Research Article

Effect of Huoxiang Zhengqi Pill on Early Neurological Deterioration in Patients with Acute Ischemic Stroke Undergoing Recanalization Therapy and Predictive Effect of Essen Score

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Early neurologic deterioration (END) in the acute phase of ischemic stroke is a serious clinical event, which is closely related to poor prognosis. Therefore, it is important to identify presentation features that predict END and take relevant treatment measures, as they could help to prevent the deterioration of high-risk patients. The prospective intervention study was carried out from January 2018 to December 2019. We included consecutive patients hospitalized for acute ischemic stroke (AIS) within 6 hours of onset. Patients were randomly assigned (1:1) to recanalization therapy plus Huoxiang Zhengqi Pill (HXZQ) (intervention group) or standard recanalization therapy alone (control group). The primary outcome was the development of END according to predefined criteria within the first 1 week of stroke onset. Poisson regression was used to identify predictors for END. Of the 155 patients enrolled in the study (age, 63 ± 11 years; 28.4% female), 20 (12.9%) developed END. Univariate analysis showed that the use of HXZQ and Essen stroke risk score (ESRS) (low risk group) were protective factors for END, while advanced age was a risk factor for END. However, in multivariate analysis, only ESRS (OR, 0.232; 95%CI, 0.058–0.928; P = 0.039) and the use of HXZQ (OR, 0.297; 95%CI, 0.096–0.917; P = 0.035) were statistically significant. ESRS can be used as the prediction factor of END. HXZQ has small side effects and wide indication. It could be used in the treatment of AIS.

1. Introduction

Stroke is a leading cause of long-term disability and death in China, of which about 70% are ischemic stroke [1]. Intravenous thrombolysis with recombinant tissue plasminogen activator within 4.5 hours after onset has been proved to be effective against acute ischemic stroke (AIS) and can reduce mortality. However, only a small number of patients were treated within the time window, and some patients still experienced neurologic deterioration after thrombolysis, which is called early neurologic deterioration (END) [2]. END is one of the most common complications of AIS, affecting about a third of ischemic stroke patients and increasing the risk of adverse functional outcomes and death [3–5]. Therefore, it is necessary to investigate the likely impact on the prediction and treatment strategies for END and to provide a novel strategy for the diagnosis and treatment of AIS.

Previous studies have found that coronary artery disease is usually associated with severe intracranial and extracranial atherosclerosis and that progressive neurological deterioration is more common in patients with coronary artery
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disease or carotid artery stenosis. This finding suggests that
patients with extensive atherosclerosis are more likely to
develop cerebrovascular disease and are more likely to have
progressive stroke [6, 7]. It is therefore necessary to explore
whether the Essen stroke risk score (ESRS) (including
atherosclerotic risk factors, such as hypertension, diabetes,
and smoking) can be used to predict END.

Huoxiang Zhengqi Pill (HXZQ) is mainly composed of
*pinellia ternata*, tangerine peel, licorice, and so on, which has
been applied to treat all kinds of diarrhea in China for
thousands of years. Pharmacological studies have shown that
composition is roughly the same as Erchen decoction, and
their main components (such as licorice) have platelet
aggregation [8], antioxidative [9], and neuroprotective
effects [10]. The main bioactive substances of licorice are
glycyrrhizin and its aglycone glycyrrhetinic acid, which
implicated in antioxidation, anti-inflammation, and neu-
roprotection [11, 12]. The regulatory effect of glycyrrhizin on
gap junction channels (GJCs) has been well demonstrated in
the previous studies. Because the electrical activity in the
brain comes from neuronal communication, GJCs not only
act as ion-transmitting neurons, but also distribute energy
flow between cells in different cell types. Stressed central
nervous system cells restore their derailment by increasing
the local expression of gap junction proteins in damaged
neurons and glial cells [13, 14]. Therefore, the aim of this
study was to explore the predictors of END and to observe
the preventive and therapeutic effects of HXZQ on END.

2. Methods

2.1. Research Design and Inclusion Criteria. This study is a
single-center, prospective, open-label, randomized con-
trolled research to blindly evaluate end-point events. The
study was supported by the ethics committee of the hospital
(No. GD2H-QR-KJ-018). This study was conducted to
evaluate the effect of HXZQ on AIS diagnosed by computed
angiography (CT) within 6 hours after onset. Subjects were
randomly assigned to HXZQ (Beijing Tongrentang,
Z13022498) treatment and control groups. Patients with
arterial occlusion confirmed by CT angiography, magnetic
resonance angiography, or digital subtraction angiography
should be admitted for endovascular treatment. Criteria for
selecting the subjects were as follows [15]: (1) prestroke
mRS Score 0-1, (2) age ≥ 18 years old, (3) treatment could
be started within 6 hours of onset (inguinal puncture), and
(4) informed consent of patients and/or agents. Exclusion
criteria: (1) intracranial hemorrhage, (2) significant mass
aggregation [8], antioxidative [9], and neuroprotective
effects [10]. The main bioactive substances of licorice are
glycyrrhizin and its aglycone glycyrrhetinic acid, which
implicated in antioxidation, anti-inflammation, and neu-
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neurons and glial cells [13, 14]. Therefore, the aim of this
study was to explore the predictors of END and to observe
the preventive and therapeutic effects of HXZQ on END.

2.2. Statistical Analysis. According to our previous research,
the incidence of END in our center is about 30%. Compared
with the control group, the incidence rate of END decreased
by 40% in the HXZQ treatment group with 80% power
(α = 0.05). PASS 11.0 software (NCSS LLC: Kaysville, UT,
USA) was used to calculate the sample size of 150 patients
with 75 cases in each group.

Numerical data were expressed as the mean ± standard
deviation or the median (quartile spacing), and categorical
variables were expressed by frequency (percentage). The
Shapiro-Wilk test was used in the normal test. Univariate
Poisson regression analysis was performed to compare the
baseline characteristics of the two groups according to
whether the END occurred or not. Fisher’s exact test was
used if the theoretical frequency ≤1. Finally, the independent
variables with P value less than 0.15 in univariate analysis
were included in multivariate analysis. Statistical analyses
were performed with the SPSS version 25.0 software package
(SPPS IBM Inc, Chicago, IL, USA). P value <0.05 was
regarded as statistically significant.

3. Results

A total of 155 patients with AIS within 6 hours of onset were
enrolled. The average age was 63 years, with 44 females
(28.4%). The main lesions of anterior and posterior circu-
lation were 111 cases (71.6%) and 44 cases (28.4%), re-
spectively. The low-risk group (0–2 ESRS) and the high-risk
group (≥3 ESRS) were 94 (60.6%) and 61 (39.4%) cases,
respectively, and 20 cases (12.9%) were diagnosed as END
within 7 days after admission. Most patients were treated
with antiplatelet (89.7%) and statins (95.5%), and 77 cases
(49.7%) were treated with HXZQ. The baseline character-
istics of the patients are shown in Table 1.

Univariate analysis showed a significant difference in age
(odds ratio (OR) 0.45; 95% confidence interval (CI)
1.006–1.085; P = 0.024), HXZQ use (OR 0.253; 95% CI
0.085–0.757; P = 0.014), and ESRS (low-risk group vs
medium–high-risk group: OR 0.162; 95% CI 0.054–0.485;
P = 0.001) between the END group and non-END Group,
in which the END group was older than the non-END Group, while the incidence of END was lower in HXZQ and ESRS low-risk groups (Table 1). As shown in Figure 1, in multivariate Poisson analysis, controlling for confounding factors, the incidence of END was lower in the low risk (ESRS) group and decreased in the HXZQ group; however, there was no statistical significance in age.

4. Discussion

END is a common and early event and is associated with poor prognosis in early-stage AIS. Therefore, the study of END predictors and the selection of appropriate therapy are related to the prognosis of AIS. Our findings indicated that patients with low-risk Essen scores have a low incidence of END, and oral administration of HXZQ may reduce the incidence of END.

In this study, END was present in 12.9% of AIS patients, which were similar to previously published studies (9.9% [17] and 13.5% [18]). One important clinically relevant finding was patients with a high risk in the ESRS are first reported to be more susceptible to END, a possible mechanism that links ESRS to END as a risk factor for cerebrovascular disease: first, previous studies have found that long-term hyperglycemia leads to lactic acid accumulation, promotes mitochondrial dysfunction, and aggravates the injury of AIS neurons, cerebral hypoperfusion, cerebral edema, and hemorrhagic transformation [19, 20]. Other studies have shown that diabetes causes an increase in metalloproteinase-9, which leads to increased permeability

| Table 1: Comparison of variables between all patients with and without END. |
|-----------------------------|---------------------|-----------------|-----------------|-----------------|
| Variable                    | All (n = 155)       | Unadjusted OR   | 95% CI          | P value         |
| Demographics                |                     |                 |                 |                 |
| Age, mean (SD), years       | 63 (11)             | 1.045           | 1.006–1.085     | 0.024           |
| Female, n (%)               | 44 (28.4)           | 0.925           | 0.355–2.407     | 0.873           |
| BMI, Median (IQR)           | 23.7 (21.5–25.4)    | 1.043           | 0.902–1.206     | 0.568           |
| Clinical characteristics    |                     |                 |                 |                 |
| Systolic blood pressure     | 142 (130–158)       | 0.991           | 0.971–1.012     | 0.409           |
| Diastolic blood pressure    | 85 (76–92)          | 0.990           | 0.955–1.027     | 0.584           |
| Sleeping well, n (%)        | 143 (92.3)          | 0.755           | 0.175–3.255     | 0.706           |
| Dysphagia, n (%)            | 43 (27.7)           | 1.536           | 0.513–4.594     | 0.443           |
| Baseline NIHSS score, median (IQR) | 6 (2–11) | 1.063           | 0.985–1.147     | 0.116           |
| Essen score classification, n (%) | 94 (60.6) | 0.162           | 0.054–0.485     | 0.001           |
| ≥3                          | 61 (39.4)           | Reference       |                 |                 |
| Cerebral infarction site, n (%) | 111 (71.6) | 0.925           | 0.355–2.407     | 0.873           |
| Anterior circulation        | 44 (28.4)           | Reference       |                 |                 |
| Medical history             |                     |                 |                 |                 |
| Smoking, n (%)              | 70 (45.2)           | 0.654           | 0.261–1.639     | 0.365           |
| Drinking, n (%)             | 41 (26.5)           | 0.309           | 0.072–1.332     | 0.115           |
| Hypertension, n (%)         | 98 (63.2)           | 2.327           | 0.778–6.959     | 0.131           |
| Diabetes, n (%)             | 51 (32.9)           | 0.874           | 0.336–2.274     | 0.782           |
| Dyslipidemia, n (%)         | 98 (63.2)           | 0.711           | 0.295–1.715     | 0.448           |
| Atrial fibrillation, n (%)  | 27 (17.4)           | 1.185           | 0.396–3.545     | 0.761           |
| Lab results                 |                     |                 |                 |                 |
| Uric acid, umol/L, mean (SD)| 376.1 (115.0)       | 1.000           | 0.997–1.005     | 0.654           |
| HDL, mmol/L, median (IQR)   | 1.0 (0.8–1.1)       | 3.333           | 0.833–13.334    | 0.089           |
| TC, mmol/L, median (IQR)    | 4.7 (3.9–5.3)       | 0.933           | 0.656–1.327     | 0.699           |
| TG, mmol/L, median (IQR)    | 1.7 (1.1–2.2)       | 0.937           | 0.560–1.568     | 0.804           |
| LDL, mmol/L, median (IQR)   | 2.9 (2.1–3.3)       | 0.749           | 0.484–1.160     | 0.194           |
| Glycated hemoglobin, %, median (IQR) | 6.0 (5.7–6.5) | 1.145           | 0.894–1.466     | 0.285           |
| D-dimer, mg/L, median (IQR) | 0.0 (0.3–1.4)       | 1.021           | 0.891–1.169     | 0.767           |
| Fibrinogen, g/L, median (IQR) | 3.6 (3.0–4.2) | 1.008           | 0.854–1.189     | 0.926           |
| FT3, Median (IQR), pmol/ml  | 4.3 (4.0–4.5)       | 0.497           | 0.172–1.433     | 0.196           |
| TSH, Median (IQR), mmol/ml  | 1.8 (0.9–2.1)       | 0.751           | 0.465–1.211     | 0.240           |
| FT4, Median (IQR), pmol/ml  | 15.2 (14.5–16.1)    | 0.907           | 0.716–1.149     | 0.419           |
| Treatment procedures        |                     |                 |                 |                 |
| Anticoagulant, n (%)        | 44 (28.4)           | 2.064           | 0.855–4.981     | 0.107           |
| Antiplatelet, n (%)         | 139 (89.7)          | 0.228*          |                 |                 |
| Statin use, n (%)           | 148 (95.5)          | 0.596*          |                 |                 |
| Antihypertensive drugs, n (%) | 87 (56.1) | 1.452           | 0.579–3.638     | 0.427           |
| Huoxiang Zhengqi Pill treatment, n (%) | 77 (49.7) | 0.253           | 0.085–0.757     | 0.014           |

* Fisher’s exact test
of the blood-brain barrier and infiltration of inflammatory cells, resulting in a poor prognosis of AIS. In addition, the increased hypercoagulability is also the key to vascular complications in patients with type 2 diabetes and is associated with poor prognosis in AIS patients. These are the potential mechanisms for poor prognosis after AIS in diabetic patients [21, 22]. Secondly, hypertension can lead to the abnormal dilation of cerebral microvessels in the ischemic area, which weakens the self-regulation function of cerebral vessels in this area and then leads to the increase of blood flow and damage of blood-brain barrier. In addition, when blood pressure decreased after successful recanalization, microvascular self-regulatory dysfunction may exacerbate the hypoperfusion of the penumbra [23]. Blood pressure maybe involved in the development of the END by changing the state of the hemodynamics, and the decrease of cerebral hemodynamic reserve and poor collateral circulation may be related to END. Therefore, early treatment of stenosis and control of arteriosclerosis can improve cerebral perfusion and thus prevent the progression and recurrence of cerebral infarction.

To the best of our knowledge, another important finding was the first report that HXZQ can reduce the incidence of END in patients with AIS. The theory of traditional Chinese medicine believes that excessive fat is easy to produce dampness and phlegm, while turbid phlegm generates the disorder of a spleen and stomach’s transportation. As a result, the body cannot absorb the essence of food or transport body fluid. If the body fluid accumulates, it is easy to produce phlegm, which can lead to vascular blockage. Animal studies showed that Erchen decoction (the same ingredients as HXZQ) could regulate the caveolae mRNA expression in rats fed with high-fat diet, lower the blood glucose, regulate the lipid metabolism, and reduce the insulin resistance; it also increased the expression of CDKAL1 mRNA and protein in mouse liver and subcutaneous fat and promoted the secretion of insulin [24]. In addition, glial cells first respond to AIS by upregulating transcription of early proinflammatory cytokines, such as TNF-α, leading to neuronal death and releasing neurotoxic substances into the bloodstream, further enhancing metalloproteinase expression [25, 26]. Moreover, animal experiments confirmed that TNF-α is closely related to acute brain edema [27]. In short, the upregulation of metalloproteinases may be a part of the vascular wall response induced by inflammatory stimuli (TNF-α) triggered by AIS. HXZQ can inhibit the expression of TNFα; therefore, it has a preventive and therapeutic effect on END [28].

Tangerine peel, one of the main components of HXZQ, contains more polymethoxy flavones, which plays a neuroprotective role by interacting with signal transduction pathways and/or altering cerebral vascular blood flow, including (1) alleviation of neurotoxin-induced neuronal damage, (2) inhibition of neuroinflammation, and (3) protecting of learning, memory, and cognition [29]. Thus, it is possible to prevent and treat END. In addition, licorice has biochemically active substances including glycyrrhizic acid and liquiritin. Glycyrrhizic acid protects neural tissue from hypoxic injury in vitro by modulating the PI3K/Akt pathway and inhibiting the glutamate-NMDA mediated neurotoxicity [30, 31]. It is reported that glycyrrhizin also plays a key role in axonal regeneration and nerve tissue repair [32]. Moreover, licochalcone was also reported to reduce platelet activation in experimental animals and humans by inhibiting cyclooxygenase-1 (COX-1) activity [33] and collagen-induced platelet aggregation [34]. These results indicated that benefits of HXZQ could be considered as a potential therapeutic agent to control or limit END.

There are some limitations in the present study. First, the study is a single-center inpatient study with a small sample size.
size, and the results need to be further validated in a large multicenter prospective study. Second, although the attending neurologists believe that endovascular therapy should be initiated immediately if the lesion vessels fail to recanalize after thrombolysis, the manner of reperfusion therapy is inconsistent. Third, the criteria for defining END have been inconsistent in the degree or time frame of neurological deficits and are still evolving [35]. In the present study, END was defined as an increase of 2 points or more in NIHSS score within 72 hours after admission. This criterion is used because it minimizes interevaluator variations and distinguishes END from late deterioration mainly caused by stroke complications such as recurrent stroke or aspiration pneumonia [36].

5. Conclusions
In conclusion, ESRS could be used as a predictor factor of END. While HXZQ, as a kind of Chinese herbal medicine, has the characteristics of multicomponent and multi-therapeutic target, which can reduce the occurrence of END and is suitable for the treatment of AIS in acute stage. A further study could assess the long-term effects of HXZQ in a large-scale prospective randomized controlled trial.

Data Availability
The study data that underlie the results of this article will be given to investigators within two months of the committee’s approval of the research proposal.

Conflicts of Interest
The authors report no conflicts of interest.

Authors’ Contributions
ZH and JL were involved in the study design; ZH, JL, CZ, YD, SL, and XL were responsible for the study implementation and data collection; ZH and CZ were involved in the data analysis and data interpretation; ZH, JL, and XL prepared the manuscript. All authors read and approved the final manuscript.

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