

Changes of serum uric acid and total bilirubin in elderly patients with major postischemic stroke depression

Jie Gao
Wei Xu
Kun Han
Lu Zhu
Lili Gao
Xiuli Shang

Department of Neurology, The First
Affiliated Hospital, China Medical
University, Shenyang, China

Background: This was a longitudinal study which investigated the relationship between serum uric acid (SUA) and total bilirubin (Tbil) upon admission in elderly stroke patients and the occurrence of postischemic stroke depression (IPSD) at 3, 6, and 9 months of post-stroke follow-up.

Subjects and methods: Data were analyzed for 525 acute ischemic stroke patients. Beck Depression Inventory (BDI) scores >17 and *Diagnostic and Statistical Manual of Mental Disorders, Fourth Edition (DSM-IV)* were used separately to screen and diagnose IPSD at 3, 6, and 9 months post-stroke. Once IPSD was diagnosed, follow-up activities were terminated.

Results: High levels of SUA (odds ratio [OR]=2.08, $P<0.01$) and Tbil (OR=2.31, $P<0.01$) in the first 3 months post-stroke and low levels of SUA (OR=2.05, $P=0.03$) and Tbil (OR=2.79, $P<0.01$) from 3 to 6 months post-stroke were identified as risk factors for major IPSD. At 3 months, patients with SUA levels $\geq 406.5 \mu\text{mol/L}$ (males with SUA levels of $\geq 409.5 \mu\text{mol/L}$ and females with SUA levels $\geq 385.5 \mu\text{mol/L}$) and Tbil levels $\geq 23.65 \mu\text{mol/L}$ were more likely to develop major IPSD. At 6 months, both SUA (area under curve [AUC]=0.625, $P=0.005$, cutoff=194.0 $\mu\text{mol/L}$) and Tbil (AUC=0.681, $P=0.004$, cutoff=6.75 $\mu\text{mol/L}$) had minor diagnostic values (AUC <0.700), although SUA levels $\leq 214.5 \mu\text{mol/L}$ (AUC=0.756, $P=0.001$) in female patients had a good diagnostic value (AUC=0.722, $P=0.006$) for major IPSD. At 9 months, major IPSD showed no statistical relationship with either SUA ($\chi^2=2.33$, $P=0.13$) or Tbil ($\chi^2=0.41$, $P=0.84$).

Conclusion: Higher levels of SUA and Tbil on admission were closely related to the occurrence of major IPSD within 3 months of stroke. Lower levels of these two biomarkers on admission were characteristic for the occurrence of major IPSD between 3 and 6 months post-stroke, while 6 months after stroke, there was no relationship between major IPSD and these two biomarkers.

Keywords: depression after stroke, stroke, total bilirubin, uric acid, longitudinal study

Introduction

Postischemic stroke depression (IPSD) is an important post-stroke complication and a chronic disorder that seriously affects the prognosis of patients.^{1,2} The biological processes and pathways of depression generally include inflammatory, oxidative, and nitrosative stress pathways; neurotransmitter systems; neurotrophins; and the regulation of neurogenesis and modulation of the hypothalamic–pituitary–adrenal (HPA) axis.^{3,4} Post-stroke depression (PSD) may have a similar pathological process to depression. Previous research has shown that the levels of some molecular markers are changed in the peripheral system of PSD patients, including C-reactive protein (CRP) and

Correspondence: Xiuli Shang
Department of Neurology,
The First Affiliated Hospital,
China Medical University,
Number 155, Nanjing Street, Heping
District, Shenyang City 110001,
Liaoning Province, China
Email xiulishang_zhao@163.com

homocysteine,^{5–7} brain-derived neurotrophic factor (BDNF),⁸ and triiodothyronine.⁹ Of these pathophysiological factors, oxidative and nitrosative stress pathways are believed to be the most crucial pathophysiological factors,¹⁰ since the brain has high metabolic rates and low antioxidant levels.¹¹

Serum uric acid (SUA) is the ultimate product of purine metabolism and undertakes 60% of the body's antioxidant reactions.¹² Uric acid (UA) can inhibit inflammatory cascades, reduce the permeability of the blood–brain barrier, and protect central nervous tissue.¹³ Similar to the situation reported for cardiovascular disease,¹⁴ hyperuricemia appears to be common in post-stroke patients and is suggestive of a better prognosis.¹⁵ However, low SUA levels after ischemic stroke have also been reported in previous experimental data and were indicative of poor outcomes.¹⁶ UA is also a known scavenger of peroxynitrite, which is the main oxidizing congener of nitric oxide and is involved in the biochemical pathogenesis of depression.¹⁷ However, both high¹⁸ and low^{19,20} SUA levels have been previously shown to be associated with depression.

Bilirubin is a powerful antioxidant and is the final catabolite of the heme fragmentation pathway.²¹ Some studies have shown that decreased serum bilirubin level could be an independent predictor of stroke incidence,^{22,23} and a high serum bilirubin level is related to the severity of stroke and a poor prognosis.²⁴ However, other studies have shown that a high level of serum bilirubin is used as a mechanism which just only reflects the intensity of initial oxidative stress rather than outcome of stroke in acute ischemic stroke.²⁵ The clear inconsistency between these earlier studies is interesting and has led to confusion with regard to our understanding of the specific relationship between bilirubin and stroke. Similar effects have been observed in studies which have evaluated the relationship between bilirubin and depression. For example, Miyaoka et al showed that high levels of biopyrrin (bilirubin oxidative metabolite) in the urine of psychiatric patients were correlated with depressive symptoms.²⁶ In contrast, there is also strong evidence suggesting that low bilirubin levels represent a risk factor for depression.²⁷

The precise relationship between IPSD and levels of SUA and Tbil has yet to be studied in depth, particularly in terms of the occurrence of IPSD at different phases after stroke. To our knowledge, only a limited number of studies have attempted to explain why IPSD may occur over different time periods following stroke.^{28,29} We hypothesized that levels of SUA and Tbil were implicated in IPSD. Consequently, the key purpose of the present study was to investigate the specific relationship between levels of SUA and Tbil at admission and the occurrence of IPSD.

Subjects and methods

Baseline participants and assessment

This was a prospective cohort study undertaken at China Medical University Affiliated Hospital between October 2014 and March 2016 involving 525 patients with ischemic stroke. Data regarding the incidence of postischemic stroke depression and physical outcomes were collected from stroke survivors every 3 months following their initial hospital admission.

Demographic and clinical data

For all subjects, a trained research assistant collected the demographic data (gender, age, marital status, and educational level) and evaluated stroke severity using the National Institute of Health Stroke Scale (NIHSS)³⁰ within 48 hours of admission. The following morning, fasting blood was collected and levels of SUA and Tbil were determined.

Our specific inclusion criteria were as follows: 1) a first occurrence of an ischemic stroke that met the standards of the World Health Organization diagnostic criteria (not exceeding 2 weeks post-stroke); 2) a duration of clinical neurological function deficit lasting over 24 hours; 3) radiological magnetic resonance imaging confirmation³¹; and 4) over 55 years of age. Patients were excluded from the study if: 1) they refused to participate in the study; 2) there was apparent nonvascular etiologies (primary or metastatic neoplasms); 3) they were suffering from pre-stroke disability or depression, or antidepressant history; 4) they had a history of liver disease, hemolytic disease (such as Rh hemolytic disease and thalassemia), serious renal disease, or gout; 5) they were suffering from severe aphasia (language score ≥ 2 using the NIHSS), total blindness, deafness, or disturbance of consciousness; 6) they had a Barthel Index score < 10 ; 7) they had an NIHSS score > 20 ; or 8) their blood indices were insufficient upon admission, such as liver function, renal function, SUA, or Tbil.

SUA and Tbil assay

Levels of SUA and Tbil were determined by Roche kit (Hoffman-La Roche Ltd., Basel, Switzerland) with the use of a Hitachi 7170 automatic biochemical analyzer test (Hitachi High-Technologies, Tokyo, Japan). In our hospital department, normal serum SUA level is known to range from 208 to 428 $\mu\text{mol/L}$ in male patients and from 155 to 357 $\mu\text{mol/L}$ in females, while normal Tbil levels range from 3.0 to 22.0 $\mu\text{mol/L}$. As raw SUA and Tbil data were skewed, we recorded and divided these data into three tertiles: (tertile 1^a ≤ 245.0 $\mu\text{mol/L}$, 245.0 $<$ tertile 2^b < 330.0 $\mu\text{mol/L}$, tertile 3^c ≥ 330.0 $\mu\text{mol/L}$) and (tertile 1^d ≤ 8.0 $\mu\text{mol/L}$, 8.0 tertile 2^e < 12.8 $\mu\text{mol/L}$, tertile 3^f ≥ 12.8 $\mu\text{mol/L}$), respectively.

Follow-up

Patients meeting our specific inclusion criteria were invited to participate in the study and were followed up until 9 months post-stroke. Follow-up measurements were taken at 3, 6, and 9 months post-stroke. All assessments were performed by trained research assistants.

Ethical statement

The study was approved by the Regional Medical Research Ethical Committee of China Medical University. All subjects, or their legal representatives, provided informed written consent.

Assessment of IPSD

Subjects were invited to complete the BDI³² by telephone interview at 3, 6, and 9 months of follow-up as the primary screening standard. Patients with scores >17 were recruited in order to consider a final diagnosis of major IPSD in accordance with the *DSM-IV*³³ by two professional clinicians who were blinded to laboratory results.

Statistical analysis

The SPSS 19.0 statistical package (SPSS Inc., Chicago, IL, USA) was used to analyze all data arising from the study. A χ^2 test was used to assess differences in categorical variables data. For continuous, normally distributed data, the Student's *t*-test was used. Risk factors ($P \leq 0.10$) were further analyzed by univariate and multivariate logistic regression. Relationships between gender and the levels of SUA and Tbil were analyzed using Spearman correlation analysis. Receiver operating characteristic curves were drawn to depict the diagnostic performance of SUA and Tbil levels in the occurrence of IPSD. All *P*-values were calculated as two-tailed, and significance level was set at $P < 0.05$.

Results

Patients

A total of 1,006 ischemic stroke patients were screened and 760 met the specific entry criteria. During the follow-up process, 235 patients withdrew; consequently, the final data analysis included 525 patients. The follow-up procedure is depicted in Figure 1.

Patients who withdrew from the study had a significantly higher mean age (mean $\pm$ SD: 68.26 $\pm$ 7.48 years vs 64.00 $\pm$ 13.19 years; $t=5.28$, $P<0.01$), and were more likely to have less than 9 years of formal education (46.6% vs 37.3%, $P=0.03$). Baseline characteristics of patients who completed the follow-up period are shown in Table 1.

Correlation analysis between the levels of SUA and Tbil upon admission and the occurrence of IPSD

When no confounding factors were considered, a significant difference was observed in terms of Tbil levels when compared between patients with major IPSD and non-major IPSD ($\chi^2=10.89$, $P<0.01$; Table 1). Next, we entered all risk factors with $P \leq 0.10$ (Table 1) into a multivariate logistic regression model (ie, female gender, age, Barthel Index [BI], NIHSS, frontal lobe, thalamus, and Tbil level [tertile 3^o]). Of these risk factors, multivariate logistic regression confirmed that Tbil levels (tertile 3^o) were independently related to the occurrence of IPSD with an odds ratio (OR) of 1.93 ($P<0.01$) (Table 2).

The relationship between the occurrence of major IPSD and the levels of serum SUA and Tbil after adjustment for other risk factors

There were 105 patients with major IPSD within 3 months of stroke, 48 patients from 3 to 6 months post-stroke, and 34 patients between 6 and 9 months post-stroke. The incidence of major IPSD was significantly different when compared between these different time periods ($\chi^2=24.96$, $P<0.01$). Therefore, the analysis of relationships between different levels of SUA and Tbil and major IPSD was stratified according to different time periods following stroke (3 months, 3–6 months, and 6–9 months post-stroke, as shown in Table 3).

When we compared patients with major IPSD and non-major IPSD across the different time periods in terms of gender, age, education, hypertension, diabetes, NIHSS, BI, location, 17-item Hamilton Depression Rating Scale, blood urea nitrogen, creatinine, alkaline phosphatase, and alanine aminotransferase, we identified several risk factors where $P \leq 0.1$: age ($t=-1.79$, $P=0.07$), thalamus ($\chi^2=5.47$, $P=0.02$), frontal lobe ($\chi^2=4.57$, $P=0.03$), and NIHSS ($t=-1.86$, $P=0.07$) in the first 3 months post-stroke; female gender ($\chi^2=5.81$, $P=0.02$), age ($t=-1.68$, $P=0.10$), and NIHSS score ($t=-2.78$, $P=0.01$) from 3 to 6 months post-stroke; female gender ($\chi^2=4.53$, $P=0.03$), age ($t=-2.43$, $P=0.02$), and living alone ($\chi^2=6.13$, $P=0.01$) from 6 to 9 months post-stroke. Next, we used univariate and multivariate analyses to investigate major IPSD, SUA, and Tbil levels and the significant risk factors identified earlier (Table 2). The results of these analyses are shown in Table 4.

The sensitivity and specificity of SUA and Tbil levels to predict major IPSD in the first 6 months post-stroke

As shown in Table 5, a negative correlation was found between female gender and SUA level ($r=-0.30$, $P<0.01$). The predictive

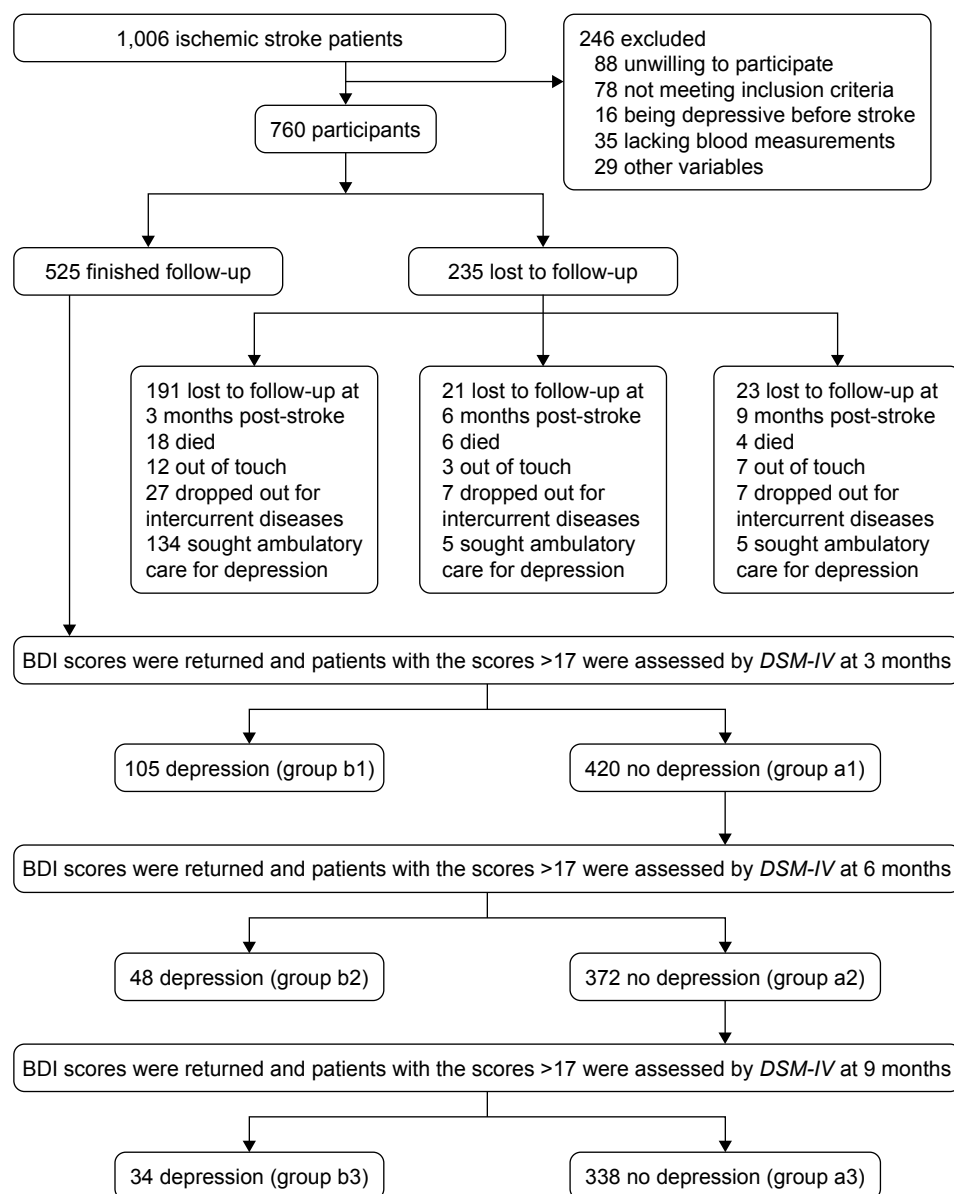


Figure 1 The follow-up chart.

Abbreviations: BDI, Beck Depression Inventory; DSM-IV, *Diagnostic and Statistical Manual of Mental Disorders, Fourth Edition*.

values of SUA and Tbil levels are shown in Table 6. In the first 3 months post-stroke, SUA levels $\geq 330.0 \mu\text{mol/L}$ (area under curve [AUC]=0.834, $P<0.001$, Figure 2A) and Tbil levels $\geq 12.8 \mu\text{mol/L}$ (AUC=0.782, $P<0.001$, Figure 2B) showed significant diagnostic value. Furthermore, patients with SUA levels $\geq 406.5 \mu\text{mol/L}$, or Tbil levels $\geq 23.65 \mu\text{mol/L}$, were more likely to be depressive. Meanwhile, male patients with SUA levels $\geq 409.5 \mu\text{mol/L}$ (AUC=0.920, $P<0.001$, Figure 2C) and females with SUA levels $\geq 385.5 \mu\text{mol/L}$ (AUC=0.756, $P=0.001$, Figure 2D) had a greater risk of suffering from major IPSP.

From 3 to 6 months post-stroke, the diagnostic performance of SUA levels $\leq 245 \mu\text{mol/L}$ (AUC=0.676,

$P=0.007$, Figure 3A) and Tbil levels $\leq 8.0 \mu\text{mol/L}$ (AUC=0.681, $P=0.004$, Figure 3B) was statistically significant, although the diagnostic value was not very high. However, compared with male patients (AUC=0.580, $P=0.465$, Figure 3C), female patients with SUA levels $\leq 214.5 \mu\text{mol/L}$ (AUC=0.722, $P=0.006$, Figure 3D) were more likely to develop major IPSP.

Discussion

Unlike previous studies, the present investigation focused on the occurrence of major IPSP across different time periods and attempted to link this information with levels of Tbil and SUA. The common characteristic of SUA and Tbil is

Table 1 Baseline subject characteristics with hemispheric lesion verified by radiologic magnetic resonance imaging

Variables	No IPSD n=338 Mean $\pm$ SD or n (%)	IPSD n=187 Mean $\pm$ SD or n (%)	t/ χ^2	P-value
Female	181 (53.6)	115 (61.5)	3.09	0.08
Age	65.4 $\pm$ 10.9	67.4 $\pm$ 11.4	-2.00	0.05
<9 years of education	130 (38.5)	67 (35.8)	0.36	0.57
Living alone	40 (11.8)	32 (16.0)	1.85	0.17
History of hypertension	224 (66.3)	130 (69.5)	0.58	0.45
History of diabetes	114 (33.7)	67 (35.8)	0.24	0.63
Lesion location				
Frontal lobe	14 (4.1)	14 (7.5)	2.67	0.10
Parietal lobe	40 (11.8)	18 (9.6)	0.60	0.44
Temporal lobe	18 (5.3)	11 (5.9)	0.07	0.79
Occipital lobe	22 (6.5)	14 (7.5)	0.18	0.67
Basal ganglia	103 (30.5)	68 (36.4)	1.90	0.17
Thalamus	19 (5.6)	26 (13.9)	10.54	<0.01
Brainstem	34 (10.1)	18 (9.6)	0.03	0.87
Cerebellum	22 (6.5)	9 (4.8)	0.62	0.43
Other location	18 (5.3)	10 (5.3)	0.00	0.99
$\geq$ Two locations	31 (9.2)	11 (5.9)	1.78	0.18
BI	71.48 $\pm$ 16.11	68.90 $\pm$ 17.34	1.71	0.09
NIHSS score	4.91 $\pm$ 2.54	5.35 $\pm$ 2.55	-1.91	0.06
BUN (mmol/L)	5.17 $\pm$ 1.56	5.23 $\pm$ 1.60	-0.43	0.67
Cr (μ mol/L)	67.15 $\pm$ 17.96	65.97 $\pm$ 18.91	0.70	0.48
ALT (U/L)	21.77 $\pm$ 9.28	20.42 $\pm$ 9.13	1.61	0.11
ALP (U/L)	71.30 $\pm$ 17.36	70.10 $\pm$ 18.94	0.74	0.46
HAMD17	2.95 $\pm$ 2.03	3.04 $\pm$ 2.00	-0.45	0.66
Uric acid (μ mol/L)			1.52	0.47
$\leq$ 245.0	108 (32.0)	69 (36.9)	1.32	0.25
245.0–330.0	117 (34.6)	57 (30.5)	0.93	0.34
$\geq$ 330.0	113 (33.4)	61 (32.6)	0.04	0.85
Total bilirubin (μ mol/L)			10.89	<0.01
$\leq$ 8.0	119 (35.2)	62 (33.2)	0.22	0.64
8.0–12.8	124 (36.7)	48 (25.7)	6.64	0.01
$\geq$ 12.8	95 (28.1)	77 (41.2)	9.34	<0.01

Abbreviations: ALP, alkaline phosphatase; ALT, alanine transaminase; BI, Barthel Index; BUN, blood urea nitrogen; Cr, creatinine; HAMD17, 17-item Hamilton Depression Rating Scale; IPSD, postischemic stroke depression; NIHSS, National Institute of Health Stroke Scale.

that both these factors exhibit antioxidant reactions within the human body. Our study also confirmed that the changes observed on the levels of these two biomarkers showed excellent consistency in major IPSD patients. While the precise mechanism underlying IPSD has yet to be established, our

present study provided significant evidence of the important pathological roles of both SUA and Tbil in IPSD.

Patients developing major IPSD in the first 3 months after stroke also showed higher levels of SUA and Tbil upon admission. In a previous study, Nanetti et al showed

Table 2 Univariate and multiple logistic regression analyses with risk factors ($P \leq 0.1$) of postischemic stroke depression

Variables	Univariate correlation analysis		Multivariate correlation analysis	
	OR (95% CI)	P-value	OR (95% CI)	P-value
Female	1.39 (0.96–1.99)	0.08	1.37 (0.94–2.00)	0.10
Age	1.02 (1.00–1.03)	0.05	1.01 (1.00–1.03)	0.23
Frontal lobe	1.87 (0.87–4.02)	0.11	2.08 (0.94–4.60)	0.07
Thalamus	2.71 (1.46–5.05)	<0.01	2.63 (1.93–4.98)	<0.01
NIHSS	1.07 (1.00–1.15)	0.06	1.08 (1.00–1.16)	0.05
BI	0.99 (0.98–1.00)	0.09	0.99 (0.98–1.00)	0.19
Total bilirubin $\geq$ 12.8 μ mol/L	1.79 (1.23–2.61)	<0.01	1.93 (1.31–2.85)	<0.01

Abbreviations: BI, Barthel Index; NIHSS, National Institute of Health Stroke Scale; OR, odds ratio.

Table 3 The relationship between serum uric acid and total bilirubin on admission and postischemic stroke depression (IPSD) in different time periods

No IPSD	Patients, n (%)	IPSD	Patients, n (%)	χ^2	P-value
Uric acid					
Group a1 (n=420)		Group b1 (n=105)			
Tertile 1 ^a	148 (35.2)	Tertile 1 ^a	29 (27.6)	2.18	0.14
Tertile 2 ^b	147 (35.0)	Tertile 2 ^b	27 (25.7)	3.27	0.07
Tertile 3 ^c	125 (29.8)	Tertile 3 ^c	49 (46.7)	10.83	<0.01
Group a2 (n=372)		Group b2 (n=48)			
Tertile 1 ^a	124 (33.3)	Tertile 1 ^a	24 (50)	5.18	0.02
Tertile 2 ^b	128 (34.4)	Tertile 2 ^b	19 (39.6)	0.50	0.48
Tertile 3 ^c	120 (32.3)	Tertile 3 ^c	5 (10.4)	9.70	<0.01
Group a3 (n=338)		Group b3 (n=34)			
Tertile 1 ^a	108 (32.0)	Tertile 1 ^a	16 (47.1)	3.17	0.08
Tertile 2 ^b	117 (34.6)	Tertile 2 ^b	11 (32.4)	0.07	0.79
Tertile 3 ^c	113 (33.4)	Tertile 3 ^c	7 (20.6)	2.23	0.13
Total bilirubin					
Group a1 (n=420)		Group b1 (n=105)			
Tertile 1 ^d	161 (38.3)	Tertile 1 ^d	20 (19.0)	13.83	<0.01
Tertile 2 ^e	138 (32.9)	Tertile 2 ^e	34 (32.4)	0.01	0.93
Tertile 3 ^f	121 (28.8)	Tertile 3 ^f	51 (48.6)	14.89	<0.01
Group a2 (n=372)		Group b2 (n=48)			
Tertile 1 ^d	136 (36.6)	Tertile 1 ^d	25 (52.1)	4.33	0.04
Tertile 2 ^e	132 (35.5)	Tertile 2 ^e	6 (12.5)	10.18	<0.01
Tertile 3 ^f	104 (28.0)	Tertile 3 ^f	17 (35.4)	1.15	0.28
Group a3 (n=338)		Group b3 (n=34)			
Tertile 1 ^d	119 (35.2)	Tertile 1 ^d	17 (50.0)	2.92	0.09
Tertile 2 ^e	124 (36.7)	Tertile 2 ^e	8 (23.5)	2.34	0.13
Tertile 3 ^f	95 (28.1)	Tertile 3 ^f	9 (26.5)	0.04	0.84

Notes: Tertile 1^a ≤ 245.0 $\mu\text{mol/L}$, $245.0 < \text{tertile } 2^b \leq 330.0$ $\mu\text{mol/L}$, tertile 3^c ≥ 330.0 $\mu\text{mol/L}$; tertile 1^d ≤ 8.0 $\mu\text{mol/L}$, $8.0 < \text{tertile } 2^e \leq 12.8$ $\mu\text{mol/L}$, tertile 3^f ≥ 12.8 $\mu\text{mol/L}$. Group a1, no IPSD at 3 months; Group b1, IPSD at 3 months; Group a2, no IPSD at 6 months; Group b2, IPSD at 6 months; Group a3, no IPSD at 9 months; Group b3, IPSD at 9 months.

Table 4 Univariate and multiple logistic regression analyses with risk factors ($P \leq 0.1$) of postischemic stroke depression (IPSD) in different time periods

Occurring time of IPSD	Variables	Univariate correlation analysis		Multivariate correlation analysis	
		OR (95% CI)	P-value	OR (95% CI)	P-value
Three months post-stroke	Age	1.02 (1.00–1.04)	0.07	1.02 (0.99–1.04)	0.11
	Frontal lobe	2.35 (1.05–5.26)	0.04	2.07 (0.88–4.87)	0.10
	Thalamus	2.17 (1.12–4.20)	0.02	2.20 (1.11–4.37)	0.03
	NIHSS	1.09 (1.01–1.19)	0.04	1.07 (0.98–1.16)	0.15
	Uric acid ≥ 330.0 $\mu\text{mol/L}$	2.07 (1.33–3.20)	<0.01	2.08 (1.33–3.27)	<0.01
	Total bilirubin ≥ 12.8 $\mu\text{mol/L}$	2.33 (1.51–3.61)	<0.01	2.31 (1.47–3.63)	<0.01
Six months post-stroke	Female	2.20 (1.15–4.24)	0.02	1.83 (0.93–3.60)	0.08
	Age	1.03 (1.00–1.06)	0.10	1.03 (0.99–1.07)	0.07
	NIHSS	1.20 (1.05–1.38)	0.01	1.27 (1.10–1.46)	<0.01
	Uric acid ≤ 245.0 $\mu\text{mol/L}$	2.00 (1.09–3.66)	0.03	2.05 (1.09–3.86)	0.03
	Total bilirubin ≤ 8.0 $\mu\text{mol/L}$	1.89 (1.03–3.45)	0.04	2.79 (1.44–5.40)	<0.01
Nine months post-stroke	Female	2.26 (1.05–4.88)	0.04	2.04 (0.93–4.51)	0.08
	Age	1.05 (1.01–1.09)	0.02	1.05 (1.01–1.09)	0.02
	Living alone	2.76 (1.20–6.34)	0.02	3.01 (1.26–7.17)	0.01
	Uric acid ≤ 245.0 $\mu\text{mol/L}$	1.89 (0.93–3.86)	0.08	2.06 (0.98–4.33)	0.06
	Total bilirubin ≥ 12.8 $\mu\text{mol/L}$	1.84 (0.91–3.74)	0.09	1.72 (0.83–3.57)	0.15

Abbreviations: NIHSS, National Institute of Health Stroke Scale; OR, odds ratio.

Table 5 Relationship between sex and serum uric acid and total bilirubin levels

	Tertiles	Patients (n)	Female (n, %)	χ^2, P	Relevance, P
Uric acid ($\mu\text{mol/L}$)	≤ 245.0	177	131 (74.0)	$\chi^2=46.47, P<0.01$	$r=-0.30, P<0.01$
	245.0–330.0	174	99 (56.9)		
	≥ 330.0	174	66 (37.9)		
Total bilirubin ($\mu\text{mol/L}$)	≤ 8.0	180	99 (55.0)	$\chi^2=0.70, P=0.71$	$r=0.03, P=0.46$
	8.0–12.8	172	95 (55.2)		
	≥ 12.8	173	102 (59.0)		

that acute ischemic stroke could cause the generation of free radicals and strong oxidative stress, and that this antioxidant capacity may limit the progression of ischemic injury.³⁴ As powerful antioxidants, high concentrations of SUA and Tbil could reflect the level of oxidative stress in the first 3 months after stroke. Meanwhile, two previous meta-analyses reported that depression was strongly correlated to high oxidative stress by demonstrating a clear imbalance between antioxidants and oxidant activity.³⁵ Previous research also implied that the pathophysiology of IPSD is probably involved in increased levels of oxidative stress in cerebral tissues.³⁶ Firstly, oxidative stress, which involves enzymatic and nonenzymatic antioxidant factors expressed in the brain and periphery, could cause damage to proteins, DNA molecules, and lipids.^{37,38} Both SUA and Tbil are nonenzymatic antioxidants, much like glutathione and vitamins A, C, and E.^{12,37} Secondly, oxidative stress has been implicated in IPSD via the regulation of inflammation.³⁵ A series of studies also provided support for the role of inflammatory factors in the development of IPSD.^{39–41} Under the influence of repeated inflammatory stimuli, microglia in the central nervous system can evolve into a source of inflammatory mediators, which may then influence brain neurotransmitter systems and neuronal integrity.⁴² Moreover, elevated levels of inflammatory cytokines can also affect the activity of 2,3-dioxygenase, which degrades tryptophan via the kynurenine

pathway and ultimately leads to a reduction in the synthesis and availability of 5-hydroxytryptamine.^{43,44} It has also been reported that interaction between oxidative stress pathways and inflammatory pathways may lead to an increased level of neuronal apoptosis and neurodegeneration and a decline in the neurogenesis and neuroplasticity of depression.^{4,45,46} Thus, we can conclude that close associations between major IPSD and higher levels of SUA and Tbil in the first 3 months after stroke may be associated with high levels of oxidative stress. It has been reported that 60% of patients with depression during this particular time period showed recovery by 12 months post-stroke.⁴⁷ It is possible that the symptoms of depression disappear at the same time as the disappearance of oxidative stress response.

However, this does not explain why patients with lower levels of SUA and Tbil at admission are more vulnerable to depression between 3 and 6 months post-stroke. One previous meta-analysis suggested that high levels of SUA at the onset of ischemic stroke could reduce levels of neurological damage and herald a better prognosis.¹⁵ Similar results were also reported by Perlstein et al for higher levels of Tbil.⁴⁸ Some studies have also reported reduced levels of antioxidants in some patients with acute ischemic stroke.⁴⁹ Thus, we can infer that lower levels of SUA and Tbil are related to more serious neurological damage than higher levels in acute stroke. It has also been reported that the more significant the

Table 6 Diagnostic performance of serum uric acid (SUA) and total bilirubin (Tbil) for the assessment of depression occurring in first 6 months post-stroke

Items	Cutoff	Specificity	Sensitivity	YI	AUC (95% CI)	P-value
Three months post-stroke						
SUA	406.5	0.760	0.816	0.576	0.834 (0.765–0.902)	<0.001
Male	409.5	0.765	1.000	0.765	0.920 (0.870–0.969)	<0.001
Female	385.5	0.739	0.762	0.501	0.756 (0.619–0.893)	0.001
Tbil	23.65	0.926	0.569	0.495	0.782 (0.704–0.860)	<0.001
Six months post-stroke						
SUA	194.0	0.694	0.750	0.444	0.676 (0.579–0.773)	0.007
Male	–	–	–	–	0.580 (0.401–0.760)	0.465
Female	214.5	0.515	0.938	0.501	0.722 (0.606–0.837)	0.006
Tbil	6.75	0.331	1.000	0.331	0.681 (0.580–0.783)	0.004

Note: YI = specificity + sensitivity – 1.

Abbreviations: AUC, area under the curve; YI, Youden Index.

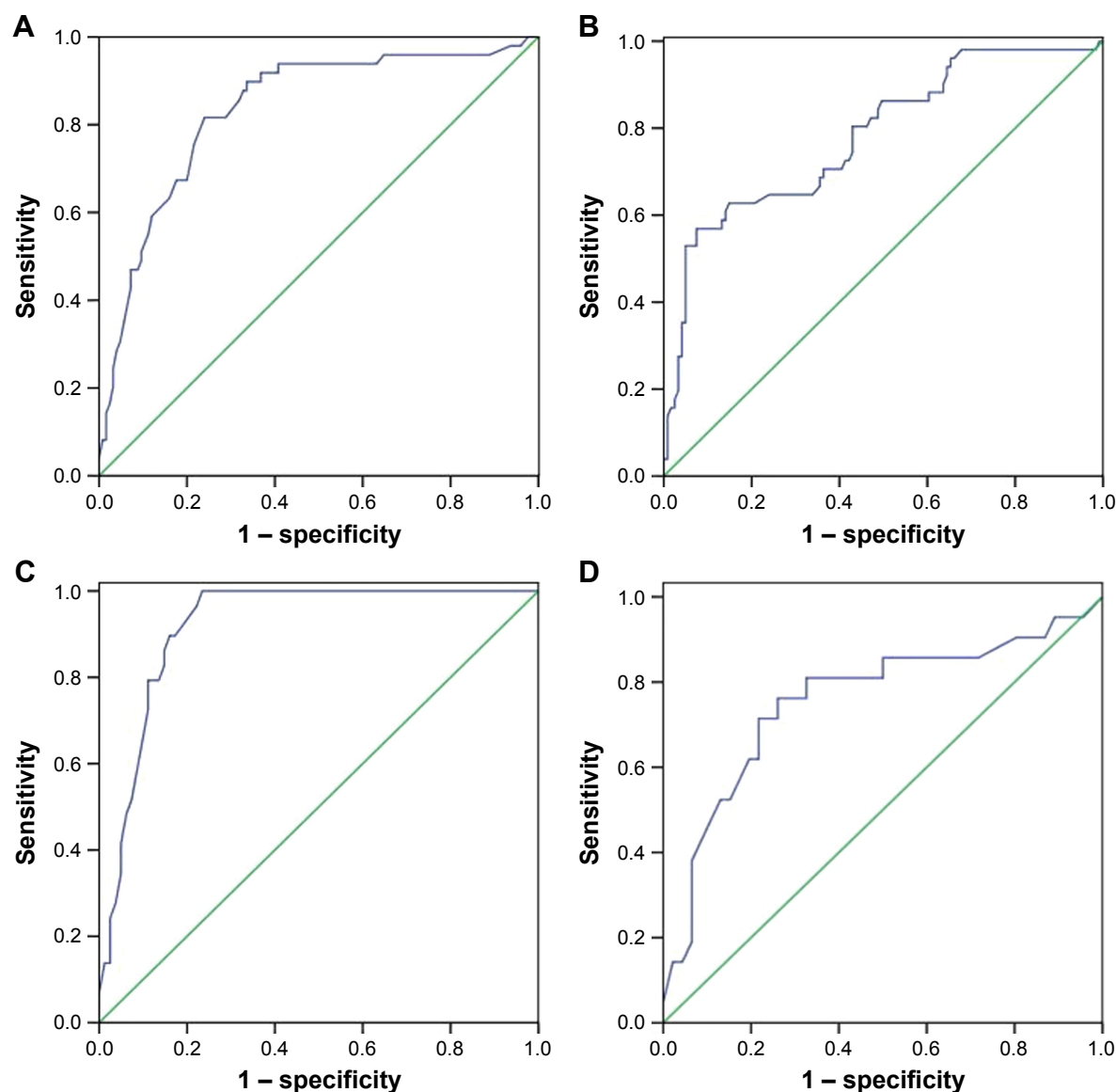


Figure 2 ROC curve analysis for SUA and Tbil used for the assessment of IPSD in the first 3 months post-stroke. (A) SUA levels, (B) Tbil levels, (C) SUA levels in males, and (D) SUA levels in females.

Abbreviations: IPSD, postischemic stroke depression; ROC, receiver operating characteristic; SUA, serum uric acid; Tbil, total bilirubin.

stroke severity and functional impairment are, the greater the probability of suffering from depression.⁵⁰ This is consistent with the higher NIHSS score in major IPSD patients observed between 3 and 6 months post-stroke. However, it is still unclear as to why patients with low levels of SUA and Tbil in our study developed depression after a 3-month delay instead of acutely. There may be several possible explanations for this phenomenon. Firstly, once the acute phase is over, patients with more severe ischemic stroke may gradually realize the seriousness of problems related to stroke,⁵¹ such as the decline in self-care and work ability, which would make such patients feel incompetent. Secondly, in patients with significant functional impairment such as

post-stroke paresthesia, early paresthesia may lead to sleep disorders, and long-term sleep disorders inevitably lead to the occurrence of depression.⁵²

From 6 to 9 months, we did not observe any significant association between major IPSD and the levels of SUA and Tbil. As we described in our previous publication, since biological changes and reconstruction in the brain tissue cease during this time frame, social psychology factors, rather than stroke severity, begin to play a more important role.⁵³

In summary, our current study provides evidence that different levels of SUA and Tbil in the blood may be helpful in the early prevention of IPSD. This information may also be useful in selecting treatments for depression considering the

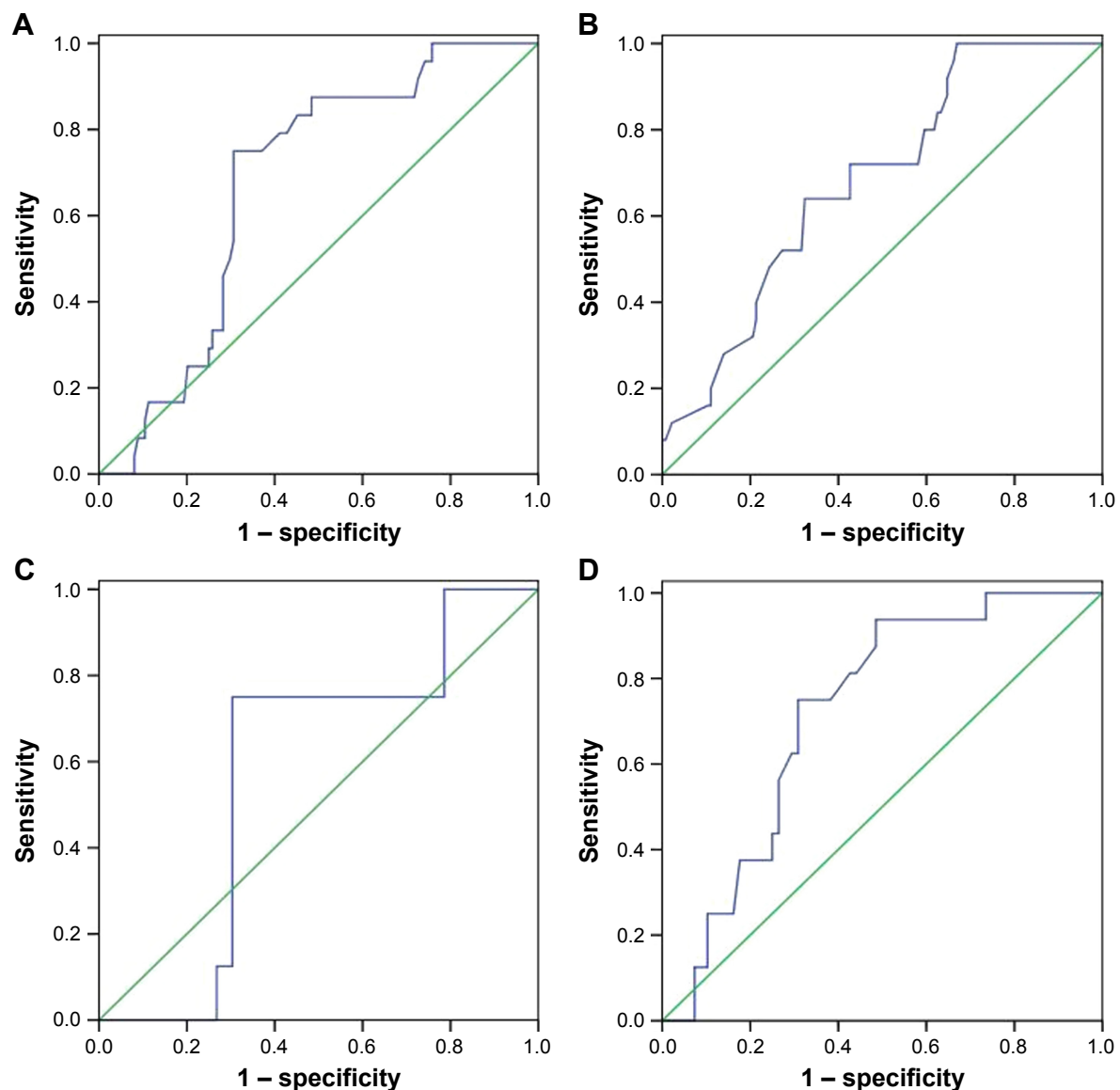


Figure 3 ROC curve analysis for SUA and Tbil used for the assessment of IPSD from 3 to 6 months post-stroke. (A) SUA levels, (B) Tbil levels, (C) SUA levels in males, and (D) SUA levels in females.

Abbreviations: IPSD, postischemic stroke depression; ROC, receiver operating characteristic; SUA, serum uric acid; Tbil, total bilirubin.

different mechanisms of depression which occur over different time periods. In the first 3 months after stroke, depression may not need excessive intervention because acute oxidative stress could disappear by itself. However, from 3 to 6 months, early post-stroke rehabilitation treatment has a more significant effect upon the prevention of depression and recovery because of the relationship between depression and stroke severity. Six months after stroke, antidepressive therapies, such as antidepressant drugs or psychotherapy, may begin to exert effects upon the treatments used for major IPSD. Compared with other biomarkers, there are several superiorities of the two biomarkers (SUA and Tbil) for IPSD. First, SUA and Tbil represent affordable routine clinical tools which do not incur

additional expense as compared with BDNF, which is limited to precise laboratory tests and not suitable for extensive clinical studies and follow-up. Second, as compared with CRP, SUA and Tbil are less affected by the following factors, such as fever, infection, immune system diseases, and so on. Third, as compared with triiodothyronine and other biomarkers, SUA and Tbil may be used to monitor the development of IPSD, since a previous report showed that the level of SUA changes as depression improves.¹¹

Acknowledgment

This work was supported by grants from the Natural Science Foundation of China (81100243 and 81502181).

Disclosure

The authors report no conflicts of interest in this work.

References

- Taylor-Piliae RE, Hepworth JT, Coull BM. Predictors of depressive symptoms among community-dwelling stroke survivors. *J Cardiovasc Nurs*. 2013;28(5):460–467.
- Babkair LA. Risk factors for poststroke depression: an integrative review. *J Cardiovasc Nurs*. 2017;49(2):73–84.
- Moylan S, Maes M, Wray NR, Berk M. The neuroprogressive nature of major depressive disorder: pathways to disease evolution and resistance, and therapeutic implications. *Mol Psychiatry*. May;18(5):595–606.
- Leonard B, Maes M. Mechanistic explanations how cell-mediated immune activation, inflammation and oxidative and nitrosative stress pathways and their sequels and concomitants play a role in the pathophysiology of unipolar depression. *Neurosci Biobehav Rev*. 2012;36(2):764–785.
- Tang CZ, Zhang YL, Wang WS, Li WG, Shi JP. Serum levels of high-sensitivity C-reactive protein at admission are more strongly associated with poststroke depression in acute ischemic stroke than homocysteine levels. *Mol Neurobiol*. 2016;53(4):2152–2160.
- Qian L, Ding L, Cheng L, et al. Early biomarkers for post-stroke cognitive impairment. *J Neurol*. Oct;259(10):2111–2118.
- Narasimhalu K, Lee J, Leong YL, et al. Inflammatory markers and their association with post stroke cognitive decline. *Int J Stroke*. 2015;10(4):513–518.
- Fang J, Cheng Q. Etiological mechanisms of post-stroke depression: a review. *Neurol Res*. 2009;31(9):904–909.
- Zhao J, Ren W, Lv D, Zhu Z, Wang Q, He J. Low triiodothyronine syndrome is a predictor of post-stroke depression. *Int J Geriatr Psychiatry*. 2017;32(3):352–353.
- Maes M. The cytokine hypothesis of depression: inflammation, oxidative & nitrosative stress (IO&NS) and leaky gut as new targets for adjunctive treatments in depression. *Neuro Endocrinol Lett*. 2008;29(3):287–291.
- Wen S, Cheng M, Wang H, et al. Serum uric acid levels and the clinical characteristics of depression. *Clin Biochem*. 2012;45(1):49–53.
- Ames BN, Cathcart R, Schwiers E, Hochstein P. Uric acid provides an antioxidant defense in humans against oxidant- and radical-caused aging and cancer: a hypothesis. *Proc Natl Acad Sci U S A*. 1981;78(11):6858–6862.
- Lucca G, Comim CM, Valvassori SS, et al. Increased oxidative stress in submitochondrial particles into the brain of rats submitted to the chronic mild stress paradigm. *J Psychiatr Res*. 2009;43(9):864–869.
- Wu CY, Hu HY, Chou YJ, et al. High serum uric acid levels are associated with all-cause and cardiovascular, but not cancer, mortality in elderly adults. *J Am Geriatr Soc*. 2015;63(9):1829–1836.
- Wang Z, Lin Y, Liu Y, et al. Serum uric acid levels and outcomes after acute ischemic stroke. *Mol Neurobiol*. 2016;53(3):1753–1759.
- Cherubini A, Polidori MC, Bregnocchi M, et al. Antioxidant profile and early outcome in stroke patients. *Stroke*. 2000;31(10):2295–2300.
- Kaster MP, Rosa AO, Santos AR, Rodrigues AL. Involvement of nitric oxide-cGMP pathway in the antidepressant-like effects of adenosine in the forced swimming test. *Int J Neuropsychopharmacol*. 2005;8(4):601–606.
- Tao R, Li H. High serum uric acid level in adolescent depressive patients. *J Affect Disord*. 2015;174:464–466.
- Wen S, Cheng M, Wang H, et al. Serum uric acid levels and the clinical characteristics of depression. *Clin Biochem*. 2012;45(1–2):49–53.
- Chaudhari K, Khanzode S, Khanzode S, Dakhale G, Saoji A, Sarode S. Clinical correlation of alteration of endogenous antioxidant-uric acid level in major depressive disorder. *Indian J Clin Biochem*. 2010;25(1):77–81.
- Stocker R, Yamamoto Y, McDonagh A, Glazer A, Ames B. Bilirubin is an antioxidant of possible physiological importance. *Science*. 1987;235(4792):1043–1046.
- Kimm H, Yun JE, Jo J, Jee SH. Low serum bilirubin level as an independent predictor of stroke incidence a prospective study in Korean men and women. *Stroke*. 2009;40(11):3422–3427.
- Targher G. Risk of ischemic stroke and decreased serum bilirubin levels: is there a causal link? *Arterioscler Thromb Vasc Biol*. 2014;34(4):702–704.
- Luo Y, Li J, Zhang J, Xu Y. Elevated bilirubin after acute ischemic stroke linked to the stroke severity. *Int J Dev Neurosci*. 2013;31(7):634–638.
- Pineda S, Bang OY, Saver JL, et al. Association of serum bilirubin with ischemic stroke outcomes. *J Stroke Cerebrovasc Dis*. 2008;17(3):147–152.
- Miyaoka T, Yasukawa R, Yasuda H, et al. Urinary excretion of biopyrrins, oxidative metabolites of bilirubin, increases in patients with psychiatric disorders. *Eur Neuropsychopharmacol*. 2005;15(3):249–252.
- Gordon WA, Hibbard MR. Poststroke depression: an examination of the literature. *Arch Phys Med Rehabil*. 1997;78(6):658–663.
- Robinson RG, Lipsey JR, Rao K, Price TR. Two-year longitudinal study of post-stroke mood disorders: comparison of acute-onset with delayed-onset depression. *Am J Psychiatry*. 1986;143(10):1238–1244.
- Kim JT, Park MS, Yoon GJ, et al. White matter hyperintensity as a factor associated with delayed mood disorders in patients with acute ischemic stroke. *Eur Neurol*. 2011;66(6):343–349.
- Brott T, Adams HP Jr, Olinger CP, et al. Measurements of acute cerebral infarction: a clinical examination scale. *Stroke*. 1989;20(7):864–870.
- Kunitz S, Gross C, Heyman A, et al. The pilot Stroke Data Bank: definition, design, and data. *Stroke*. 1984;15(4):740–746.
- Beck AT, Steer R, Brown G. *Beck Depression Inventory Manual [M]*. Psychological Corporation: San Antonio, TX; 1996.
- American Psychiatric Association. *Diagnostic and Statistical Manual of Mental Disorders*. 4th ed. Washington, DC: American Psychiatric Association; 1994.
- Nanetti L, Raffaelli F, Vignini A, et al. Oxidative stress in ischaemic stroke. *Eur J Clin Invest*. 2011;41(12):1318–1322.
- Palta P, Samuel LJ, Miller ER 3rd, Szanton SL. Depression and oxidative stress: results from a meta-analysis of observational studies. *Psychosom Med*. 2014;76(1):12–19.
- Nabavi SF, Dean OM, Turner A, Sureda A, Daglia M, Nabavi SM. Oxidative stress and post-stroke depression: possible therapeutic role of polyphenols? *Curr Med Chem*. 2015;22(3):343–351.
- Valko M, Leibfritz D, Moncol J, Cronin MT, Mazur M, Telser J. Free radicals and antioxidants in normal physiological functions and human disease. *Int J Biochem Cell Biol*. 2007;39(1):44–84.
- Girotti AW. Lipid hydroperoxide generation, turnover, and effector action in biological systems. *J Lipid Res*. 1998;39(8):1529–1542.
- Rawdin BJ, Mellon SH, Dhabhar FS, et al. Dysregulated relationship of inflammation and oxidative stress in major depression. *Brain Behav Immun*. 2013;31:143–152.
- Galecki P. Peripheral markers of inflammation, oxidative and nitrosative stress pathways and memory functions as a new target of pharmacotherapy in depression. *Prog Neuropsychopharmacol Biol Psychiatry*. 2018;80(Pt C):167.
- Doolin K, Farrell C, Tozzi L, Harkin A, Frodl T, O'Keane V. Diurnal hypothalamic-pituitary-adrenal axis measures and inflammatory marker correlates in major depressive disorder. *Int J Mol Sci*. 2017;18(10):pii: E2226.
- Chen Z, Jalabi W, Shpargel KB, et al. Lipopolysaccharide-induced microglial activation and neuroprotection against experimental brain injury is independent of hematogenous TLR4. *J Neurosci*. 2012;32(34):11706–11715.
- Li W, Ling S, Yang Y, Hu Z, Davies H, Fang M. Systematic hypothesis for post-stroke depression caused inflammation and neurotransmission and resultant on possible treatments. *Neuro Endocrinol Lett*. 2014;35(2):104–109.

44. Fujigaki H, Saito K, Fujigaki S, et al. The signal transducer and activator of transcription 1alpha and interferon regulatory factor 1 are not essential for the induction of indoleamine 2,3-dioxygenase by lipopolysaccharide: involvement of p38 mitogen-activated protein kinase and nuclear factor-kappaB pathways, and synergistic effect of several proinflammatory cytokines. *J Biochem.* 2006;139(4):655–662.
45. Maes M, Mihaylova I, Kubera M, Uytterhoeven M, Vrydags N, Bosmans E. Lower whole blood glutathione peroxidase (GPX) activity in depression, but not in myalgic encephalomyelitis/chronic fatigue syndrome: another pathway that may be associated with coronary artery disease and neuroprogression in depression. *Neuro Endocrinol Lett.* 2011;32(2):133–140.
46. Maes M, Galecki P, Chang YS, Berk M. A review on the oxidative and nitrosative stress (O&NS) pathways in major depression and their possible contribution to the (neuro)degenerative processes in that illness. *Prog Neuropsychopharmacol Biol Psychiatry.* 2011;35(3):676–692.
47. Astrom M, Adolfsson R, Asplund K. Major depression in stroke patients: a 3-year longitudinal study. *Stroke.* 1993;24(7):976–982.
48. Perlstein TS, Pande RL, Creager MA, Weuve J, Beckman JA. Serum total bilirubin level, prevalent stroke, and stroke outcomes: NHANES 1999–2004. *Am J Med.* 2008;121(9):781–788 e781.
49. Cojocaru IM, Cojocaru M, Sapira V, Ionescu A. Evaluation of oxidative stress in patients with acute ischemic stroke. *Rom J Intern Med.* 2013; 51(2):97–106.
50. Ferro JM, Caeiro L, Figueira ML. Neuropsychiatric sequelae of stroke. *Nat Rev Neurol.* 2016;12(5):269–280.
51. Berg A, Palomaki H, Lehtihalmes M, Lonnqvist J, Kaste M. Poststroke depression: an 18-month follow-up. *Stroke.* 2003;34(1):138–143.
52. Vandeputte M, de Weerd A. Sleep disorders and depressive feelings: a global survey with the Beck Depression Scale. *Sleep Med.* 2003;4(4):343–345.
53. Gao J, Lin M, Zhao J, Bi S, Ni Z, Shang X. Different interventions for post-ischaemic stroke depression in different time periods: a single-blind randomized controlled trial with stratification by time after stroke. *Clin Rehabil.* 2017;31(1):71–81.

Neuropsychiatric Disease and Treatment

Publish your work in this journal

Neuropsychiatric Disease and Treatment is an international, peer-reviewed journal of clinical therapeutics and pharmacology focusing on concise rapid reporting of clinical or pre-clinical studies on a range of neuropsychiatric and neurological disorders. This journal is indexed on PubMed Central, the 'PsycINFO' database and CAS,

Submit your manuscript here: <http://www.dovepress.com/neuropsychiatric-disease-and-treatment-journal>

Dovepress

and is the official journal of The International Neuropsychiatric Association (INA). The manuscript management system is completely online and includes a very quick and fair peer-review system, which is all easy to use. Visit <http://www.dovepress.com/testimonials.php> to read real quotes from published authors.