Effect of Shengbai Decoction on Chemotherapy-Induced Myelosuppression and Survival of Gastric Cancer Patients After Radical Resection: A Retrospective Study

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Background: Myelosuppression is one of the most common chemotherapy-induced adverse events and results in a series of clinical symptoms. This study aimed to evaluate the effect of Shengbai decoction (SD) on chemotherapy-induced myelosuppression and survival of gastric cancer (GC) patients after radical resection.

Material/Methods: We retrospectively analyzed data from 115 patients with stage II-III GC who underwent adjuvant chemotherapy after radical resection between May 2015 and June 2017 in our hospital. Among these patients, 57 received Shengbai decoction along with adjuvant chemotherapy (SD group), while 58 received adjuvant chemotherapy alone (control group). Medical records, including adverse events, the treatment completion rate of adjuvant chemotherapy, 3-year overall survival (OS), and 3-year recurrence-free survival (RFS), were compared.

Results: Patient characteristics did not differ significantly between the 2 groups. No adverse events related to Shengbai decoction were reported in the SD group. Patients in the SD group had less neutropenia (P=0.0430), thrombocytopenia (P=0.0323), and anemia (P=0.0497). The SD group had a significantly lower probability of dose reduction (P=0.0448). The completion rate of adjuvant chemotherapy of the SD group was considerably higher than that of the control group (P=0.0398). The SD group had a significantly better 3-year RFS (P=0.0369) and 3-year OS (P=0.0455) than the control group.

Conclusions: Shengbai decoction effectively improved postoperative survival of patients with GC by alleviating chemotherapy-induced myelosuppression and improving the completion rate of adjuvant chemotherapy.

Keywords: Medicine, Chinese Traditional • Survival • Chemotherapy, Adjuvant

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**Background**

Gastric cancer (GC), one of the most prevalent malignancies, caused nearly 770,000 deaths in 2020 [1]. D2 gastrectomy is considered a standardized surgical strategy for localized GC in Asian countries [2]. It is also widely proposed by treatment guidelines of Western countries due to the 15-year findings of the Dutch D1/D2 study [3-5]. Nevertheless, approximately 40% of patients develop recurrence within 2 years despite radical resection [6-8]. Various adjuvant chemotherapy regimens have been implemented to control postoperative relapse and improve long-term survival over the past 40 years [9].

However, all the chemotherapy regimens cause toxic adverse events, which harm patient quality of life and treatment compliance [10]. Myelosuppression is one of the most common chemotherapy-induced adverse events and results in a series of clinical symptoms, such as anemia, thrombocytopenia, and neutropenia [11]. Severe myelosuppression even leads to treatment dose reduction or discontinuation, which limit treatment efficacy.

Chinese herbal medicine (CHM) has been widely used in cancer treatment as integrative therapy [12]. Several studies revealed that CHM can prevent chemotherapy-induced myelosuppression [13-15]. However, the efficacy of CHM varies greatly with the composition [16]. This retrospective study was performed to assess the effect of Shengbai decoction on chemotherapy-induced myelosuppression and survival of GC patients after radical resection.

**Material and Methods**

**Patients**

A total of 115 patients with stage II-III GC who underwent adjuvant chemotherapy after radical resection between May 2015 and June 2017 in our hospital were retrospectively analyzed. Among these patients, 57 received Shengbai decoction along with adjuvant chemotherapy (SD group), and 58 received adjuvant chemotherapy alone (control group).

Inclusion criteria were: undergoing adjuvant chemotherapy after D2 gastrectomy, pathological TNM stage II-III GC [17], ECOG 0-1, age 18-75 years, no previous anti-cancer treatment, adequate organ function, and complete follow-up data. Exclusion criteria were: allergic to any ingredient of the Shengbai decoction, recurrence or death within 6 months after surgery, a history of other malignant tumors, and lost to follow-up.

We conducted this study following the principles of the Declaration of Helsinki and “Good Clinical Practice” guidelines. The Institutional Review Board of the college approved this study (approval no. HZYY-2020100811). All patients signed the informed consent.

**Administration of Adjuvant Chemotherapy and Shengbai Decoction**

Postoperative chemotherapy was started within 3 weeks after surgery. The regimens of chemotherapy referred to the NCCN guidelines for GC. Patients decided by themselves whether to receive Shengbai decoction after doctors clarified the potential risks and benefits. Patients signed an additional consent form if they chose to receive Shengbai decoction. The cost of the Shengbai decoction was covered by medical insurance.

The Shengbai decoction was administered concurrently with the chemotherapy to the patients of the SD group (taken morning and evening, 30 min after meals, 100 mL per time). The formula of Shengbai decoction consisted of *Radix Astragali 10 g, Radix Salviae Miltiorrhiza 10 g, Ganoderma Lucidumseu Sinensis 6 g, Rhizoma Atractylodis Macrocephalae 10 g, Radix Angelicae Sinensis 10 g, Caulis Spatholobi 15 g, Fructus Ligustri Lucidi 10 g, Rhizoma Polygonati 10 g, Fructus Psoraleae 10 g and Herba Dendrobii 10 g* (Table 1).

Treatment-related adverse events were estimated based on the Common Terminology Criteria for Adverse Events (CTCAE v4.0). The treatment dose would be reduced to 75% in subsequent treatment courses in case of grade 3-4 hematologic or acute non-hematologic adverse events or grade 2-3 hand-foot syndrome (HFS). Treatments for myelosuppression were not implemented until grade 3 or worse hematologic adverse events occurred. Chemotherapy was discontinued in the event of disease progression or life-threatening adverse events, or patients’ request for discontinuation. Patients who received fewer than half of the scheduled cycles of chemotherapy were excluded from analysis.

**Assessment and Follow-up**

All patients underwent comprehensive disease assessment and health monitoring during the chemotherapy phase and the follow-up phase. Follow-up visits were initiated after the chemotherapy ended. In the first postoperative year, patients underwent monthly follow-up visits and then every 3 months until death or last follow-up.

Recurrence was diagnosed by imaging examination and, if necessary, cytologic analysis or biopsy. Once recurrence was identified, chemotherapy, radiofrequency ablation, or palliative care were implemented.
### Table 1. Formula of Shengbai decoction.

| Name in Chinese | Name in English       | Name in Latin                          | Dose |
|-----------------|-----------------------|----------------------------------------|------|
| Huang Qi        | Milkvetch Root        | Radix Astragali                        | 10 g |
| Dan Shen        | Danshen Root          | Radix Salviae Militiorrhizae           | 10 g |
| Ling Zhi        | Lucid Ganoderma       | Ganoderma Lucidumseu Sinensis         | 6 g  |
| Bai Shu         | Largehead Atractylodes Rhizome | Rhizoma Atractylodis Macrocephalae | 10 g |
| Dang Gui        | Chinese Angelica      | Radix Angelicae Sinensis               | 10 g |
| Ji Xue Teng     | Suberect Spatholobus Stem | Caulis Spatholobi              | 15 g |
| Nv Zheng Zi     | Glossy Privet Fruit   | Fructus Ligustri Lucidi               | 10 g |
| Huang Jing      | Siberian Solomonseal Rhizome | Rhizoma Polygonati     | 10 g |
| Bu Gu Zhi       | Malaytea Scurfpea Fruit | Fructus Psoraleae          | 10 g |
| Shi Hu          | Dendrobium            | Herba Dendrobii                     | 10 g |

### Table 2. Patient characteristics.

|                        | Control group (n=58) | SD group (n=57) | P value |
|------------------------|----------------------|-----------------|---------|
| Age (year)             | 55.24±6.28           | 54.36±6.12      | 0.4483  |
| Tumor size (cm)        | 3.57±1.13            | 3.61±1.25       | 0.8574  |
| Gender                 |                      |                 |         |
| Male                   | 39                   | 40              | 0.7344  |
| Female                 | 19                   | 17              |         |
| Tumor Stage            |                      |                 |         |
| II                     | 26                   | 24              | 0.7694  |
| III                    | 32                   | 33              |         |
| Tumor differentiation  |                      |                 |         |
| Well                   | 8                    | 9               | 0.9284  |
| Moderately             | 18                   | 16              |         |
| Poorly                 | 27                   | 26              |         |
| Signet ring cell       | 5                    | 6               |         |
| Tumor location         |                      |                 | 0.9642  |
| Lower                  | 28                   | 27              |         |
| Middle                 | 12                   | 15              |         |
| Upper                  | 14                   | 12              |         |
| Entire                 | 4                    | 3               |         |
| Regimen of chemotherapy|                      |                 | 0.8550  |
| SOX                    | 18                   | 12              |         |
| CapOX                  | 22                   | 23              |         |
| FOLFOX4                | 18                   | 19              |         |
| Underlying diseases    |                      |                 |         |
| Cardiovascular disease | 5                    | 7               | 0.5209  |
| Respiratory disease    | 6                    | 8               | 0.5451  |
| Diabetes               | 7                    | 7               | 0.9723  |
Statistical analysis and visualization were performed using MedCalc software (version 15.2.2, MedCalc Software, Ltd). Quantitative variables were analyzed using the *t* test and are presented as mean±standard deviation. The *χ²* test or Fisher’s exact test was used to analyze enumeration variables. Ranked data such as treatment-related adverse events, tumor stage, and tumor differentiation were analyzed by Ridit analysis. Recurrence-free survival (RFS) was calculated from the gastrectomy to (i) first recurrence, (ii) last follow-up, or (iii) death from any cause. Overall survival (OS) was calculated from the gastrectomy to death from any cause or the last follow-up. Survival curves were obtained via the Kaplan-Meier method and compared by the log-rank test. A *P* value less than 0.05 was considered to be statistically significant.

### Results

#### Patient Characteristics

Of these patients, 50 had stage II disease and 65 had stage III disease. Patient characteristics did not differ significantly between the 2 groups (Table 2).

#### Adverse Events and Treatment Completion Rate

Treatment-related adverse events were shown in Table 3. There were no significant differences found in non-hematologic adverse events between the 2 groups. No adverse events related to Shengbai decoction were reported in the SD group. Shengbai decoction significantly relieved chemotherapy-induced myelosuppression. Patients in the Shengbai decoction group had less neutropenia (*P* = 0.0430), thrombocytopenia (*P* = 0.0323), and anemia (*P* = 0.0497). No treatment-related deaths occurred. Most of the adverse events were controlled by symptomatic treatment and dose reduction. Dose reduction was documented in 16 patients from the control group and 7 patients from the SD group. The SD group had a significantly lower probability of dose reduction (*P* = 0.0448). Despite administration of dose reduction and thorough monitoring and symptomatic treatment, chemotherapy was finally discontinued in 9 patients (8 from the control group and 1 from the SD group). The completion rate of adjuvant chemotherapy of the SD group was significantly higher than that of the control group (56/57 vs 50/58, *P* = 0.0398).

#### Survival Outcomes

In 30 patients (20 from the control group and 10 from the SD group), recurrence was reported within the first 3 postoperative years. The 3-year RFS rate was 65.52% and 82.76% for the control group and SD group, respectively. The SD group had a significantly better 3-year RFS than the control group (*P* = 0.0369; hazard ratio (HR) for recurrence, 0.4570; 95% confidence interval (CI), 0.2232 to 0.9355) (Figure 1).
characterized by anemia, thrombocytopenia, and neutropenia. of the most common chemotherapy-induced adverse events, dis-continuation of chemotherapy. Myelosuppression is one reported, and overwhelming adverse events can even lead to is consistently implemented when severe adverse events are quality of life and low treatment compliance. Dose reduction GC but also causes various adverse events, which result in poor Postoperative chemotherapy improves the survival of advanced of surgical technique.

In China, the prognosis of GC in China is poor despite the advancement in tumor biology and surgical technique. The main factors for poor prognosis include advanced stage at diagnosis, presence of high-risk factors, and organ metastases. Patients with advanced GC typically have a poor outlook, and the 5-year survival rate is less than 20%. However, with advances in personalized treatment, the prognosis for advanced GC has improved in recent years. The use of combination chemotherapy, targeted therapies, and immunotherapies has shown promising results in improving survival outcomes.

Discussion

China is one of the countries with the highest incidence rates of GC, and about half of the world’s GC-related deaths occur in this country [1]. Due to the inadequacy of nationwide screening programs for cancer, nearly 80% of GC cases are metastatic or locally advanced at the time of diagnosis [18]. Therefore, the prognosis of GC in China is poor despite the advancement of surgical technique.

Postoperative chemotherapy improves the survival of advanced GC but also causes various adverse events, which result in poor quality of life and low treatment compliance. Dose reduction is consistently implemented when severe adverse events are reported, and overwhelming adverse events can even lead to discontinuation of chemotherapy. Myelosuppression is one of the most common chemotherapy-induced adverse events, characterized by anemia, thrombocytopenia, and neutropenia.

Without appropriate treatment, myelosuppression will cause serious consequences [19]. Currently, the main treatments for chemotherapy-induced myelosuppression include dose adjustment, blood transfusion, and recombinant human stimulating factors. However, these treatments can cause organ injuries and vascular events, and they can even contribute to cancer progression [20,21].

Chinese herbal medicine is one of the treatment strategies for myelosuppression and is widely used in China. The Shengbai decoction used in this study was composed of 10 kinds of herbal plants. In the formula, *Radix Astragali*, *Rhizoma Atractylodis Macrocephalae*, and *Ganoderma Lucidum* mainly invigorate qi for consolidating superficial and strengthen immu-nity supplemented by *Rhizoma Polygonati*. *Radix Angelicae Sinensis* enriches the blood and alleviates anemia by promoting erythropoiesis. *Caulis Spatholobi* stimulates the production of platelets by the bone marrow. *Fructus Ligustri Lucidi* has the effect of promoting neutrophil proliferation. *Radix Astragali* and *Fructus Psoraleae* nourish Yin and Yang to achieve balance. *Radix Salviae Miltiorrhizae* improves blood circulation and can acti-vate hematopoietic stem cells. Those components were rati-o-nally assembled and integrated into SD for enhanced functions.

Our results showed that the incidence and severity of myelo-suppression were significantly reduced by Shengbai decoction. Patients in the SD group experienced significantly less neutro-penia (*P*=0.0430), thrombocytopenia (*P*=0.0323), and anemia
Our study has several limitations. First, bias in patient selection was inevitable for this non-randomized and retrospective study. Secondly, this study was performed at a single medical center in China with a limited sample size. Therefore, we are preparing a random clinical trial with a larger sample size to confirm the results of this study. Moreover, further basic research is needed to elucidate the mechanism underlying the therapeutic effects of Shengbai decoction on chemotherapy-induced myelosuppression.

Conclusions

Our results suggest that Shengbai decoction can effectively improve postoperative survival of patients with GC by alleviating chemotherapy-induced myelosuppression and improving the completion rate of adjuvant chemotherapy.

Declaration of Figures’ Authenticity

All figures submitted have been created by the authors, who confirm that the images are original with no duplication and have not been previously published in whole or in part.

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