

# Using Pegaspargase in Combination with Chemotherapy in the Treatment of Lymphoma

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**Abstract:** *Objective:* To analyze the clinical effect of pegaspargase combined with chemotherapy on patients with lymphoma. *Methods:* Seventy patients with lymphoma admitted to Shaanxi Provincial People's Hospital between December 2020 and June 2021 were selected as study subjects and were equally divided into the control group and the intervention group using the lottery method, with 35 cases in each group; the control group received conventional treatment, while the intervention group received pegaspargase combined with chemotherapy. The treatment satisfaction, quality of life, psychological status, and incidence of adverse reactions of the patients in the two groups were compared. *Results:* The differences in the indicators between the two groups were statistically significant ( $p < 0.05$ ). *Conclusion:* Pegaspargase combined with chemotherapy can effectively improve the treatment effect and satisfaction of lymphoma patients; hence, it is worthy of promotion in clinical treatment.

**Keywords:** Pegaspargase combined with chemotherapy; Conventional treatment; Lymphoma

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## 1. Case study

Lymphoma is a malignant tumor originating from the lymphatic system. Lymphomas can be grouped into Hodgkin's lymphoma or non-Hodgkin's lymphoma. Clinically, painless enlargement of lymph nodes is the main feature, and examination would reveal liver and spleen enlargement, along with symptoms of fever and anemia in the late stage. Chemotherapy is the mainstay of treatment in lymphoma, and it can cure lymphoma in some patients [1-4]. Several studies have shown that the use of pegaspargase in combination with chemotherapy is clinically effective in the treatment of lymphoma [5-8]. In this study, 70 patients with lymphoma admitted to Shaanxi Provincial People's Hospital between December 2020 and June 2021 were used as study subjects. The main objective of the study was to investigate the treatment effect of pegaspargase combined with chemotherapy on patients with lymphoma.

## 2. Materials and methods

### 2.1. General information

Seventy patients with lymphoma admitted to Shaanxi Provincial People's Hospital between December 2020 and June 2021 were selected as study subjects and divided into two groups using the lottery method. Thirty-five patients in the control group were treated with conventional treatment, while the other 35 patients in the intervention group were treated with pegaspargase in combination with chemotherapy. There were 19 male patients and 16 female patients in the control group, age ranging from 46 to 78, with a mean age of  $58.85 \pm 1.84$ . In the intervention group, the patients aged 48 to 76, with a mean age of  $59.65 \pm 1.93$ ,

of whom 20 were male and 15 were female. The differences in the general information, including age and gender, between the two groups were not statistically significant ( $p > 0.05$ ) and were comparable. This study was approved by the Ethics Committee of Shaanxi Provincial People's Hospital, and the patients and their families were informed of the study and signed the informed consent form.

## 2.2. Methods

Both groups were treated with oxaliplatin (Qilu Pharmaceutical [Hainan] Co., Ltd., Guodianshi H20093168) at a dose of 130 mg/m<sup>2</sup> in 250–500 ml 5% glucose solution for 2–6 hours; gemcitabine (Jiangsu Haosen Pharmaceutical Group Co., Ltd., Guodianshi H20030104) at a dose of 1000 mg/m<sup>2</sup> intravenously once a week for three weeks, followed by a one-week break, and repeated every four weeks; dexamethasone (Guangdong Nanguo Pharmaceutical Co., Ltd., GMP H44024618) at a dose of 0.75–3.00 mg (1–4 tablets) once for adults, two to four times a day.

- (1) In the control group, L-asparaginase for injection (Kyowa Hakko Kogyo Co., Ltd., H20090520) was administered intravenously at a dose of 50-200 KU per kg body weight on 1 day or every other day. It was increased or decreased as appropriate for age and systemic status.
- (2) The intervention group was treated with pemesterase in combination with chemotherapy: pemesterase (Jiangsu Hengrui Pharmaceutical Co., Ltd., H20153215) 2500 IU/m<sup>2</sup>, administered intramuscularly every 14 days.

## 2.3. Observed indicators

- (1) The patients were surveyed using a self-made satisfaction questionnaire with a total score of 100, with very satisfied being 80-100, satisfied being 60-79, and unsatisfied being 59 or less; satisfaction = (very satisfied + satisfied)/total\*%.
- (2) The Self-Rating Depression Scale (SDS) was used to rate the depression of the two groups, with 20 items in total, 4 points each, and a cut-off score of 53. The Self-Rating Anxiety Scale (SAS) was used to rate the anxiety of patients, with 20 items in total, 4 points each, and a cut-off score of 50, above which anxiety was indicated; the lower the score, the better a patient's psychological state.
- (3) The Generic Quality of Life Inventory-74 (GQOL-74) was used to assess the quality of life of both groups, including physical, psychological, somatic, and social aspects; the total scores were compared.
- (4) The incidence of adverse reactions, including hyperglycemia, gastrointestinal reactions, and granulocytopenia, was compared between the two groups.

## 2.4. Statistical analysis

SPSS 20.0 was used to process the data; t-test was performed on the data obtained and expressed as *s*, while X<sup>2</sup> test was performed on the count data and expressed as %.  $p < 0.05$  was considered statistically significant.

## 3. Results

### 3.1. Patient satisfaction

The difference in patient satisfaction between the intervention and control groups was statistically significant ( $p < 0.5$ ), as shown in **Table 1**.

**Table 1.** Comparison of patient satisfaction between the two groups (n/%)

Group	Number of cases	Very satisfied	Satisfied	Unsatisfied	Satisfaction
Intervention group	35	34 (97.14)	0 (0.00)	1 (2.86)	34 (97.14)
Control group	35	21 (60.00)	7 (20.00)	7 (20.00)	28 (80.00)
X <sup>2</sup>					5.0806
<i>p</i>					0.0242

### 3.2. Mental state

There was no significant difference in the SAS and SDS scores between the two groups before treatment ( $p > 0.05$ ), and they both improved significantly after treatment ( $p < 0.05$ ). The data comparison between the two groups showed that the SAS and SDS scores of patients in the intervention group were significantly lower than those in the control group ( $p < 0.05$ ), and the difference between the groups was statistically significant (**Table 2**).

**Table 2.** Comparison of SAS and SDS scores before and after treatment (n = 35, ± s)

Group	Number of cases	SAS		SDS	
		Pre-treatment	Post-treatment	Pre-treatment	Post-treatment
Control group	35	50.55 ± 2.26	43.63 ± 0.57	49.35 ± 2.65	42.26 ± 0.03
Intervention group	35	50.35 ± 1.58	30.42 ± 0.18	49.35 ± 2.52	33.67 ± 0.53
t		0.4291	130.7436	0.0000	95.7319
p		0.6692	0.0000	1.0000	0.0000

### 3.3. Quality of life

Before treatment, there was no significant difference in the physical, psychological, somatic, and social scores between the two groups ( $p > 0.05$ ); however, after treatment, the physical, psychological, somatic, and social scores of the patients in the intervention group were significantly higher than those of the control group ( $p < 0.05$ ), and the difference was statistically significant (**Table 3**).

**Table 3.** Comparison of quality-of-life scores before and after treatment (n = 35, ± s)

Group	Number of cases	Physical life		Somatic functions	
		Pre-treatment	Post-treatment	Pre-treatment	Post-treatment
Control group	35	83.55 ± 2.26	87.63 ± 2.57	81.35 ± 3.65	86.26 ± 3.03
Intervention group	35	83.35 ± 2.58	90.42 ± 2.18	80.35 ± 2.52	91.67 ± 2.53
t		0.3450	4.8978	1.3338	8.1082
p		0.7312	0.0000	0.1867	0.0000
Group	Number of cases	Social functions		Psychological functions	
		Pre-treatment	Post-treatment	Pre-treatment	Post-treatment
Control group	35	80.55 ± 3.26	87.63 ± 2.57	79.35 ± 2.65	86.26 ± 2.03
Intervention group	35	81.35 ± 2.58	91.42 ± 2.18	80.35 ± 3.52	91.67 ± 2.53
t		1.1384	6.6533	1.3427	9.8670
p		0.2589	0.0000	0.1838	0.0000

### 3.4. Incidence of adverse reactions

The incidence of hyperglycemia, gastrointestinal reactions, and granulocytopenia was significantly lower in the intervention group than in the control group ( $p < 0.05$ ), as shown in **Table 4**.

**Table 4.** Comparison of adverse reactions in the two groups (n, %)

Group	Number of cases	Hyperglycemia	Gastrointestinal reactions	Granulocytopenia	Incidence rate
Intervention group	35	1	0	0	1 (2.86)
Control group	35	3	1	2	6 (17.14)
X <sup>2</sup>					3.9683
<i>p</i>					0.0464

#### 4. Discussion

Lymphoma can be caused by a number of factors. Most patients are affected by viral infections, mainly related to EBV infection, which is one of the main causes of Hodgkin's lymphoma [9-14]; others are affected by retroviral infections, which can cause lymphoma as well as t-cell leukemia. There is also a strong relationship between lymphoma and individual immune factors, with a higher incidence of disease in cases of immune deficiencies [15,16]. The systemic symptoms of malignant lymphoma vary widely depending on the type of disease and the time of onset. Some patients may have no systemic symptoms [17-19]. Fever, weight loss (more than 10% weight loss), and night sweats are the most common symptoms, followed by loss of appetite, fatigue, and pruritus. Systemic symptoms are related to age of onset, extent of the tumor, and the body's immunity. Systemic symptoms are significant in elderly patients, those who are immunocompromised, or patients with multiple focal attacks [20]. The survival rate of patients without systemic symptoms is three times higher than that of patients with symptoms. In terms of management, surgery is superior to conventional chemotherapy, as the latter has significant side effects and can be very disruptive or damaging. In the current study, pegaspargase combined with chemotherapy was found effective in treating lymphoma.

In conclusion, by using pegaspargase combined with chemotherapy for lymphoma patients, it can effectively improve the treatment effect and psychological state of patients; thus, it is worthy of promotion in clinical practice.

#### Disclosure statement

The authors declare no conflict of interest.

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