



A pilot study in acute subarachnoid haemorrhagic patients after aneurysm clipping with complementary therapies of Chinese medicine

Han-Chung Lee^{a,b,c}, Ching-Liang Hsieh^{d,e,f,*}, Chun-Chung Chen^a,
Der-Yang Cho^a, Kuang-Fu Cheng^g, Pao-Hsuan Lin^{g,h}

^a Department of Neurosurgery, China Medical University Hospital, Taichung, Taiwan

^b Graduate Institute of Integrated Medicine, China Medical University, Taichung, Taiwan

^c School of Medicine, China Medical University, Taichung, Taiwan

^d Graduate Institute of Acupuncture Science, China Medical University, Taichung, Taiwan

^e Acupuncture Research Center, China Medical University, Taichung, Taiwan

^f Department of Chinese Medicine, China Medical University Hospital, Taichung, Taiwan

^g Biostatistics Center, China Medical University, Taichung, Taiwan

^h Institute of Environmental Health, China Medical University, Taichung, Taiwan

Available online 1 September 2010

KEYWORDS

Complementary therapies;
Chinese medicine;
Acute subarachnoid haemorrhage;
Glasgow Outcome Scale;
Total admission day

Summary

Objectives: Acute subarachnoid haemorrhage still has high mortality and morbidity despite the use of modern standard treatment. In Taiwan, complementary therapies of Chinese medicine are usually used to treat stroke patients. The aim of this study was to investigate the effect of complementary therapies of Chinese medicine on patients with acute subarachnoid haemorrhage after aneurysm clipping.

Design: This study was designed as a pilot study. A total of 32 patients with acute subarachnoid haemorrhage were randomly assigned to either a Chinese herbs extra group (CH) in which the patients were given complementary therapies of Chinese medicine and standard treatment or a standard treatment only group (ST) in which patients were given standard treatment only.

Main outcome measures: Glasgow Outcome Scale scores, which were assessed by an evaluator who was blinded to the groups, 3 months after admission, and total admission days including intensive care unit stay days.

Results: The average Glasgow Outcome Scale score 3 months after admission was 3.7 ± 1.4 in the CH was greater than 3.0 ± 1.7 in the ST ($p=0.041$). Average total admission days were 53.9 ± 28.6 (median 61) in the ST longer than 28.1 ± 19.1 (median 20.5) in the CH ($p=0.004$).

Conclusion: Traditional Chinese medicine for the treatment of patients with acute subarachnoid haemorrhage is of value because they can increase Glasgow Outcome Scale scores 3 months after admission and also because they can reduce total admission days.

© 2010 Elsevier Ltd. All rights reserved.

* Corresponding author at: Graduate Institute of Acupuncture Science, China Medical University, 91 Hsueh-Shih Road, Taichung 40402, Taiwan. Tel.: +886 4 22053366x3600; fax: +886 4 22035191.

E-mail address: clhsieh@mail.cmuh.org.tw (C.-L. Hsieh).

Introduction

Subarachnoid haemorrhage (SAH) is an acute emergency. About 85% of cases result from the rupture of an aneurysm.¹ The incidence of SAH is about 6 cases per 100,000 patient years.¹ The overall case mortality rate of SAH is 42% during the first 28 days.² Although ultra-early aneurysm clipping (within 3 days after onset) is used for the treatment of SAH, higher mortality and morbidity rates are still noted compared with the other cerebral diseases due to occurrence of vasospasm following SAH, which may cause cerebral ischaemia.³ The use of complementary therapies of Chinese medicine (CM) including Chinese herbs and acupuncture to treat patients with chronic or subacute stage of stroke is popular in Taiwan. *Salviae miltiorrhizae* may increase recovery rate of patients with acute SAH had been reported.⁴

Aneurysm rupture with SAH may cause inflammation resulting in fatal vasospasm and central pyrexia.⁵ Patients with SAH who have complicated fever may have a prolonged stay in the ICU and a poorer outcome.^{6,7} Mechanical compression of the brainstem and hypothalamus may induce the production and release of pro-inflammatory cytokines, including interleukin-1 β (IL-1 β), IL-6, tumor necrosis factor- α (TNF- α), and S100B as a calcium-binding protein of astrocytes, causing central pyrexia.^{8–10} The pro-inflammatory cytokine levels increase in brain tissues, cerebrospinal fluid (CSF), and blood in patients with traumatic brain injury or stroke.^{11–25}

Traditional Chinese medicine (TCM) such as acupuncture, Chinese herbs, and CM formula has been used to treat stroke for centuries in China. Our previous studies have shown that *Gastrodia elata* plays a neuroprotective role in kainic acid-treated rats²⁶ and may reduce IL-1 β and TNF- α levels in brain tissue of kainic acid-treated rats.²⁷ *Paeonol*, a major component of Moutan Cortex of *Paeonia suffruticosa* Andrews and the root of *Paeonia lactiflora* Pall, has anti-inflammation and anti-oxidant effects in transient ischaemia-reperfusion injury rats.²⁸ Therefore, we designed a pilot study to investigate the effect of complementary therapies of CM on patients with acute SAH.

Materials and methods

Subjects

A total of 53 patients with acute SAH were treated at China Medical University Hospital, Taichung, Taiwan from January 2007 and December 2007. Thirty-two patients who underwent craniotomy were included in the study. The inclusion criteria were the following: (1) SAH due to cerebral aneurysm rupture that was confirmed by sequential computed tomography angiography (CTA) scanning within 6 h after the episode; (2) the neurological deficit was between grades 2 and 4 of Hunt and Hess (H&H) grade.²⁹ The exclusion criteria were as follows: (1) patients with pregnancy; (2) age <12 years or >70 years; (3) H&H grade of 1 and 5; (4) patients or their families refuse participation in trial.

Study design

All the experimental procedures were according to the ethical principles dictated in the Declaration of Helsinki. The

protocol of the trial was approved by the institutional review board of the China Medical University Hospital, Taichung City, Taiwan (DMR95: IRB80), and informed consent regarding the experimental procedures and purpose was obtained prior to the trial.

After undergoing an aneurysm clipping operation the patients were randomized by an on duty doctor who takes a lot of Chinese herbs extra group (CH) or standard treatment only group (ST) from a dark box to either an CH which received complementary therapies of CM and standard treatment or a ST which received standard treatment only. Each group had 16 subjects. Because this study was a pilot study, there was no basis for calculating its power and sample size (Fig. 1).

Standard treatment

Standard treatment of acute SAH according to the guidelines of the Stroke Council, American Heart Association³⁰ is based on clipping of the aneurysm as early as possible and the prevention of secondary insults to the brain. External ventricular drainage was performed routinely during aneurysm clipping. It was used not only to monitor the postoperative ICP but also for drainage of the intraventricular haemorrhage and was maintained for not more than 7 days to avoid related infection. All patients were intubated and placed on volume-controlled ventilation under sedation to maintain partial pressure of oxygen in arterial blood (PaO₂) of at least 100 mm Hg and arterial carbon dioxide pressure or tension (PaCO₂) of approximately 35–40 mm Hg after the operation. The endotracheal tube was not removed until the consciousness of the patient was clear and the ICP was stable. Hypertension, hyperperfusion, haemodilution and calcium-channel blocking agent (nimodipine) were started immediately after arrival in the ICU after surgery. ICP was treated by elevating the patient's head by raising one end of the bed, sedation, paralysis, and mannitol. Nutritional support was started as soon as possible and was maintained by administering adequate parenteral or enteral solutions. Oral acetaminophen was given regularly to prevent further pyrexia.

The complementary therapies of CM

The complementary therapies of CM were given every day continuously for 2 weeks after the patients started to take food on the second day after the surgical operation. Patients were mainly given the following four essential Chinese herbs: (1) *Astragalus membranaceus* (Fisch) Bunge (Radix Astragalli, 12 g/day; Shaanxi, China); (2) *G. elata* Blume (Rhizoma Gastrodiae, 12 g/day; Sichuan, China); (3) *Acorus gramineus* Soland (Rhizoma Acori Graminei, 7.5 g/day; Sichuan, China); and (4) *Pheretima aspergillum* (E. Perrier) (Lumbricus, 12 g/day; Thailand). In addition, Chinese herbs such as *P. suffruticosa* Andr (Cortex Moutan, 12 g/day; Zhejiang, China), *Lonicera japonica* Thunb (Flos Lonicerae, 19 g/day; Anhui, China), raw *Rehmannia glutinosa* Libosch (Radix Rehmaniae, 12 g/kg; Henan, China), and *Scutellaria baicalensis* Georgi (Radix Scutellariae, 12 g/kg; Hebei, China) were given when patients had fever or a heat phenomenon in TCM such as quickened radial pulse

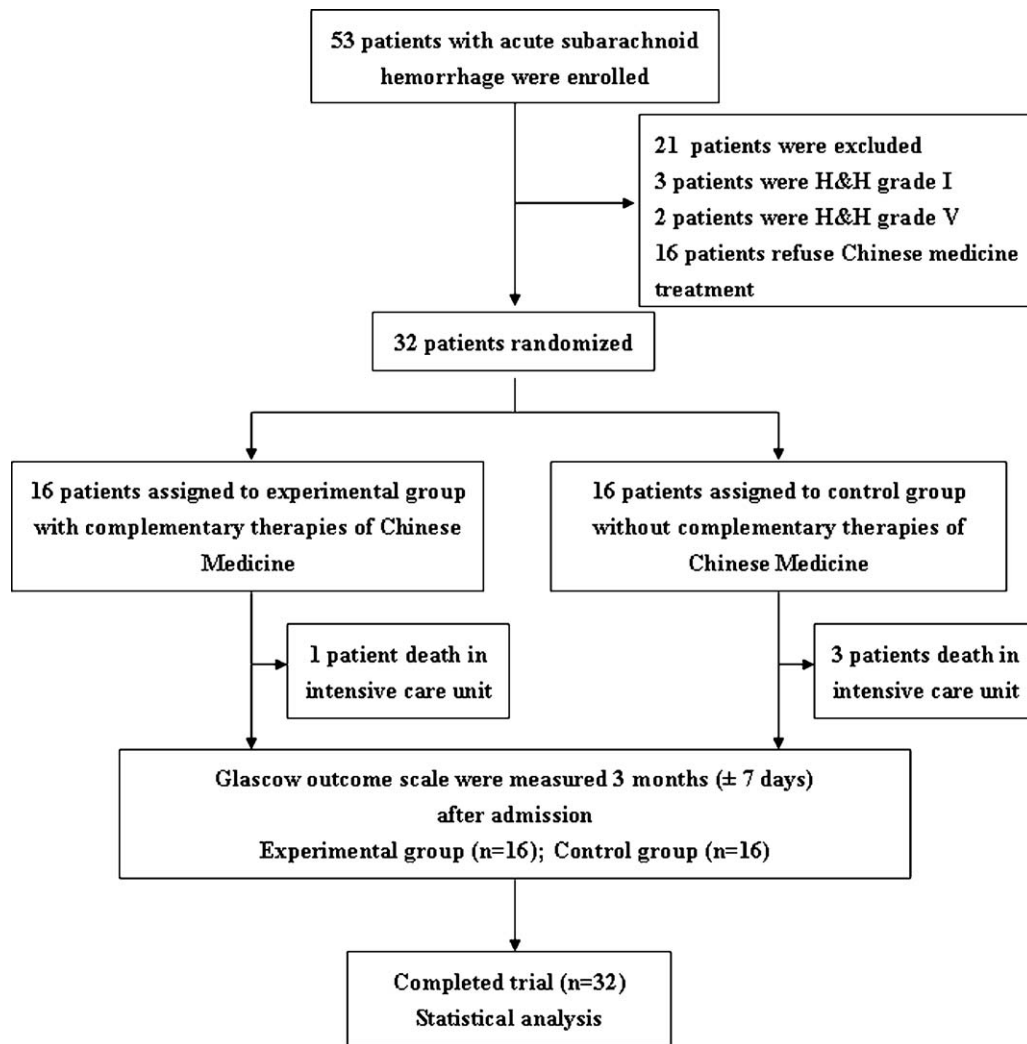


Figure 1 Flowchart.

(more than 85 beats/min), tongue color was fresh red, etc. *Rheum palmatum* Linn (Radix and Rhizoma Rhei, 4g/kg; Sichuan, China) and *Citrus aurantium* Linn (Fructus Aurantii Immaturus, 12g/kg; Sichuan, China) were added when patients had no defecation for more than 3 days. The Chinese herbs were authenticated according to the characteristics and shape even histological section, and decocted by the Chinese herb specialist in the China Medical University Hospital. These herbs were mixed with 600cm³ of water, and then decocted to 300cm³. Patients were given 100cm³ of the solution of herbs three times a day.

Clinical characteristics and basic data recording

The age, gender, GCS scores, H&H grade and SAH grade of the patients were recorded on the day of admission.

Main outcome measure

GOS scores were assessed and recorded on the personal medical record in the outpatient door 3 months (± 7 days) after

admission by an evaluator who was blinded to the group. The GOS scores were divided into five grades from 1 to 5: score 1, death; score 2, vegetative state; score 3, severe disability; score 4, moderate disability; score 5, mild or no disability.³¹ In addition, total admission days were used as an outcome measure including ICU stay days.

Secondary outcome measure

The Glasgow Coma Scale (GCS) scores and H&H grade were recorded on the day of admission, and the GCS scores were also recorded on the day of discharge. The daily body temperature (BT), taken with an ear thermometer, and daily ICP were recorded. In addition, cytokine levels including IL-1 β , IL-6, TNF- α , and S100B were measured in the cerebrospinal fluid on the 1st and 5th day after the operation.

Daily BT and ICP recordings

The BT and ICP were monitored every 2h continuously for 5 days after surgery. The average BT and ICP were calculated. The variation in BT was calculated (daily highest

BT – average daily BT)² and the variation in ICP was calculated (daily highest ICP – average daily ICP).²

The measurement of cytokine levels

Three-milliliter samples of CSF were collected on the 1st and 5th days from the external ventricle drainage of the lateral ventricle after surgery. The samples were centrifuged for 20 min at 2000 rpm, and the supernatant was immediately stored at -80°C until analysis. The levels of IL-1 β , IL-6, and TNF- α were determined by using a commercial enzyme-linked immunosorbent assay (ELISA) kit (Bender MedSystems, Inc., USA) and an ELISA reader (Dynex MRX, Virginia, USA). The sensitivity of the assay was typically 0.124 pg/ml for IL-1 β , 0.094 pg/ml for IL-6, and 0.081 pg/ml for TNF- α . The S100B level in CSF was quantified by sandwich ELISA. The samples were analyzed in duplicate and compared with known concentrations. The lower limit of detection of the ELISA is 0.01 ng/ml. No cross-reactivity or interference with other related interleukins was observed. The data were represented in pg/ml and all assays were performed in duplicate.

Statistical analysis

The means and standard deviations were used to summarize the continuous variables in Tables 1–3. Wilcoxon's rank-sum test and log rank test were applied to compare the two groups. The Spearman correlation coefficients between BT and IL-1 β Df, IL-6 Df, TNF Df and S100B Df were computed to strength association.

Results

The analysis of basic data

The age, gender, GCS score on admission, H&H grade score on admission, and SAH grade score on admission were not significantly different between the CH and ST (Table 1).

Main outcome measures

The GOS score was assessed in all the 32 patients 3 months after admission. The average GOS score was 3.7 ± 1.4 in the CH which was significantly greater than the score of 3.0 ± 1.7 in the ST ($p = 0.041$). A total of four patient's death

due to intractable increased ICP in ICU, and the complication included pulmonary infection, urinary tract infection, gastrointestinal tract haemorrhage, etc., but no patient withdrew or adverse events were noted in the trial. The mortality rate during the stay in the ICU was 6.25% (1/16) in the CH which was similar to the rate of 18.8% (3/16) in the ST ($p = 0.473$). The average admission days was 53.9 ± 28.6 (median 61) in the ST was longer than 28.1 ± 19.6 (median 20.5) in the CH ($p = 0.004$). The mean ICU stay days was 26.9 ± 17.1 (median 26) in the ST longer than 11.9 ± 4.3 (median 11) in the CH ($p = 0.002$).

Secondary outcome measure

The average GCS score at discharge was 12.3 ± 2.5 in the CH greater than 10.3 ± 2.9 in the ST ($p = 0.037$).

Effect of complementary therapies of CM on BT and ICP

The average daily BT and variation of daily BT, and the average daily ICP and variation of daily ICP (VDICP) from the 1st to 5th days after surgery were not significantly different between the CH and ST (Tables 2 and 3) except VDICP of day 4 ($p = 0.038$).

Effect of complementary therapies of CM on IL-1 β , IL-6, TNF- α , and S100B in CSF

The difference between IL-1 β levels on the 1st day and IL-1 β levels on the 5th day was positively correlated with the average daily BT on the 1st, 4th, and 5th days after surgery in the CH ($p = 0.015$, 0.022 and 0.023, respectively; Table 4), whereas there was no correlation with average daily BT on the 2nd and 3rd days after surgery in the CH (Table 4). The difference between IL-1 β levels on the 1st day and IL-1 β levels on the 5th day had no correlation with the average BT on the 1st, 2nd, 3rd, 4th, and 5th days after surgery in the ST (Table 4).

The difference between IL-6 levels on the 1st day and IL-6 levels on the 5th day had no correlation with the average daily BT on the 1st, 2nd, 3rd, 4th, and 5th days after surgery in the CH and ST (Table 4).

The difference between TNF- α levels on the 1st day and TNF- α levels on the 5th day had no correlation with the average BT on the 1st, 2nd, 3rd, 4th, and 5th days after surgery in the CH and ST (Table 4).

Table 1 Clinical characteristics and basic data in acute subarachnoid haemorrhagic patients (mean \pm standard deviation).

	CH ($n = 16$)	ST ($n = 16$)	p -Value
Age (years)	55.1 ± 12.9	59.9 ± 12.9	0.417
Gender (female/male)	8/8	8/8	1.000
GCS score in admission	9.6 ± 4.8	8.5 ± 4.4	0.436
H&H grade in admission	2.9 ± 1.2	3.3 ± 1.2	0.382
SAH grade in admission	2.1 ± 0.9	2.6 ± 0.9	0.266

n : patients number; CH: Chinese herbs extra group, acute subarachnoid haemorrhagic patient with complementary therapies of Chinese medicine; ST: standard treatment only group, acute subarachnoid haemorrhagic patient with standard treatment only; GCS: Glasgow coma scale; H&H: Hunt & Hess; SAH: subarachnoid haemorrhage.

Table 2 The daily body temperature changes in acute subarachnoid haemorrhagic patients (mean \pm standard deviation).

	ADBT			VDBT		
	CH	ST	<i>p</i> -Value	CH	ST	<i>p</i> -Value
Day 1	36.88 \pm 0.54	37.13 \pm 0.77	0.526	0.18 \pm 0.23	0.23 \pm 0.46	0.679
Day 2	37.19 \pm 0.46	37.26 \pm 0.54	0.641	0.13 \pm 0.19	0.21 \pm 0.42	0.503
Day 3	37.06 \pm 0.59	37.17 \pm 0.40	0.373	0.09 \pm 0.10	0.13 \pm 0.18	0.421
Day 4	37.25 \pm 0.64	37.31 \pm 0.56	0.403	0.12 \pm 0.14	0.12 \pm 0.17	0.956
Day 5	37.36 \pm 0.54	37.31 \pm 0.69	0.852	0.15 \pm 0.22	0.24 \pm 0.42	0.445

CH: Chinese herbs extra group, acute subarachnoid haemorrhagic patient with complementary therapies of Chinese medicine; ST: standard treatment only group, acute subarachnoid haemorrhagic patient with standard treatment only; ADBT: averaged daily body temperature; VDBT: variation of daily body temperature; Day 1: 1st day after surgical operation; Day 2: 2nd day after surgical operation; Day 3: 3rd day after surgical operation; Day 4: 4th day after surgical operation; Day 5: 5th day after surgical operation; Wilcoxon's signed rank test.

Table 3 The daily intracranial pressure in acute subarachnoid haemorrhagic patients (mean \pm standard deviation).

	ADICP			VDICP		
	CH	ST	<i>p</i> -Value	CH	ST	<i>p</i> -Value
Day 1	7.06 \pm 4.52	6.89 \pm 4.23	0.914	25.25 \pm 47.91	16.44 \pm 18.81	0.695
Day 2	9.16 \pm 4.40	11.00 \pm 8.62	0.456	5.91 \pm 10.55	7.24 \pm 15.14	0.628
Day 3	9.18 \pm 4.36	10.26 \pm 7.61	0.623	2.17 \pm 2.23	4.24 \pm 5.86	0.551
Day 4	9.78 \pm 6.03	10.10 \pm 7.23	0.891	1.70 \pm 3.75	13.06 \pm 22.22	0.038
Day 5	7.93 \pm 2.66	10.71 \pm 8.05	0.205	4.17 \pm 9.79	5.96 \pm 9.07	0.147

CH: Chinese herbs extra group, acute subarachnoid haemorrhagic patient with complementary therapies of Chinese medicine; ST: standard treatment only group, acute subarachnoid haemorrhagic patient with standard treatment only; ADICP: averaged daily intracranial pressure; VDICP: variation of daily intracranial pressure; Day 1: 1st day after surgical operation; Day 2: 2nd day after surgical operation; Day 3: 3rd day after surgical operation; Day 4: 4th day after surgical operation; Day 5: 5th day after surgical operation; Wilcoxon's signed rank test.

Table 4 The correlation coefficient of cytokines between averaged daily body temperatures in acute subarachnoid haemorrhage patients.

Group	BT				
	1	2	3	4	5
IL-1 β Df					
CH	0.59*	0.12	0.29	0.57**	0.56***
ST	-0.48	-0.41	-0.38	-0.14	-0.25
IL-6 Df					
CH	0.23	-0.39	-0.27	0.11	0.15
ST	0.01	0.05	0.52	-0.10	-0.54
TNF- α Df					
CH	0.08	-0.17	0.40	0.42	0.38
ST	0.4	0.00	0.06	0.00	0.30
S100B Df					
CH	0.31	0.17	0.31	-0.03	-0.04
ST	0.18	0.26	0.12	0.26	0.19

CH: Chinese herbs extra group, acute subarachnoid haemorrhagic patient with complementary therapies of Chinese medicine; ST: standard treatment only group, acute subarachnoid haemorrhagic patient with standard treatment only; Df: the difference of concentration between 1st and 5th day; BT: averaged daily body temperatures; 1: 1st day after surgical operation; 2: 2nd day after surgical operation; 3: 3rd day after surgical operation; 4: 4th day after surgical operation; 5: 5th day after surgical operation; Spearman's correlation coefficients.

* $p=0.015$.

** $p=0.022$.

*** $p=0.023$.

The difference between S100B levels on the 1st day and S100B levels on the 5th day had no correlation with the average BT on the 1st, 2nd, 3rd, 4th, and 5th days after surgery in the CH and ST (Table 4).

Discussion

Our results indicated that complementary therapies of CM for patients with acute SAH may increase the GOS score 3 months after admission, and reduce the total number of admission days, including both ICU stay days, which suggests that complementary therapies of CM provide an advantage in outcome for such patients. Critically ill patients with SAH commonly have fever, a factor known to worsen neurologic injury due to vasospasm.¹⁴ In these patients, fever and vasospasm may be both associated with the production and release of pro-inflammatory cytokines, including IL-1 β , IL-6, and TNF- α .¹⁸⁻²² Increased protein levels of pro-inflammatory cytokines have been reported in brain tissues, cerebrospinal fluid, and blood of patients with SAH, traumatic brain injury, stroke, and other neurological conditions.^{23,24}

Interleukin-1 β is thought to play an important role in mediating inflammation and neuronal damage after traumatic brain injury, spontaneous SAH, and stroke by enhancing the inflammatory reactions via the release of other inflammatory mediators such as prostaglandins, collagenase, and phospholipase A₂.¹² Additionally, IL-1 β has been implicated in apoptotic cell death,^{13,14} leukocyte-endothelium adhesion,¹⁵ blood-brain barrier disruption,¹⁶ edema formation,^{16,17} astrogliosis, and neovascularization.¹⁸ Experimentally, intracerebroventricular administration of IL-1 β is associated with marked stimulation of circulating IL-6 and TNF- α levels.^{19,21} Inhibition of IL-1 β has been shown to reduce the incidence of central pyrexia, vessel spasm, and early edema formation.^{22,23} Our studies showed that complementary therapies of CM in patients with acute SAH did not significantly change the average daily BT or variation of daily BT, or the average daily ICP or variation of daily ICP except day 4 due to two patients death with increased ICP. The difference in IL-1 β concentration between the 1st day and 5th day after surgery showed a positive correlation with daily BT on the 1st, 4th, and 5th day after surgery in the complementary therapies of CM, whereas there were no similar results in the ST which did not receive complementary therapies of CM. These results suggest that complementary therapies of CM may decrease IL-1 β concentration in CSF which reduces the inflammation and fever caused by SAH. Unfortunately, this tendency was not observed for IL-6, TNF- α , and S100B. More frequent checking of the concentration of cytokines and a longer observation period may be helpful for demonstrating a significant difference.

That *G. elata* and its component *vanillyl alcohol* may inhibit the production and scavenging of oxygen free radicals, and inhibit microglia activation in kainic acid-induced epileptic rats was shown in our previous studies.^{26,32,33} *Astragaloside IV* is a component of *Astragalus membranaceus* that can reduce the cerebral infarction area induced by middle cerebral artery occlusion in rats, and this effect of *Astragaloside IV* results from its anti-oxidative properties.³⁴ *Acorus gramineus* has the action of resolving

phlegm to open orifices in TCM, and can enhance learning and memory.³⁵ *Lumbrokinase* is a component of *P. aspergillum* (earthworm), and can mediate via antiplatelet activity, attenuating the calcium release from calcium storage, inhibiting intracellular adhesion molecular-1 (ICAM-1) expression to protect against cerebral ischaemia.³⁶ The *Paeonol* component of *P. suffruticosa* may reduce the cerebral infarction area and neurological deficit, and also has anti-oxidative action in cerebral ischaemia-reperfusion injured rat.²⁸ The component of *Lonicera japonica* may inhibit microglia activation to protect dopaminergic neurons from lipopolysaccharide (LPS)-induced injury,³⁷ and also may inhibit nuclear factor- κ B (NF- κ B) and activator protein-1 (AP-1) to suppress inflammatory reaction in mouse alveolar macrophage.³⁸ *Catapol* is a component of *R. glutinosa* that may reduce lipid peroxidation and also may increase glutathione and superoxide dismutase activities in MPP⁺-induced oxidative stress in mesencephalic neurons.³⁹ *Catapol* also can reduce the formation of intracellular reactive oxygen species in astrocytes with H₂O₂-induced oxidative stress.⁴⁰ The baicalin component of *S. baicalensis* can mediate via binding to chemokines to produce anti-inflammatory activity in human peripheral blood leucocytes,⁴¹ and *baicalein* also can maintain brain mitochondrial homeostasis and function in rats with chronic cerebral hypoperfusion-induced oxidative damage.⁴² *R. palmatum* is a laxative, and its emodin component can through the inhibition of AP-1 and NF- κ B suppress matrix metalloproteinase in human cancer cells.⁴³ *C. aurantium* is qi-regulating and is medicinal for digesting, and it has anti-oxidant activity.⁴⁴ To sum up, the Chinese herbs taken together may produce anti-oxidation and anti-inflammation including the inhibiting generation and scavenging of oxygen free radicals, and the inhibition of IL-1 β , suggesting that these action may improve functional recovery of SAH patients.

The present study was a pilot study and therefore there were some limitations as follows: (1) treatment with the complementary therapies of CM had to be agreed by the patients or their families, thus randomly assigning the patients to the ST or CH by a completely blind method was difficult; (2) the sample size was small; (3) there was no fixed CM formula. Future research using a randomized double-blind study design, an increased number of patients, and a fixed CM formula is needed.

In conclusion, TCM in patients with acute SAH is a worthy extension to treatment because they can increase GOS scores at 3 months after admission and also reduce total admission days including ICU stay days.

Acknowledgments

This study was supported by grant DMR 95-044 from the China Medical University Hospital, Taichung, Taiwan, and was supported in part by Taiwan Department of Health Clinical Trial and Research Center of Excellence (DOH99-TD-B-111-004).

References

1. van Gijn J, Rinkel GJ. Subarachnoid haemorrhage: diagnosis, causes and management. *Brain* 2001;124:249–78.

2. Ingall T, Asplund K, Mähönen M, Bonita R. A multinational comparison of subarachnoid hemorrhage epidemiology in the WHO MONICA stroke study. *Stroke* 2000;**31**:1054–61.
3. Kassell NF, Sasaki T, Colohan ART, Nazar G. Cerebral vasospasm following aneurysmal subarachnoid hemorrhage. *Stroke* 1985;**16**(4):562–72.
4. Gui JP. Observation of effects of salvia injection in the treatment of acute subarachnoid haemorrhage. *Modern Journal of Integrated Traditional Chinese and Western Medicine* 2004;**13**(6):768.
5. Oliveira-Filho J, Ezzeddine MA, Segal AZ, Buonanno FS, Chang Y, Ogilvy CS, et al. Fever in subarachnoid hemorrhage: relationship to vasospasm and outcome. *Neurology* 2001;**56**:1299–304.
6. Commichau C, Scarmeas N, Mayer SA. Risk factors for fever in the neurologic intensive care unit. *Neurology* 2003;**60**:837–41.
7. Gaetani P, Pasqualin A, Baena RR, Borasio E, Marzatico F. Oxidative stress in the human brain after subarachnoid hemorrhage. *Journal of Neurosurgery* 1998;**89**:748–54.
8. Marik PE. Fever in the ICU. *Chest* 2000;**117**(3):855–69.
9. Cunha BA. Fever in the critical care unit. *Critical Care Clinics* 1998;**14**(1):1–14.
10. Kassell NF, Torner JC, Haley EC, Jane JA, Adams HP, Kongable GL, et al. The international cooperative study on the timing of aneurysm surgery. Part 1: Overall management results. *Journal of Neurosurgery* 1990;**73**:18–36.
11. De Simoni MG, Sironi M, De Luigi A, Manfredi A, Mantovani A, Ghezzi P. Intracerebroventricular injection of interleukin 1 induces high circulating levels of interleukin 6. *Journal of Experimental Medicine* 1990;**171**:1773–8.
12. Fan L, Young PR, Barone FC, Feuerstein GZ, Smith DH, McIntosh TK. Experimental brain injury induces expression of interleukin-1 β mRNA in the rat brain. *Molecular Brain Research* 1995;**30**:125–30.
13. Holmin S, Schalling M, Höjeberg B, Nordqvist ACS, Skeftruna AK, Mathiesen T. Delayed cytokine expression in rat brain following experimental contusion. *Journal of Neurosurgery* 1997;**86**:493–504.
14. Morganti-Kossmann MC, Lenzlinger PM, Hans V, Stahel P, Csuka E, Ammann E, et al. Production of cytokines following brain injury: beneficial and deleterious for the damaged tissue. *Molecular Psychiatry* 1997;**2**:133–6.
15. Shohami E, Ginis I, Hallenbeck JM. Dual role of tumor necrosis factor alpha in brain injury. *Cytokine & Growth Factor Reviews* 1999;**10**:119–30.
16. Shohami E, Novikov M, Bass R, Yamin A, Gallily R. Closed head injury triggers early production of TNF α and IL-6 by brain tissue. *Journal of Cerebral Blood Flow and Metabolism* 1994;**14**:616–9.
17. Bell MJ, Kochanek PM, Doughty LA, Carcillo JA, Adelson PD, Clark RSB, et al. Interleukin-6 and interleukin-10 in cerebrospinal fluid after severe traumatic brain injury in children. *Journal of Neurotrauma* 1997;**14**(7):451–7.
18. Boutin H, LeFeuvre RA, Horai R, Asano M, Iwakura Y, Rothwell NJ. Role of IL-1 α and IL-1 β in ischemic brain damage. *The Journal of Neuroscience* 2001;**21**(15):5528–34.
19. Friedlander RM, Gagliardini V, Rotello RJ, Yuan J. Functional role of interleukin 1 β (IL-1 β) in IL-1 β -converting enzyme-mediated apoptosis. *Journal of Experimental Medicine* 1996;**184**:717–24.
20. Holmin S, Mathiesen T. Intracerebral administration of interleukin-1 β and induction of inflammation, apoptosis, and vasogenic edema. *Journal of Neurosurgery* 2000;**92**:108–20.
21. Bevilacqua MP, Pober JS, Wheeler ME, Cotran RS, Gimbrone MA. Interleukin 1 acts on cultured human vascular endothelium to increase the adhesion of polymorphonuclear leukocytes, monocytes, and related leukocyte cell lines. *Journal of Clinical Investigation* 1985;**76**:2003–11.
22. Quagliarello VJ, Wispelwey B, Long WJ, Scheld WM. Recombinant human interleukin-1 induces meningitis and blood–brain barrier injury in the rat: characterization and comparison with tumor necrosis factor. *Journal of Clinical Investigation* 1991;**87**:1360–6.
23. Yamasaki Y, Matsuura N, Shozuhara H, Onodera H, Itoyama Y, Kogure K. Interleukin-1 as a pathogenetic mediator of ischemic brain damage in rats. *Stroke* 1995;**26**:676–81.
24. Giulian D, Woodward J, Young DG, Krebs JF, Lachman LB. Interleukin-1 injected into mammalian brain stimulates astrogliosis and neovascularization. *The Journal of Neuroscience* 1988;**8**(7):2485–90.
25. De Simoni MG, De Luigi A, Gemma L, Sironi M, Manfredi A, Ghezzi P. Modulation of systemic interleukin-6 induction by central interleukin-1. *American Journal of Physiology* 1993;**265**:R739–42.
26. Hsieh CL, Chen CL, Tang NY, Chuang CM, Hsieh CT, Chiang SY, et al. *Gastrodia elata* BL mediates the suppression of nNOS and microglia activation to protect against neuronal damage in kainic acid-treated rats. *The American Journal of Chinese Medicine* 2005;**33**(4):599–611.
27. Huang YH, Chang YM, Shen JJ, Hsieh CL. Relationship between the anticonvulsion effect of *Gastrodia elata* and interleukin-1 β , tumor necrosis factor- α and nitric oxide. *Mid-Taiwan Journal of Medicine* 2005;(Suppl. 10):S1–8.
28. Hsieh CL, Cheng CY, Tsai TH, Kin IH, Liu CH, Chiang SY, et al. Paeonol reduced cerebral infarction involving the superoxide anion and microglia activation in ischemia-reperfusion injured rats. *Journal of Ethnopharmacology* 2006;**106**:208–15.
29. Hunt WE, Hess RM. Surgical risk as related to time of intervention in the repair of intracranial aneurysms. *Journal of Neurosurgery* 1968;**28**(1):14–20.
30. Bederson JB, Connolly Jr ES, Batjer HH, Dacey RG, Dion JE, Diringer MN, et al. Guidelines for the management of aneurysmal subarachnoid hemorrhage: a statement for healthcare professionals from a special writing group of the Stroke Council, American Heart Association. *Stroke* 2009;**40**:994.
31. Jennett B, Bond M. Assessment of outcome after severe brain damage. *Lancet* 1975;**1**(7905):480–4.
32. Hsieh CL, Chang CH, Chiang SY, Li TC, Tang NY, Pon CZ, et al. Anticonvulsive and free radical scavenging activities of vanillyl alcohol in ferric chloride-induced epileptic seizures in Sprague–Dawley rats. *Life Sciences* 2000;**67**:1185–95.
33. Hsieh CL, Chiang SY, Cheng KS, Lin YH, Tang NY, Lee CJ, et al. Anticonvulsive and free radical scavenging activities of *Gastrodia elata* BL in kainic acid-treated rats. *The American Journal of Chinese Medicine* 2001;**29**(2):331–41.
34. Luo Y, Qin Z, Hong Z, Zhang X, Ding D, Fu JH, et al. Astragaloside IV protects against ischemic brain injury in a murine model of transient focal ischemia. *Neuroscience Letters* 2004;**363**:218–23.
35. Wu X, Huang Y. Review of experimental research on Calamus chemical components functional mechanism on CNS. *Journal of Zhejiang University of Traditional Chinese Medicine* 2007;**31**(6):789–91.
36. Ji H, Wang L, Bi H, Sun L, Cai B, Wang Y, et al. Mechanisms of lumbrokinase in protection of cerebral ischemia. *European Journal of Pharmacology* 2008;**590**:281–9.
37. Chen HQ, Jin ZY, Wang XJ, Xu XM, Deng L, Zhao JW. Luteolin protects dopaminergic neurons from inflammation-induced injury through inhibition of microglial activation. *Neuroscience Letters* 2008;**448**:175–9.
38. Chen CY, Peng WH, Tsai KD, Hsu SL. Luteolin suppresses inflammation-associated gene expression by blocking NF- κ B and AP-1 activation pathway in mouse alveolar macrophage. *Life Sciences* 2007;**81**:1602–14.
39. Tian YY, Jiang B, An LJ, Bao YM. Neuroprotective effect of catalpol against MPP⁺-induced oxidative stress in mes-

- encephalic neurons. *European Journal of Pharmacology* 2007;**568**:142–8.
40. Bi J, Jiang B, Liu JH, Lei C, Zhang XL, An LJ. Protective effects of catalpol against H₂O₂-induced oxidative stress in astrocytes primary cultures. *Neuroscience Letters* 2008;**442**:224–7.
41. Li BQ, Fu T, Gong WH, Dunlop N, Kung HF, Yan Y, et al. The flavonoid baicalin exhibits anti-inflammatory activity by binding to chemokines. *Immunopharmacology* 2000;**49**: 295–306.
42. He XL, Wang YH, Gao M, Li XX, Zhang TT, Du GH. Baicalein protects rat brain mitochondria against chronic cerebral hypoperfusion-induced oxidative damage. *Brain Research* 2009;**1249**:212–21.
43. Huang Q, Shen HM, Ong CN. Inhibitory effect of emodin on tumor invasion through suppression of activator protein-1 and nuclear factor- κ B. *Biochemical Pharmacology* 2004;**68**:361–71.
44. Su MS, Shyu YT, Chien PJ. Antioxidant activities of citrus herbal product extracts. *Food Chemistry* 2008;**111**:892–6.

Reproduced with permission of the copyright owner. Further reproduction prohibited without permission.