thickness = 5 mm, slice interval = 0 mm, 18 slices, field of view (FOV) = 24×24 cm, matrix size = 256×256). For <sup>1</sup>H-MRS studies, the location of the PFC volume of interest (VOI) was showed in Figure 2. The prefrontal lobe VOI was localized at the midline and included the dorsolateral prefrontal white matter and anterior cingulate gray regions bilaterally avoiding the striatum, ventricle, scalp, skull base, or sinuses. The posterior boundary of the VOI was placed adjacent to the anterior margin of the frontal horn of the lateral ventricle. The VOI was placed in a uniform manner by the same investigator. Forty-nine nominal voxels (7.5×7.5×10mm<sup>3</sup>) was included in the VOI. Single section 2D multivoxel <sup>1</sup>H-MRS was acquired using a point-resolved spectroscopy sequence (PRESS) with water suppression by a chemical shift selective saturation (CHESS) pulse. The acquisition parameters were as follows: TR = 1000 ms, TE = 144 ms, NEX = 1, matrix size = 18x18, FOV = 24×24 cm, slice thickness = 10 mm. Additional saturation bands were placed outside the VOI to minimize lipid contamination from the scalp. Automatic pre-scanning was performed before each spectroscopic scan to achieve an optional full width half maximum of 10 Hz. As a general quality standard spectrum, a line width of above 10 Hz or water suppression of above 98% was excluded. The acquisition time for <sup>1</sup>H-MRS sequence was 5 min and 28s.

The analyses of spectral dataset were performed using the Function Tool of GE 3.0 T workstation (Sun, Advantage Windows ADW4.5). The distribution maps of different metabolites were obtained, and then the distribution maps and MRI images were fused to obtain <sup>1</sup>H-MRS curves. The software was used to automatically conduct the phase, frequency coding and baseline correction for the spectrum. The built-in software of MRI scanner automatically conducted the baseline calibration, signal average and metabolite identification, and measured the peak area of NAA, Cho, MI and Cr, and calculated the NAA/Cr, Cho/Cr and MI/Cr ratios. Each spectrum was evaluated for the peak area of NAA at 2.02 ppm, Cho at 3.22 ppm, MI at 3.0 ppm and Cr at 3.03 ppm. The voxel placements for spectroscopy and all data



**Figure 2** Magnetic resonance image (MRI) scan of healthy controls showing the location of magnetic resonance spectroscopy (MRS) of volume of interest (VOI) placed in the PFC and proton magnetic resonance spectra in the left and right PFC. The large white box represents the VOIs for MRS acquisition, and small white boxes depict the individual VOIs for spectral analysis.

**Abbreviations:** NAA, N-acetylaspartate; Cho, choline; ml, myo-inositol; Cr, creatine.

analyses were carried out by a trained radiologist who was blinded to the diagnosis of each subject.

### Statistical Analysis

SPSS Statistical Package Version 20.0 was used for statistical analysis, and two-tailed significance level was set at  $p < 0.05$ . All indicators were measured for normal distributions by the Kolmogorov–Smirnov goodness-of-fit test. When comparing group differences in terms of demographics and clinical data, the  $t$ -test was used if continuous variables were normal. The Mann–Whitney  $U$ -test was used if continuous variables

were skewed. Independent sample  $t$ -test (normal variables) or Mann–Whitney  $U$  (skewed variables) was used to test the differences in WM performance and neurometabolite ratios between OCD and control groups. Bonferroni correction for multiple comparisons was applied and  $p < 0.004$  ( $0.05/14$ ) was considered significant. Multivariate linear regression analysis was performed using WM of the case group as the dependent variable, and NAA/Cr, Cho/Cr, and MI/Cr ratios of the PFC as independent variables to screen the influential factors of WM, with an introduction level of 0.05 and rejection level of 0.10.

**Table 1** Characteristics of Patients with OCD and HCs

Factors	OCD (n=55)	Control (n=55)	Statistical Analysis	
			$F/X^2$	$p$ value
Gender (male, female)	29, 26	21, 34	$X^2=2.347$	0.126
Age (years)	26.70±7.43	25.33±5.28	$F=1.122$	0.265
Education (years)	14.40±2.66	14.62±1.95	$F=-0.335$	0.739
Y-BOCS score				
Total	19.51±3.70	—	—	—
Obsession scores	10.53±2.23	—	—	—
Compulsion scores	9.00±2.33	—	—	—
Duration of illness (months)	40.58±5.41	—	—	—

**Abbreviation:** Y-BOCS, Yale–Brown Obsessive–Compulsive Scale.

**Table 2** Comparison of WM Indices Between Drug-Naive OCD and HCs

Working Memory Tests	OCD	Control	t	PI	P2	ES
DSF	8.22±1.46	9.07±1.12	3.45	0.001	OCD<Control <sup>f</sup>	-0.31
DSB	5.96±1.63	6.76±1.89	2.38	0.019		-0.22
DST (forwards+backwards)	14.19±2.53	15.83±2.54	3.43	0.001	OCD<Control <sup>f</sup>	-0.31
2-back task (reaction time ms)	3058±1198	2671±1226	1.70	0.093		0.16
2-back task (accuracy rate)	16.53±3.56	18.69±1.98	3.96	<0.001	OCD<Control <sup>f</sup>	-0.35
VRST (reaction time ms)	1781±923	1388±687	2.57	0.011		0.23
VRST (accuracy rate)	3.04±1.87	4.52±0.84	5.39	<0.001	OCD<Control <sup>f</sup>	-0.45
SCWT (execution time s)	16.13±3.51	13.49±3.54	3.97	<0.001	OCD<Control <sup>f</sup>	0.35

**Abbreviations:** DSF, digit span forward; DSB, digit span backward; DST, Digit Span Test; VRS, Visual Reproduction Subtest; SCWT, Stroop Color and Word test; P2, Bonferroni test; ES, effect size.

**Table 3** Comparison of Neurometabolite Ratios in the PFC Between Drug-Naive OCD and HCs

	OCD	Control	t	PI	P2	ES
IPFC						
NAA/Cr	2.06±0.22	2.38±0.28	6.31	<0.001	OCD<Control <sup>f</sup>	-0.54
Cho/Cr	1.09±0.10	1.41±0.40	5.72	<0.001	OCD<Control <sup>f</sup>	-0.48
MI/Cr	0.15±0.18	0.25±0.27	2.29	0.024		-0.21
rPFC						
NAA/Cr	2.33±0.30	2.31±0.25	0.51	0.611		0.04
Cho/Cr	1.26±0.28	1.20±0.17	1.35	0.179		0.13
MI/Cr	0.10±0.08	0.16±0.11	3.002	0.003	OCD<Control <sup>f</sup>	0.30

**Abbreviations:** l/rPFC, left/right of prefrontal cortex; NAA, N-acetyl-aspartate; Cho, choline; MI, myo-inositol, Cr, creatine; ES, effect size; P2, Bonferroni test.

## Results

### Sample Characteristics

The demographics and clinical data of OCD group and HCs are shown in Table 1. A total of 55 patients with drug-naive OCD and 55 HCs were enrolled in this study. The two groups were matched by age ( $F = 1.122$ ,  $p = 0.265$ ), gender ( $X^2 = 2.347$ ,  $p = 0.126$ ) and educational level ( $F = -0.335$ ,  $p = 0.739$ ). In the Y-BOCS score of all patients, the minimum score was 12 and the maximum score was 32 points.

### The Comparison of WM Indices Between Drug-Naive OCD and HCs

The independent sample *t*-test was used to compare the WM indices between the two groups, and the results are shown in Table 2. Compared to HCs, patients with OCD had significantly worse DST, VSMT, 2-back task (accuracy rate), and SCWT (execution time) (all  $p < 0.01$ ). No statistical differences of 2-back task (reaction time) were observed between the two groups ( $p > 0.05$ ).

### Comparison of Neurometabolite Ratios in the PFC Between Drug-Naive OCD and HCs

The independent sample *t*-test was used to compare the NAA/Cr, Cho/Cr, and MI/Cr ratios of IPFC and rPFC between the two groups, and the results are shown in Table 3. Compared to HCs, patients with OCD had more reduction of NAA/Cr, Cho/Cr, MI/Cr ratios in the IPFC, while MI/Cr ratio in the rPFC, showed statistically significant differences ( $p < 0.05$ ). There were no significant differences in the ratios of NAA/Cr and Cho/Cr of the rPFC between the two groups ( $p > 0.05$ ).

### Correlations Between Abnormal Neurometabolite Ratios and WM Indices, Demographics and Clinical Symptoms in Patients with Drug-Naive OCD

The standardized residual histogram of WM showed an approximated normal distribution. A simple linear regression model was performed with NAA/Cr, Cho/Cr, and MI/Cr ratios of PFC as independent variables and age, gender, education

**Table 4** Multiple Regression Analysis in Patients with OCD

Dependent Variables	Independent Variables	Standardized Coefficients	t	P	
DSF	NAA/Cr of IPFC	-0.306	-2.306	0.025	
	Cho/Cr of IPFC		-1.043	0.302	
	MI/Cr IPFC		0.953	0.345	
	NAA/Cr of rPFC		0.767	0.446	
	Cho/Cr of rPFC		0.982	0.331	
	MI/Cr of rPFC		0.278	2.095	0.041
	DSB		NAA/Cr of IPFC		-0.515
Cho/Cr of IPFC		-0.181	0.857		
MI/Cr of IPFC		0.965	0.339		
NAA/Cr of rPFC		0.292	0.771		
Cho/Cr of rPFC		0.767	0.447		
MI/Cr of rPFC		0.748	0.458		
DST (forwards+backwards)		NAA/Cr of IPFC			
	Cho/Cr of IPFC				
	MI/Cr of IPFC	-1.484		0.144	
	NAA/Cr of rPFC	-0.622		0.537	
	Cho/Cr of rPFC	0.665		0.509	
	MI/Cr of rPFC	0.303		0.763	
		1.112		0.271	
VRST (accuracy rate)	NAA/Cr of IPFC	0.301	-0.36	0.72	
	Cho/Cr of IPFC		0.01	0.992	
	MI/Cr of IPFC		-0.82	0.416	
	NAA/Cr of rPFC		-0.81	0.422	
	Cho/Cr of rPFC		-1.078	0.286	
	MI/Cr of rPFC		2.295	0.026	
	VRST (reaction time ms)		NAA/Cr of IPFC	0.253	1.79
Cho/Cr of IPFC		1.114	0.27		
MI/Cr of IPFC		0.468	0.642		
NAA/Cr of rPFC		-0.259	0.797		
Cho/Cr of rPFC		0.637	0.527		
MI/Cr of rPFC		1.251	0.217		
2-back task(accuracy rate)		NAA/Cr of IPFC	0.358		-1.13
	Cho/Cr of IPFC	2.791		0.007	
	MI/Cr of IPFC	0.67		0.506	
	NAA/Cr of rPFC	-0.604		0.548	
	Cho/Cr of rPFC	1.198		0.236	
	MI/Cr of rPFC	1.207		0.233	
	Compulsive symptom scores	MI/Cr of rPFC		0.270	2.045

**Note:** \* $p < 0.05$  significant.

**Abbreviations:** l/rPFC, left/right of prefrontal cortex; NAA, N-acetyl-aspartate; Cho, choline; MI, myo-inositol, Cr, creatine; DSF, digit span forward.

level, duration of illness, Y-BOCS score and WM as a dependent variable. Multivariate linear regression stepwise regression method was used to screen the influential factors of WM. The results are shown in Table 4.

For OCD patients, the DSF showed a positive correlation with MI/Cr ratios of the rPFC and a negative correlation with NAA/Cr ratios of the IPFC ( $F = 4.062$ ,  $R^2 = 10.4\%$ ,  $p < 0.05$ ),

implying that approximately 10.4% of the DSF scores can be explained by the MI/Cr ratios of the rPFC and NAA/Cr ratios of the IPFC. The VSMT (accuracy rate) showed a positive correlation with MI/Cr ratios of the rPFC ( $F = 5.266$ ,  $R^2 = 0.09$ ,  $p < 0.05$ ), implying that approximately 9% of the VSMT (accuracy rate) scores can be explained by MI/Cr ratios of the rPFC. The 2-back task (accuracy rate) showed a positive

correlation with Cho/Cr ratios of the IPFC ( $F = 7.79$ ,  $R^2 = 0.128$ ,  $p < 0.05$ ), implying that approximately 12.8% of the 2-back task (accuracy rate) scores can be explained by Cho/Cr ratios of the IPFC.

The compulsive symptoms showed a positive correlation with MI/Cr ratios of the rPFC ( $F = 3.96$ ,  $R^2 = 0.132$ ,  $p < 0.05$ ), which implied that approximately 13.2% of the compulsive symptom scores can be explained by the MI/Cr ratios of the rPFC. We found no significant correlation between neurometabolite ratios and the other demographics and clinical symptoms.

## Discussion

The major study findings are as follows. Patients with OCD performed significantly worse on DST (score), VSMT (score), 2-back task (accuracy rate), SCWT (execution time) when compared with HCs, and showed significantly lower NAA/Cr and Cho/Cr ratios in the left PFC (IPFC) and lower MI/Cr ratios in the bilateral PFC. For OCD patients, lower NAA/Cr ratio in the IPFC was negatively correlated with the score of DST (forwards), lower Cho/Cr ratio in the IPFC was positively correlated with the accuracy rate of 2-back task, and the MI/Cr ratio in the right PFC (rPFC) was positively correlated with the score of DST (forwards) and the accuracy rate of VSMT.

The verbal WM is measured by DST and the 2-back task methods. Results showed significant deficits in the performance with DST and 2-back task (accuracy rate), suggestive of a significant impairment of verbal WM in the OCD. This result was consistent with that of previous research findings.<sup>9,15</sup> DSF showed a significant correlation with MI/Cr ratios of the rPFC and NAA/Cr ratios of IPFC, while the 2-back task (accuracy rate) showed a significant correlation with Cho/Cr of IPFC. Patients showed a significant decrease in NAA/Cr and Cho/Cr ratios in the IPFC, and MI/Cr ratios in the rPFC when compared to HCs. NAA is one of the most prominent metabolites in magnetic resonance spectrograms of the brain, and is considered as a marker of neuronal integrity and neuron density.<sup>35</sup> Furthermore, the patients showed a significant decrease in NAA/Cr ratios in the IPFC and showed a negative correlation with DSF. Lentz et al investigated postmortem frontal lobe tissue by using high-spectral-resolution 1H MR spectroscopy of brain extracts and found NAA/Cr as a sensitive marker of neuronal injury.<sup>36</sup> 1H-MRS studies demonstrated decreased NAA concentration during psychiatric conditions, including affective disorder, OCD, schizophrenia and dementia,

and the decrease of NAA levels is often located in regions of gray matter volume reduction, indicating neuronal reduction.<sup>35,37</sup> Additionally, a study with the effect of 12-week treatment using citalopram on NAA/Cr ratios in 13 drug-naive patients with OCD showed a significant increase in the bilateral prefrontal cortex, frontal white matter and anterior cingulate when compared to those before treatment.<sup>38</sup> So, the decrease of NAA/Cr ratios in the IPFC might indicate gray matter volume reduction and neuronal damage in OCD, which is closely related to verbal working memory impairment.

In the literature, there are rare reports on the decrease of choline levels of IPFC in OCD. Choline plays a central role in membrane structure and many physiological pathways, including the membrane-mediated cell signaling via phospholipases, acetylcholine synthesis, lipid transport, myelination, growth factor signaling, and methyl-group metabolism.<sup>39</sup> Studies have found that choline deficiency can cause nerve cell damage or induce apoptosis, while choline supplementation can inhibit apoptosis and promote neural plasticity.<sup>40,41</sup> There are many studies on choline metabolism in patients with OCD, and most of these focused on thalamus, parietal white matter, hippocampus, striatal, caudate nucleus, and the anterior cingulate cortex among others.<sup>32,33</sup> In this study, the patients showed a significant decrease in Cho/Cr ratios in the IPFC and showed a negative correlation with the 2-back task (accuracy rate). The 2-back task was carried out to investigate verbal WM and updating the function of the central executive system. Updating is the constant monitoring and rapid addition/deletion of WM contents, which can be measured through tests such as the n-back, which are typically considered as the probes of WM.<sup>42</sup> The ability to acquire new information and manipulate it in real time is critical to adjust one's behavior to meet the demands of the changing environment.<sup>25</sup> So, the impairment of updating ability results in behavioral and cognitive inflexibility and this seems to be consistent with the clinical symptoms of OCD. Moreover, in this study, we also found that SCWT (execution time) showed significant deficits in the performance of the drug-naive OCD group, but we did not find its relationship to the neurometabolites in the brain. SCWT (execution time) can reflect the inhibition function and mainly involves prevention of target task being intervened by irrelevant information, suppressing reactions that are not in conformity with the situation and preventing secondary activation of unwanted internal information in memory. Our results in agreement with those of the previous studies, in which a prominent disturbance in the ability to inhibit the WM of OCD has been reported.<sup>16</sup> We speculate that the lack of choline



in IPFC may be closely related to the defects of verbal WM and the central executive system in OCD. Choline supplementation might be able to improve this condition, which may provide another option for the treatment of obsessive-compulsive disorder, but further research is needed in the future.

The visuospatial WM is measured by VSMT. In the current study, the results showed significant deficits in performance with VRST (accuracy rate), suggesting a significant impairment of the visuospatial WM in the OCD. When compared to HCs, patients with OCD showed a significant decrease of MI/Cr ratios in the rPFC, which showed a positive correlation with VSMT (accuracy rate). Visual-spatial WM is mainly involved in the maintenance and control of visual and spatial information. Disturbance of the visuospatial WM suggests that OCD has disturbances in maintaining and controlling external information, similar results were found in previous study, but measured it with Cambridge Neuropsychological Automated Battery (CANTAB).<sup>43</sup> Myoinositol (MI) is located mainly in the glial cells of the brain, and is considered as a glial marker, and its content is believed to relate with glial cells proliferation.<sup>44,45</sup> It might participate in the protection against osmotic shock, signaling pathways, vesicle endocytosis and exocytosis.<sup>44</sup> Alterations in the MI levels of brain are associated with diseases of central nervous system, such as schizophrenia, bipolar disorders, and cognitive impairment.<sup>45–47</sup> 1H-MRS findings in OCD have reported less on the alterations of MI in OCD subjects before 2012.<sup>33</sup> Some recent studies on OCD have also found changes in the levels of inositol metabolism in the brain. Parmar et al<sup>38</sup> have found significant increase in inositol levels in the caudate nucleus and anterior cingulate cortex of OCD. It is positively correlated with the compulsive symptoms and disease severity, and it is speculated to be related to the potential reversible abnormalities in regulating the neuro-osmotic pressure. Another study<sup>48</sup> measured the metabolite levels in the left and right prefrontal white matter and found that children and adolescents with OCD did not differ from those of controls in MI metabolite levels. The metabolic concentration of MI in patients with OCD is increased significantly by 52% when compared with HCs. However, this study included a small sample size (only 14 OCD patients and 14 healthy individuals), and some patients received medications (escitalopram or fluvoxamine). Also, we found out that the levels of MI in bilateral PFC were reduced, which has not been reported in previous studies. MI is also a brain metabolite related to important functions of phosphatidylinositol second messenger system (PI-cycle), and is associated with mental disorders such as OCD, generalized anxiety disorder or schizophrenia.<sup>49</sup> MI is transformed by

various receptors attached to the PI-cycle, and its concentration or function changes the way the PI-cycle affects specific neurons.<sup>50</sup> The abnormal metabolism of MI has extensive effects on neurologic pathways and changes in PI-cycle activation, which might induce OCD. According to our results, verbal and visual-spatial impairments in OCD patients may be closely related to decreased MI metabolism.

Therefore, this study suggests an extensive and significant impairment of WM in OCD including verbal WM, visuospatial WM and central executive system, which may be closely related to the abnormal metabolism of PFC in patients with OCD and may be an important cause of WM damage.

## Limitations

This study might offer laboratory markers for etiological studies, and perhaps enhance the identification of WM impairment in OCD. However, some potential limitations in the current study should be taken into account when conducting future research. Firstly, due to statistical effect of the results, the sample size might be reduced, and future studies are needed to expand the sample size, allowing us to verify the results again. Secondly, this study is a cross-sectional study. Longitudinal studies are required to validate these findings. Thirdly, all patients had both obsessive thinking and compulsive behavior, and so the subtype analysis could not be carried out. Fourthly, although a battery of common neuropsychological WM tasks was used in our study, it was still not comprehensive enough to address various aspects of WM. In addition, we investigated WM functions without controlling for executive dysfunction and intelligence quotient (IQ) that could affect WM results (eg short-term memory, attention, vigilance etc.). Lastly, OCD is a disease with abnormal brain regions. In this study, only one ROI region was selected using <sup>1</sup>H MRS, and we could not clarify the metabolism of other brain regions.

## Conclusion

In conclusion, these preliminary data suggest that the WM of drug-naive OCD is damaged, involving verbal WM, visual WM and central executive system. Also, our study supported the hypothesis of PFC hypometabolism, especially the IPFC, which showed close relation to the WM impairment, and this might be the pathological mechanism of WM.

## Ethics Committee Approval

The study was approved by the Regional Ethics Committee of the First Affiliated Hospital of Jinan University District and performed in accordance with the

ethical standards laid down in the 1964 Declaration of Helsinki and its later amendments.

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## Disclosure

The authors report no conflicts of interest in this work. None of the authors have any actual or potential conflicts of interest including any financial, personal or other relationships with other people or organizations during the study period.

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