

CORRELATION OF LEVELS OF LACTIC ACID AND GLUCOSE IN CEREBROSPINAL FLUID OF CEREBRAL HEMORRHAGE PATIENTS WITH THE OCCURRENCE OF POSTOPERATIVE INTRACRANIAL INFECTION AND CLINICAL PROGNOSIS

KORELACIJA NIVOVA MLEČNE KISELINE I GLUKOZE U CEREBROSPINALNOJ TEČNOSTI KOD PACIJENATA SA CEREBRALNIM KRVARENJEM SA POJAVOM POSTOPERATIVNE INTRAKRANIJALNE INFEKCIJE I KLINIČKA PROGNOZA

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Summary

Background: Cerebral haemorrhage is a critical condition that often requires surgical treatment, and postoperative intracranial infection can significantly impact patient outcomes. The aim of the study was to examine the relationship between the levels of lactic acid and glucose in cerebrospinal fluid (CSF) of patients with cerebral haemorrhage and their postoperative intracranial infection and clinical prognosis.

Methods: The study selected the clinical data of 324 patients with cerebral haemorrhage who underwent surgical treatment in our hospital from March 2020 to March 2022 for retrospective analysis and divided these patients into the intracranial infection group (Group A, n=22, leukocyte values in CSF > 5 × 10⁶/L) and the non-intracranial infection group (Group B, n=302, leukocyte values in CSF ≤ 5 × 10⁶/L) according to the occurrence of postoperative intracranial infection in patients to detect the levels of lactic acid and glucose in CSF at different times in the two groups. Pearson method was adopted to analyze the correlation of the levels of lactic acid and glucose in CSF of patients with intracranial infection, and the Glasgow Outcome Scale (GOS) was used to assess the clinical prognosis of patients. According to their scores, these patients were divided into the good prognosis group (GPG, scores of 4–5 points, n=178) and the poor prognosis group

Kratak sadržaj

Uvod: Cerebralno krvarenje je kritično stanje koje često zahteva hirurško lečenje, a postoperativna intrakranijska infekcija može značajno da utiče na ishod. Cilj studije je bio da se analizira korelacija nivoa mlečne kiseline i glukoze u cerebrospinalnoj tečnosti (CSF) pacijenata sa cerebralnim krvarenjem sa pojavom postoperativne intrakranijske infekcije i kliničkom prognozom.

Metode: U studiji su selektovani klinički podaci 324 pacijenta sa cerebralnom hemoragijom koji su podvrgnuti hirurškom lečenju u našoj bolnici od marta 2020. do marta 2022. godine radi retrospektivne analize. Pacijenti su podeljeni u grupu intrakranijske infekcije (Grupa A, n=22, vrednosti leukocita u CSF > 5 × 10⁶/L) i grupu bez intrakranijske infekcije (Grupa B, n=302, vrednosti leukocita u CSF ≤ 5 × 10⁶/L) prema pojavi postoperativne intrakranijske infekcije kod pacijenata. U obe grupe su mereni nivoi mlečne kiseline i glukoze u CSF u različitim vremenima. Korišćena je Personova metoda za analizu korelacije nivoa mlečne kiseline i glukoze u CSF kod pacijenata sa intrakranijskom infekcijom, a za procenu kliničke prognoze pacijenata je korišćena Glasgow Outcome Scale (GOS). Prema njihovim rezultatima, pacijenti su podeljeni u dve grupe: grupu sa dobrim ishodom (GPG, 4–5 poena, n=178) i grupu sa lošim ishodom (PPG, 1–3 poena, n=146). Mereni su nivoi mlečne kiseline i glukoze u CSF

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(PPG, scores of 1–3 points, $n=146$). The levels of lactic acid and glucose in the CSF of patients in the two groups were measured, and the Pearson method was adopted to analyze the relationship between these levels and clinical prognosis.

Results: Compared with Group B, Group A had markedly higher lactic acid levels in CSF of patients at T1, T2 and T3 ($P<0.001$) and notably lower glucose levels ($P<0.001$). Patients in the PPG had notably higher lactic acid levels in CSF at T1, T2 and T3 ($P<0.001$) and overtly lower glucose levels than those in the GPG ($P<0.001$). Pearson's correlation analysis results showed that lactic acid levels in the CSF of patients were positively correlated with leukocyte values but negatively correlated with glucose levels ($P<0.05$). The lactic acid levels in CSF were negatively correlated with GOS scores but positively correlated with glucose levels ($P<0.05$).

Conclusion: The levels of lactic acid and glucose in CSF of patients with cerebral haemorrhage are correlated with postoperative intracranial infection and clinical prognosis, and the detection of the above indicators will help doctors better understand patients' condition, thus providing a scientific basis for the formulation of clinical treatment plans.

Keywords: cerebral haemorrhage, lactic acid, glucose, postoperative intracranial infection, clinical prognosis, correlation

Introduction

Cerebral haemorrhage is a primary disease that causes adult disability with high mortality. The main mechanism of early brain injury after cerebral haemorrhage is that blood accumulates locally after cerebral haemorrhage to form a hematoma, which can directly stimulate and compress the surrounding brain tissues, and cerebral haemorrhage often causes various symptoms of neurologic impairment, such as limb paralysis, language disorder and disturbance of consciousness (1). Surgery is still the first choice for treating cerebral haemorrhage, whose therapeutic objective is reducing the compression of hematoma on patients' brain tissues. At the same time, the toxic substances such as batroxobin and 5-hydroxytryptamine produced after haemorrhage lead to aggravation of cerebral oedema, and clearing the hematoma can alleviate the secondary pathological and physiological vicious cycles and effectively control the vasogenic and cytogenic brain oedema (2). However, this surgical treatment and retention of the postoperative drainage tube may lead to intracranial infection. In contrast, intracranial infection, one of the major factors causing a poor prognosis, can aggravate the neurological impairment of patients, prolong hospitalization time and increase the mortality of patients (3, 4). Bacteriological detection of cerebrospinal fluid (CSF) is the »golden standard« for clinical diagnosis of intracranial infection, but it is somewhat difficult to diagnose early intracranial infection because of longer culture time and a lower rate of positive cultivation (5).

kod pacijenata u obe grupe, a Personova metoda je korišćena za analizu veze između tih nivoa i kliničke prognoze.

Rezultati: U poređenju sa Grupom B, Grupa A je imala značajno više nivoa mlečne kiseline u CSF pacijenata u vreme T1, T2 i T3 ($P<0,001$), i primetno niže nivoa glukoze ($P<0,001$). Pacijenti u PPG grupi su imali primetno više nivoa mlečne kiseline u CSF u vreme T1, T2 i T3 ($P<0,001$) i jasno niže nivoa glukoze u poređenju sa GPG ($P<0,001$). Rezultati Personove metode su pokazali da su nivoi mlečne kiseline u CSF pacijenata pozitivno korelisani sa vrednostima leukocita, ali negativno korelisani sa nivoima glukoze ($P<0,05$). Nivoi mlečne kiseline u CSF su u negativnoj korelaciji sa rezultatima GOS skale, ali u pozitivnoj korelaciji sa nivoima glukoze ($P<0,05$).

Zaključak: Nivoi mlečne kiseline i glukoze u cerebrospinalnoj tečnosti (CSF) kod pacijenata sa cerebralnim krvarenjem su povezani sa postoperativnom intrakranijalnom infekcijom i kliničkom prognozom, a detekcija gore navedenih pokazatelja pomaže lekarima da bolje razumeju stanje pacijenata, pružajući naučnu osnovu za formulisanje kliničkih terapijskih planova.

Cljučne reči: cerebralno krvarenje, mlečna kiselina, glukoza, postoperativna intrakranijalna infekcija, klinička prognoza, korelacija

How to develop more scientific and objective diagnostic indicators becomes a critical subject facing clinicians. Studies have pointed out (6) that lactic acid, a product of anaerobic glycolysis of tissues, has notably higher levels of CSF compared with serum levels in the occurrence of intracranial bacterial infection. When the blood perfusion of brain tissues decreases, the oxygen supply of the brain will decrease, and the anaerobic glycolysis of brain tissues and lactic acid levels will increase (7). In addition, studies have also found (8, 9) that there is a correlation between glucose content in CSF and central nervous system (CNS) infection, and the permeability of the CSF barriers is related to the extent of glucose glycolysis in CSF. Moreover, when infection occurs in CNS, glucose content in CSF decreases under the enzymolysis released by pathogens and damaged cells. There is little literature at home and abroad to confirm the correlation of levels of lactic acid and glucose in CSF of cerebral haemorrhage patients with postoperative intracranial infection and clinical prognosis. By conducting clinical trials, this study adopts levels of lactic acid and glucose in CSF to predict postoperative intracranial infection and prognosis of patients with cerebral haemorrhage, thereby further accurately assessing the disease progression and prognosis of such patients, which is reported as follows.

Materials and Methods

General information

The study selected the clinical data of 324 patients with cerebral haemorrhage who underwent surgical treatment in our hospital from March 2020 to March 2022 for retrospective analysis, and the study has been reviewed and approved by the ethical committee of our hospital, conforming to the Declaration of Helsinki (2013) (10). All family members of patients were informed about the purpose and process of this pilot study and signed informed consent.

Inclusion and exclusion criteria

Inclusion criteria: (1) Patients met the diagnostic criteria of cerebral haemorrhage according to the Chinese Stroke Association Guidelines for Clinical Management of Cerebrovascular Disorders (11) and were diagnosed through clinical, magnetic resonance imaging (MRI) and computed tomography (CT) examinations. (2) Patients received treatment within 12 h at first morbidity (3). Patients had no disease affecting the physiological status of cerebral vessels, such as arteriosclerosis and cerebrovascular malformation, and no other neurological disease before the injury.

Exclusion criteria: (1) Patients with severe dysfunctions in the kidney, heart, liver and other organs; (2) patients with previous diseases such as brain tumour and cerebral infarction; (3) patients with haematological diseases; and (4) patients who were in pregnancy or lactation.

Methods

The first (T1) sample, derived from the cerebral ventricles, was immediately detected after ventricular drainage. The supernatant was extracted through centrifugation and precipitation before detection. Samples with hemolysis or more than 5% of interfering redundant erythrocytes were subsampled. Lactic acid levels were measured using the enzyme kinetics method with an automatic blood gas electrolyte analyzer, the Hitachi 7600 automatic biochemical analyzer (manufacturer: Shanghai Huanxi Medical Instruments Co., Ltd.), and matching reagents. The glucose levels in patient's cerebrospinal fluid (CSF) were measured using the glucose oxidase method.

The second (T2) sample was detected before removing the ventricular drainage tube, and the detection methods and procedures were the same as above.

The third (T3) sample was detected postoperatively, lumbar puncture was adopted to obtain CSF from patients, the matching reagents of Gem Premier 3500 (Instrumentation Laboratory, US) were used to detect lactic acid levels, and the methods and proce-

dures of glucose detection were the same as above. The normal Lactic acid levels in CSF range from 0.999 to 2.775 mmol/L, and normal glucose levels in CSF range from 2.5 to 4.4 mmol/L in humans.

Statistical analysis

The experimental data were all processed by Statistical Product and Service Solutions (SPSS) 26.0 software (SPSS Inc., Chicago, IL, USA) for statistical analysis, PASS software was adopted to calculate the study's sample size, and GraphPad Prism 7 (GraphPad Software, San Diego, USA) was used for image drawing. The enumeration data and the measurement data were detected by χ^2 and t-test, indicated by (n (%)) and ($\bar{x} \pm$ standard deviation (Sd)), respectively. The Pearson method was used to analyze the correlation of each indicator with GOS scores and leukocyte values, and the difference was considered statistically significant when $P < 0.05$.

Results

Comparison of clinical data of patients

There was no notable difference in clinical data such as age, BMI values and residence between the intracranial infection group (Group A) and the non-intracranial infection group (Group B) ($P > 0.05$), with comparability, as shown in *Table I*. There was no significant difference in age, BMI values and residence between the good prognosis group (GPG) and the poor prognosis group (PPG) ($P > 0.05$), with comparability, as shown in *Table II*.

Comparison of levels of lactic acid and glucose in CSF of patients at different times between Group A and Group B

At T1, T2, and T3, patients in Group A had significantly higher lactic acid levels in CSF ($P < 0.001$), whereas they had notably lower glucose levels than those in Group B ($P < 0.001$), as shown in *Table III* and *Table IV*.

Comparison of levels of lactic acid and glucose in CSF of patients between the GPG and the PPG

At T1, T2, and T3, patients in the PPG had notably higher lactic acid levels in CSF ($P < 0.001$), whereas they had notably lower glucose levels than those in the GPG ($P < 0.001$), as shown in *Table V* and *Table VI*.

Correlation of levels of lactic acid and glucose in CSF with intracranial infection

Results of Pearson showed that lactic acid levels in CSF of patients were positively correlated with

Table I Comparison of clinical data between Group A and Group B.

| Projects | Group A (n=22) | Group B (n=302) | χ^2/t | P |
|---|-------------------|-------------------|------------|-------|
| Gender | | | 0.774 | 0.379 |
| Male | 15 (68.18) | 231 (76.49) | | |
| Female | 7 (31.82) | 71 (23.51) | | |
| Age ($\bar{x}\pm s$, years old) | 62.05 \pm 11.45 | 58.17 \pm 10.20 | 1.874 | 0.075 |
| BMI values ($\bar{x}\pm s$, kg/m ²) | 21.20 \pm 1.48 | 21.03 \pm 1.50 | 0.032 | 0.975 |
| Time from injury to admission ($\bar{x}\pm s$, h) | 4.41 \pm 1.05 | 4.69 \pm 1.10 | 0.499 | 0.623 |
| Causes of injury | | | 0.001 | 0.981 |
| Spontaneous cerebral haemorrhage | 20 (90.91) | 275 (91.06) | | |
| Cerebral haemorrhage caused by brain trauma | 2 (9.09) | 27 (8.94) | | |
| Bleeding parts | | | 2.442 | 0.486 |
| Putaminal regions | 14 (63.64) | 231 (76.49) | | |
| Thalamus | 4 (18.18) | 43 (14.24) | | |
| Cerebral ventricles | 2 (9.09) | 12 (3.97) | | |
| Others | 2 (9.09) | 16 (5.30) | | |
| Residence | | | 0.454 | 0.500 |
| City | 9 (40.91) | 146 (48.34) | | |
| Countryside | 13 (59.09) | 156 (51.66) | | |

Table II Comparison of clinical data between the GPG and the PPG.

| Projects | GPG (n=178) | PPG (n=146) | χ^2/t | P |
|---|-------------------|-------------------|------------|-------|
| Gender | | | 0.008 | 0.927 |
| Male | 96 (53.93) | 78 (53.42) | | |
| Female | 82 (46.07) | 68 (46.58) | | |
| Age ($\bar{x}\pm s$, years old) | 58.37 \pm 10.55 | 57.87 \pm 10.71 | 0.275 | 0.784 |
| BMI values ($\bar{x}\pm s$, kg/m ²) | 21.04 \pm 1.55 | 21.23 \pm 1.61 | 0.883 | 0.379 |
| Time from injury to admission ($\bar{x}\pm s$, h) | 4.48 \pm 1.16 | 4.47 \pm 1.18 | 0.307 | 0.759 |
| Causes of injury | | | 0.654 | 0.419 |
| Spontaneous cerebral hemorrhage | 160 (89.89) | 135 (92.47) | | |
| Cerebral hemorrhage caused by brain trauma | 18 (10.11) | 11 (7.53) | | |
| Bleeding parts | | | 5.120 | 0.163 |
| Putaminal regions | 118 (66.29) | 97 (66.44) | | |
| Thalamus | 42 (23.60) | 28 (19.18) | | |
| Cerebral ventricles | 6 (3.37) | 13 (8.90) | | |
| Others | 12 (6.74) | 8 (5.48) | | |
| Residence | | | 0.003 | 0.955 |
| City | 92 (51.69) | 75 (51.37) | | |
| Countryside | 86 (48.31) | 71 (48.63) | | |

Table III Comparison of lactic acid levels in CSF of patients at different times between the two groups.

| Groups | n | T1 | T2 | T3 |
|---------|-----|-----------|-----------|-----------|
| Group A | 22 | 5.40±0.54 | 5.20±0.64 | 5.25±0.63 |
| Group B | 302 | 1.47±0.23 | 1.48±0.23 | 1.46±0.22 |
| t | | 31.080 | 24.944 | 26.962 |
| P | | <0.001 | <0.001 | <0.001 |

Table IV Comparison of glucose levels in CSF of patients at different times between the two groups.

| Groups | n | T1 | T2 | T3 |
|---------|-----|-----------|-----------|-----------|
| Group A | 22 | 2.47±0.44 | 2.56±0.36 | 2.45±0.37 |
| Group B | 302 | 3.66±0.56 | 3.64±0.56 | 3.64±0.52 |
| t | | 5.825 | 7.851 | 9.000 |
| P | | <0.001 | <0.001 | <0.001 |

Table V Comparison of lactic acid levels in CSF of patients at different times between the two groups.

| Groups | n | T1 | T2 | T3 |
|--------|-----|-----------|-----------|-----------|
| GPG | 178 | 2.59±0.23 | 2.59±0.21 | 2.60±0.25 |
| PPG | 146 | 4.03±0.73 | 3.95±0.68 | 4.00±0.74 |
| t | | 23.688 | 23.988 | 22.418 |
| P | | <0.001 | <0.001 | <0.001 |

Table VI Comparison of glucose levels in CSF of patients at different times between the two groups.

| Groups | n | T1 | T2 | T3 |
|--------|-----|-----------|-----------|-----------|
| GPG | 178 | 3.96±0.41 | 3.89±0.40 | 3.95±0.43 |
| PPG | 146 | 3.12±0.37 | 3.08±0.35 | 3.11±0.36 |
| t | | 17.954 | 19.842 | 17.606 |
| P | | <0.001 | <0.001 | <0.001 |

Table VII Correlation of levels of lactic acid and glucose in CSF of patients with leukocyte values.

| Indexes | Leukocyte values | |
|-------------|------------------|-------|
| | r | P |
| Lactic acid | 0.400 | <0.05 |
| Glucose | -0.973 | <0.05 |

Table VIII Correlation of levels of lactic acid and glucose in CSF of patients with GOS scores.

| Indexes | GOS scores | |
|-------------|------------|-------|
| | r | P |
| Lactic acid | -0.848 | <0.05 |
| Glucose | 0.921 | <0.05 |

leukocyte values but negatively correlated with glucose levels ($P<0.05$), as shown in *Table VII*.

Correlation of levels of lactic acid and glucose in CSF with GOS scores

Results of Pearson showed that lactic acid levels in CSF of patients were negatively correlated with GOS scores but positively correlated with glucose levels ($P<0.05$), as shown in *Table VIII*.

Discussion

Cerebral haemorrhage, which can have multiple underlying causes, is predominantly attributed to the rupture of hypertensive blood vessels with arteriosclerosis (12). It is a frequent neurosurgical emergency that leads to brain damage, including localized compression from the hematoma, injuries caused by releasing inflammatory factors and free radicals, and delayed cerebral injury characterized by brain cell oedema and metabolic dysfunction (13). Patients with cerebral haemorrhage often experience complex and severe injuries, with rapidly changing disease conditions and challenging treatment scenarios. Despite the advancements in clinical therapeutic techniques and the implementation of aseptic and antibacterial surgical practices, postoperative intracranial infection remains a concern. This underscores the need for clinicians to establish a reliable prognostic evaluation system to guide treatment decisions during the management of these patients (14, 15).

Lactic acid is a metabolic end product of glycolysis, and the concentration of lactic acid in CSF can reflect the glycolytic metabolism of CNS. When infections occur in patients (16), leukocytes will increase the anaerobic metabolism of glucose, thereby producing lactic acid and reducing PH values. The lactic acid in CSF is not affected by blood lactate levels, so the concentration in CSF reflects the extent of CNS infection in the body to some extent (17, 18). Some scholars have found (19) that when the body is infected, capillary endothelial damage and oedema of local tissues cause the decrease of microvessel density in tissues. Accumulation of lactic acid and the subsequent increase in its concentration occur due to metabolic dysfunction in local tissues and the impaired local clearance of lactic acid. Therefore, lactic acid levels are considered to be positively correlat-

ed with body infection. Through examining lactic acid levels in CSF between Group A and Group B after surgery, the study found that patients in Group A had overtly higher lactic acid levels in CSF than those in Group B ($P < 0.001$), predicting that patients with postoperative infection after traumatic brain injury have ischemia and hypoxia of local nerve cells to cause a massive release of lactic dioxygenases from nerve cells (20, 21) and an increase of the lactic acid concentration in CSF. It was further found that lactic acid levels in patients' CSF were positively correlated with leukocyte values ($r = 0.400$, $P < 0.05$), indicating that the detection of lactic acid levels in patients with cerebral haemorrhage can effectively predict whether patients have intracranial infection postoperatively.

Glucose is an important energy source for nerve cells, while bacteria also produce energy for their use by decomposing glucose (22, 23). When a bacterial infection occurs, bacteria decompose glucose in CSF, leading to lower glucose levels in the CSF. Therefore, it was found that patients in Group A had overtly lower glucose levels in CSF than those in Group B, indicating that glucose is involved in developing and progressing postoperative intracranial infection in patients. The study further found that glucose levels in CSF were negatively correlated with leucocyte values ($r = -0.973$, $P < 0.05$), indicating that the detection of glucose levels in CSF of patients can effectively predict whether patients have intracranial infection postoperatively. GOS score is a common scoring standard in neurosurgery, which has some auxiliary values in evaluating the prognosis of patients with conscious changes (24). This study found that patients in the PPG had notably higher lactic acid levels in CSF and significantly lower glucose levels than those in the GPG ($P < 0.001$). Results of Pearson showed that lactic acid levels in CSF of patients were negatively correlated with GOS scores but positively correlated with glucose levels ($P < 0.05$), indicating that the detection of levels of lactic acid and glucose in CSF of patients with cerebral haemorrhage can effectively predict clinical prognosis.

References

1. Magid-Bernstein J, Girard R, Polster S, Srinath A, Romanos S, Awad IA, et al. Cerebral Hemorrhage: Pathophysiology, Treatment, and Future Directions. *Circ Res* 2022; 130(8): 1204–29.
2. Powers WJ. Stroke/lore: Intracranial volumes and pressures following cerebral hemorrhage. *J Stroke Cerebrovasc* 2022; 31(9): 106637.
3. Lap B, Rai M, Tyagi W. Playing with colours: genetics and regulatory mechanisms for anthocyanin pathway in cereals. *Biotechnol Genet Eng* 2021; 37(1): 1–29.
4. McClelland SR. Postoperative intracranial neurosurgery infection rates in North America versus Europe: a systematic analysis. *Am J Infect Control* 2008; 36(8): 570–3.
5. Hristeva L, Bowler I, Booy R, King A, Wilkinson AR. Value of cerebrospinal fluid examination in the diagnosis of meningitis in the newborn. *Arch Dis Child* 1993; 69(5 Spec No): 514–7.
6. Cunha BA. Cerebrospinal fluid lactic acid levels: accurate, fast, and inexpensive. *Crit Care Med* 2011; 39(10): 2383–4, 2384–5.

Conflict of interest statement

All the authors declare that they have no conflict of interest in this work.

7. Annoni F, Peluso L, Gouvea BE, Creteur J, Zanier ER, Taccone FS. Brain Protection after Anoxic Brain Injury: Is Lactate Supplementation Helpful? *Cells-Basel* 2021; 10(7): 1714.
8. Sanjith S, S MK, G TP, M SS. Correlation between CSF Glucose Estimation using Glucometers against the Conventional Laboratory Technique in Determining Bacterial Meningitis: An Indian Study. *J Assoc Physicians India* 2020; 68(2): 43–7.
9. Yin L, Han Y, Miao G, Jiang L, Xie S, Liu B. CSF leukocyte, polykaryocyte, protein and glucose: Their cut-offs of judging whether post-neurosurgical bacterial meningitis has been cured. *Clin Neurol Neurosur* 2018; 174: 198–202.
10. World Medical Association Declaration of Helsinki: ethical principles for medical research involving human subjects. *Jama-J Am Med Assoc* 2013; 310(20): 2191–4.
11. Cao Y, Yu S, Zhang Q, Yu T, Liu Y, Sun Z, et al. Chinese Stroke Association guidelines for clinical management of cerebrovascular disorders: executive summary and 2019 update of clinical management of intracerebral haemorrhage. *Stroke Vasc Neurol* 2020; 5(4): 396–402.
12. An Z, Yin Y, Zhang L, Wang B, Cui T, Li M, et al. Effect of Ulinastatin Combined with Xingnaojing Injection on Severe Traumatic Craniocerebral Injury and Its Influence on Oxidative Stress Response and Inflammatory Response. *Biomed Res Int* 2022; 2022: 2621732.
13. Tian Y, Du HG, Fan CP, Wang C, Zhang GJ, Chen L, et al. Clinical significance of percutaneous endoscopic gastrostomy for patients with severe craniocerebral injury. *Chin J Traumatol* 2014; 17(6): 341–4.
14. Kim YJ, Moon KS, Kim SK, Kang SJ, Lee KH, Jang WY, et al. The difference in diffusion-weighted imaging with apparent diffusion coefficient between spontaneous and postoperative intracranial infection. *Brit J Neurosurg* 2014; 28(6): 765–70.
15. Berndt M, Lange N, Ryang YM, Meyer B, Zimmer C, Hapfelmeier A, et al. Value of Diffusion-Weighted Imaging in the Diagnosis of Postoperative Intracranial Infections. *World Neurosurg* 2018; 118: e245–53.
16. Aguirre M, Collins MD. Lactic acid bacteria and human clinical infection. *J Appl Bacteriol* 1993; 75(2): 95–107.
17. Nagaroor V, Gummadi SN. An overview of mammalian and microbial hormone-sensitive lipases (lipolytic family IV): biochemical properties and industrial applications. *Biotechnol Genet Eng* 2022: 1–30.
18. Boer K, Pfister W, Kiehntopf M. Lactic acid is of low predictive value for the diagnosis of bacterial infection in ventricular cerebrospinal fluid samples containing residual blood. *Clin Chem Lab Med* 2010; 48(12): 1777–80.
19. Valente DSL, Hoffmann A, Weiss G. Impact of bacterial infections on erythropoiesis. *Expert Rev Anti-Infe* 2021; 19(5): 619–33.
20. Wroblewski F, Ladue JS. Lactic dehydrogenase activity in blood. *Proc Soc Exp Biol Med* 1955; 90(1): 210–3.
21. Zhang W, Wang G, Xu ZG, Tu H, Hu F, Dai J, et al. Lactate Is a Natural Suppressor of RLR Signaling by Targeting MAVS. *Cell* 2019; 178(1): 176–89.
22. Tang BL. Glucose, glycolysis, and neurodegenerative diseases. *J Cell Physiol* 2020; 235(11): 7653–62.
23. Lopez-Gambero AJ, Martinez F, Salazar K, Cifuentes M, Nualart F. Brain Glucose-Sensing Mechanism and Energy Homeostasis. *Mol Neurobiol* 2019; 56(2): 769–96.
24. Yin W, Weng S, Lai S, Nie H. GCS score combined with CT score and serum S100B protein level Can evaluate severity and early prognosis of acute traumatic brain injury. *Nan Fang Yi Ke Da Xue Xue Bao* 2021; 41(4): 543–8.

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