Abstract. The present study aimed to investigate the curative effect of high-dose methotrexate (HD-MTX) combined with teniposide (Vm26) vs HD-MTX alone in the treatment of primary central nervous system lymphoma (PCNSL), in order to provide data for assisting decisions associated with clinical treatment. Data from 56 patients with PCNSL admitted in Shanghai Huashan Hospital (Shanghai, China) from January 2009 to December 2014 were included into the present study. Clinical data, curative effects and prognosis of patients in these two groups were retrospectively analyzed using SPSS 20 statistical software. In the HD-MTX+Vm26 group, 12 patients (42.85%) achieved complete remission (CR) and 10 patients (35.71%) achieved partial remission (PR), while in the HD-MTX group 7 patients (25%) achieved CR and 11 patients (39.29%) achieved PR (P=0.158). The median progression-free survival (PFS) time was 22 months in the HD-MTX+Vm26 group and 12 months in the HD-MTX group (P=0.019). The median overall survival time was 57 months in the HD-MTX+Vm26 group, and 28 months in the HD-MTX group (P=0.019). Compared with HD-MTX alone, the combined treatment of HD-MTX+Vm26 had an improved curative effect in the treatment of PCNSL, effectively controlled tumor progression in patients, prolonged survival time and improved prognosis. Age was an independent prognostic factor in patients with PCNSL. Patients with an age of ≤60 years exhibited longer PFS compared with patients with an age of >60 years.

Introduction

Primary central nervous system lymphoma (PCNSL) is a malignant tumor confined to the brain, pia mater, spinal cord or eye (1). A study at the Massachusetts General Hospital from 1958 to 1989 showed that a total of >90% of patients with PCNSL have diffuse large B cell lymphoma (2), which exhibits high invasion rates and poor prognosis. Poor prognostic factors include age, Karnofsky score, serum lactate dehydrogenase (LDH), cerebrospinal fluid protein, B cell lymphoma 6 protein expression (3) and reduced lymphocytes (4).

PCNSL cells usually present with diffuse infiltration. Therefore, it is difficult to completely remove by surgery; and its recurrence rate is high (5). Previously, whole brain radiotherapy was performed as a first-line therapy; however, treatment with radiotherapy alone confers disease recurrence and neurotoxicity (6,7), which limits its clinical application. High-dose methotrexate (HD-MTX) has an affirmed curative effect, but the complications, such as neurotoxicity, limit its dose (8,9). In addition, the 2-year overall survival (OS) rate of HD-MTX monotherapy is only 61-63% (10,11), which is not satisfying. At present, the preferred regimen is the combined treatment based on HD-MTX (combined with cytarabine, rituximab and idarubicin) (12-18). However, the combination of HD-MTX based chemotherapy is still under debate. Teniposide (Vm26) has high liposolubility, which makes it is able to pass the blood-brain barrier, and it has a slower elimination rate (19). Fan et al (20) Revealed that MTX+Vm26 did not significantly improve the median progression‑free survival of patients, but complete remission (CR) and overall response (OR) were significantly increased. In the present study, the clinical curative effects of HD-MTX+Vm26 vs. Vm26 alone were compared, with the aim of providing a useful reference for clinical practice.

Materials and methods

Data acquisition. A total of 56 patients with PCNSL who were admitted, but had not been treated, to the Department of Hematology of Huashan Hospital Affiliated to Fudan University (Shanghai, China) from January 2009 to December 2014 were enrolled into the present study. Due to the retrospective nature of the present study, the requirement for ethical approval was waived by the Ethics Committee of Huashan Hospital (Shanghai, China). The cohort of 56 patients comprised 38 males and 18 females. The age of these patients...
ranged between 25-74 years, with a median age of 54.5 years. A total of 13 patients (23.21%) were ≥60 years.

According to the treatment regimens received, patients were divided into two groups: A HD-MTX+Vm26 group and HD-MTX group (n = 28 in each group). A retrospective analysis method was adopted. Inclusion criteria were as follows: i) Patients who were first diagnosed with PCNSL; ii) patients with a negative HIV test result; iii) patients with diagnosis supported by pathological evidences; iv) patients who had completed at least three courses of chemotherapy; v) patients with changes in intracranial lesions that had been followed up by imaging tests (MRI or PET-CT) prior to diagnosis and subsequent to chemotherapy; and vi) patients who were diagnosed based on the 2008 World Health Organization classification of hematopoietic and lymphoid neoplasms (21). Exclusion criteria were as follows: Congenital or acquired immunodeficiency, including patients with previous organ transplantation, concurrent treatment with immunosuppressive drugs, and AIDS-related PCNSL; disease confined to the eye without another localization in the central nervous system (CNS); the presence or history of systemic lymphoma; any prior malignancy with the exception of adequately treated nonmelanoma skin cancer and carcinoma in situ of the cervix uteri; a serious impairment of cardiac, renal, or liver function; pregnancy; any severe uncontrolled infection; prior chemotherapy, with the exception of corticosteroids, for a maximum period of 6 weeks before and after diagnosis or surgery; and Burkitt's lymphomas of low-grade T-cell lymphomas.

Tissue and reagents. 56 cases of excised tissue or biopsy tissue of brain tumor were collected, fixed by 3.7% neutral formaldehyde at room temperature for 6–12 h, until further processing, routinely dehydrated, embedded by paraffin, sliced into 3-µm thick sections, and then subjected to H&E staining (stained with Gill's hematoxylin for 2-3 min and eosin for 10-20 sec at room temperature) and immunohistochemical staining. In immunohistochemical staining, the tissue was embedded by paraffin, sliced into 4-µm thick sections, dewaxed with 100% xylene and 95% EtOH for 4 min twice, 80% EtOH for 4 min) according to concentration gradients of alcohol (100% EtOH for 5 min twice, 95% EtOH for 4 min twice, 80% EtOH for 4 min) according to the standard operation process, and repaired in citrate buffer (pH 6.0; Shanghai Xin Yu Biotech Co., Ltd., Shanghai, China) and boiled for 10 min. After natural cooling, the endogenous peroxidase activity was blocked using 3% hydrogen peroxide at room temperature for 10 min, and then treated with the primary antibodies and incubated at 4˚C overnight: CD10 (1:50; cat. no. ab951), CD19 (1:200; cat. no. ab108403), CD3 (1:30; cat. no. ab16669), Bcl-2 (undiluted; cat. no. ab172610) and Bcl-2 (1:100; cat. no. ab32124), which were purchased from Abcam; CD38 (1:200; cat. no. or677046), which were purchased from Zymed; TDT (undiluted; cat. no. PL0403179) and CyclinD1 (1:50; cat. no. or677046), which were purchased from Neomarkers (https://univ.biomart.cn/). After washing with PBS, the tissue sections were placed to react with biotin-labeled rabbit anti-guinea pig IgG secondary antibody (1:1000; cat. no. QN2726; Beijing Baiaoleboke Co., Ltd., Beijing, China) at 37˚C for 30 min. The sections were washed with PBS for 5 min, repeated three times, then incubated with VECTA STAIN Elite ABC standard kit (Vector Laboratories, Inc., Burlingame, CA, USA) according to the manufacturer's protocol at 37˚C for 30 min, finally the signals were visualized with a chromogenic kit (Vector Laboratories, Inc.) according to the manufacturer's protocol or by microscopic examination: Optical microscope (Olympus BX41, Olympus Corporation, Tokyo, Japan) was used, and x4, x10 and x40 objectives were used for observation, respectively.

Statistical analysis. The collected data were analyzed using SPSS 20.0 software (IBM Corp., Armonk, NY, USA). Overall survival (OS) time was defined as the period from the time the patient was diagnosed to the time the patient succumbed, or up to the follow-up deadline. Progression-free survival (PFS) was defined as the period from the time the patient was diagnosed to disease recurrence/progression or last point of contact with the patient. Uniivariate analysis was conducted using Kaplan-Meier curves and log-rank test and multivariate analysis was conducted using a Cox's regression. P<0.05 was considered to indicate a statistically significant difference.

Results

Patients. The cohort of 56 patients comprised 38 males and 18 females. The age of these patients ranged between
Clinical characteristics of patients

Clinical symptoms. The common symptoms of PCNSL include elevated intracranial pressure (53.6%), somatic dysfunction (53.6%), cognitive dysfunction (33.9%), impaired vision (30.3%) and language disorder (17.9%).

Imaging tests. All patients underwent cranial magnetic resonance imaging, and 17 additionally underwent positron emission tomography-computed tomography. Patients with multiple lesions accounted for 60.7% of the total cohort, and patients with deep lesions accounted for 60.7%. The frontal lobe was the most common lesion site, accounting for 46.4% patients, followed by the parietal lobe (25%) and basal ganglia (23.2%).

ECOG score results. A total of 12 patients received 1 point, 13 patients received 2 points, 26 patients received 2 points, 4 patients received 3 points and 2 patients received 5 points. Patients who received a score within 1-3 points accounted for 91% of the total cohort.

Histopathology and laboratory tests

Histopathology. The histopathology of all 56 patients was diffuse large B cell lymphoma.

Cerebrospinal fluid examination. Cerebrospinal fluid examination was performed on 26 patients. Among 26 patients, a Pandy's test exhibited a positive result in 19 patients (73.1%), the protein content increased in 18 patients (69.1%), and spinal cord involvement, such as chest-back or lower-back pain, weakness of lower limbs, paraplegia and superficial sensory dullness or disappearance, occurred in 11 patients (42.3%). Spinal cord involvement: i) The patients had periodical lumbar pain and compression symptoms; ii) Enhanced MRI revealed lesions; iii) The lesions shrank after treatment.

Serum LDH and serum and urinary β2-microglobulin (MG) examination. Serum LDH (normal range from 109 to 245 U/l) was increased in 21 patients (37.5%), serum β2-MG (normal range from 1 to 2.8 µg/ml) was increased in 32 patients (57.1%), and urinary β2-MG (normal range from 0 to 0.3 µg/ml) was increased in 21 patients (37.5%).

Treatment regimen

Treatment regimen. Treatment regimens were divided into three categories: i) Radiotherapy + chemotherapy (radiotherapy followed by chemotherapy); ii) chemotherapy + radiotherapy (chemotherapy followed by radiotherapy); iii) chemotherapy alone. All had been done. 7 patients received radiotherapy + chemotherapy, 20 patients received chemotherapy + radiotherapy, and 29 patients received chemotherapy alone. MTX group: 3 patients received radiotherapy + chemotherapy, 13 patients received chemotherapy + radiotherapy, and 13 patients received chemotherapy alone.

The 56 patients in the present study all received chemotherapy regimens with HD-MTX. Among these patients, 51.8% patients were treated with chemotherapy alone, and the remaining 48.2% patients were treated with chemotherapy and radiotherapy. The majority of the patients received radiotherapy followed by chemotherapy; only 7 patients (12.5%) received chemotherapy followed by radiotherapy. These results revealed that the median PFS was 20 months and the median OS time was 31 months in patients who received chemotherapy alone. The median PFS and median OS were 21 and 37 months, respectively, in patients who received chemotherapy followed by radiotherapy; while median PFS and median OS were 22 and 52 months, respectively, in patients who received radiotherapy followed by chemotherapy. Between patients who received chemotherapy alone and the patients who received chemotherapy combined with radiotherapy, the difference in PFS was not statistically significant.

Curative effect analysis

Treatment response. The evaluation of disease progression was performed following three courses of chemotherapy for all patients. In the MTX+Vm26 group, 12 patients (42.85%) achieved CR, 10 patients (35.71%) achieved PR and the OR rate was 78.57%. In the MTX group, 7 patients (25%) achieved CR, 11 patients (39.29%) achieved PR and the OR rate was 64.3%. Differences in CR and OR between these two groups were not statistically significant (Table II).

PFS and OS. At the end of the follow-up, 26 patients succumbed, 29 patients survived, and 1 patient in the MTX+Vm26 group was lost to follow-up subsequent to confirmation of relapse. The median PFS were 22 and 12 months in the MTX+Vm26 and MTX groups, respectively; the median OS rates were 57 and 28 months, respectively. These two indices were significantly increased in the MTX+Vm26 group compared with the MTX group (P<0.05; Table II). The median OS was 40 months in 55 patients (total patient cohort minus 1 patient lost to follow up) (Fig. 1A). Comparison of the median OS between the two treatment groups revealed that the OS of the MTX+Vm26 group was significantly increased compared with the MTX group (Fig. 1B).

Mortality. Overall, mortality was 47.3% at the end of follow-up, in which 13 patients (47.3%) were from the MTX group and 13 patients (48.1%) were from the MTX+Vm26 group, and the difference between the mortality rates between the two treatment groups was not statistically significant (P=0.898).

Disease progression. A total of 30 patients exhibited disease progression at the end of the follow-up, in which 16 patients (57.14%) were from the MTX group and 14 patients (50%) were from the MTX+Vm26 group; the difference between these two groups was not statistically significant (P=0.592).

Prognosis analysis

Univariate analysis. Univariate analysis was performed to analyze factors that may be associated with the prognosis of disease. The disease progression curve was evaluated using the Kaplan-Meier method and the factors were compared using a log-rank test. Detailed results are indicated in Table III. Univariate analysis revealed that age was a prognostic factor in patients with PCNSL, and the median PFS of patients with an age >60 years was 10 months shorter compared with that in
patients with an age of ≤60 years (Fig. 1C). These are consistent with the results of previous studies (24-26). The median PFS was 69 months in patients who achieved CR following three courses of treatment, and was 20 months in patients who did not achieve CR. The difference between these two was marked, but was not statistically significant (P=0.061). The median PFS was 12 months in the MTX group, and 22 months in the HD-MTX+Vm26 group, and the difference between these two groups was statistically significant (P=0.033; Fig. 1D).

**Discussion**

PCNSL is a rare type of extra-nodal lymphoma. The initial symptoms include changes in cognitive function, increased intracranial pressure and hemiplegia. During the early stage, lesions are confined to the CNS; and in the late stage, <10% of patients exhibit tumors that spread outside the CNS (27). Although the cancer cells originates from lymphocytes, PCNSL usually is considered as a brain tumor in practice due to its therapeutic challenges, such as drug delivery through blood-brain barrier, and cerebral toxicity (28). Due to these unique characteristics of the clinical symptoms and biological behaviors of the tumor cells in PCNSL, the 2008 WHO Hematopoietic and Lymphoid Classification lists it as an independent disease entity (21). In the past three decades, epidemiological data in Western countries, such as Norway and Sweden revealed that the incidence of PCNSL continues to increase (29-31). The prognosis of patients with PCNSL is relatively poor, and the prognosis of patients with HIV

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**Table I. Criteria for assessment of therapeutic effects.**

| Treatment response | Imaging examination | Dosage of glucocorticoid | Ophthalmologic examination | Cerebrospinal fluid examination |
|--------------------|---------------------|--------------------------|----------------------------|--------------------------------|
| CR                 | No enhancement      | Nil                      | Normal                     | Negative                       |
| CRu                | No enhancement      | Any                      | Normal                     | Negative                       |
|                    | No abnormal lesions | Any                      | Micro RPE was abnormal     | Negative                       |
| PR                 | Enhanced lesion was | No association           | Micro RPE was abnormal or  | Negative                       |
|                    | reduced by 50%      |                          | normal                     |                                |
| PD                 | No enhancement      | No association           | Vitreous cell or retinal infiltration, reduction | Persistent positive or suspicious positive |
|                    | Tumor was reduced by 25% or new lesion appeared | | Recurrence or new eye lesions | Recurrence or positive |

CR, complete remission; CRu, unconfirmed complete remission; RPE, retinal pigment epithelial cells; PR, partial remission; PD, disease progression; SD, stable disease, in a state between PR and PD. In the present study, CR included CR and CRu.

**Table II. Comparison of the curative effects between treatments groups.**

| Group              | MTX + Vm26 | MTX   | χ²     | P-value |
|--------------------|------------|-------|--------|---------|
| CR                 | 12 (42.85%)| 7 (25%)| 1.991  | 0.158   |
| PR                 | 10 (35.71%)| 11 (39.29%)| 0.076 | 0.783   |
| SD                 | 4 (14.29%) | 4 (14.29%)| 0.000 | 1.000   |
| PD                 | 2 (7.14%)  | 6 (21.43%)| 1.313 | 0.252   |
| OR                 | 22 (78.57%)| 18 (64.3%)| 1.400 | 0.237   |
| Median PFS, months | 22         | 12    | 5.462  | 0.019   |
| Median OS, months  | 57         | 28    | 6.196  | 0.013   |

CR, complete response; PR, partial remission; SD, stable disease; PD, progressive disease; OR, overall response; PFS, progression-free survival; OS, overall survival; MTX, methotrexate; Vm26, Teniposide.
infection is particularly poor (32). In the present study, no patient had HIV infection.

Statistical data from the United States of America between 2000-2004 revealed that the average age of patients diagnosed with PCNSL was 60 years, and the age of the majority of patients ranged within 50-65 years (5,33,34). A study conducted in Spain revealed that within the period 1990-2010, the average age of patients diagnosed with PCNSL was 59.3 years (35). In the present study, the average age of patients was 53.5 years, which was decreased compared with that in other countries.

PCNSL exhibits three pathological types: DLBCL, T cell lymphoma and other types of B cell lymphoma. DLBCL accounts for >90% of PCNSL (36). The pathological types of the 56 patients were all DLBCL in the present study.

The clinical symptoms of patients with PCNSL were associated with the location of tumor lesions (37,38). A study in Spain revealed that patients with cognitive function decline accounted for 33.4%, and those with cephalgia accounted for 25% (35). The present study revealed that increased intracranial pressure and physical movement disorders were common, and the rates for the two symptoms were 53.6%; followed by cognitive dysfunction (33.9%), vision impairment (30.3%) and language disorder (17.9%).

Statistics from the guidelines of the National Comprehensive Cancer Network (NCCN) revealed that >90% of patients with PCNSL exhibited lesions affecting the brain parenchyma, and over one-half of these patients possessed multiple lesions (39). Fraser et al (40) demonstrated that 35% of patients with PCNSL exhibited multiple lesions, and up to 38% of patients exhibited lesions in the cerebral hemispheres. In the present study, frontal lobe involvement (46.4%) accounted for the largest proportion of lesion sites. The proportion of patients with deep involvement was 60.7%. The majority of patients exhibited multiple lesions (60.7%).

The physical fitness score of patients was one of the factors that affected the choice of treatment regimens. It was not recommended that patients with poor physical condition use the HD-MTX+Vm26 regimen (41). In the present study, the ECOG scores of patients were concentrated within 1-3 points.

Results of the cerebrospinal fluid examination in 26 patients indicated that the Pandy’s test demonstrated a positive result in 19 patients (73.1%), and that the protein level in cerebrospinal fluid was increased in 18 patients (not examined using a Pandy’s test) (69.2%). The examinations of serum LDH, serum β2-MG and urine β2-MG revealed that 37.5% of patients exhibited increased serum LDH, 57.1% of patients exhibited
Table III. Univariate analysis.

| Potential prognostic factors                  | PD/total cases | Median PFS, months | $\chi^2$ | P-value |
|-----------------------------------------------|----------------|--------------------|----------|---------|
| Age, years                                    |                |                    |          |         |
| $\leq$60                                       | 19/43          | 20                 | 6.021    | 0.014   |
| >60                                           | 11/13          | 22                 |          |         |
| ECOG score, points                            |                |                    | 0.017    | 0.896   |
| 0-1                                           | 6/12           | 22                 |          |         |
| $\geq$1                                       | 24/44          | 20                 |          |         |
| Deep lesions accumulating                     |                |                    | 1.117    | 0.291   |
| No                                            | 14/22          | 16                 |          |         |
| Yes                                           | 16/34          | 20                 |          |         |
| Number of lesions                             |                |                    | 0.741    | 0.389   |
| Single                                        | 14/22          | 16                 |          |         |
| Multiple                                      | 16/34          | 20                 |          |         |
| Serum LDH                                     |                |                    | 1.100    | 0.294   |
| Normal (106-211 U/l)                          | 15/35          | 24                 |          |         |
| Abnormally increasing                         | 15/21          | 18                 |          |         |
| Blood $\beta_2$-MG                            |                |                    | 0.793    | 0.373   |
| Normal (0.70-1.80 mg/l)                       | 14/24          | 17                 |          |         |
| Abnormally increasing                         | 16/32          | 22                 |          |         |
| Urine $\beta_2$-MG                            |                |                    | 0.076    | 0.783   |
| Normal ($<0.25$ mg/l)                         | 19/35          | 18                 |          |         |
| Abnormally increasing                         | 11/21          | 21                 |          |         |
| CSF; Pandy's test                             |                |                    | 1.670    | 0.196   |
| Negative                                      | 4/6            | 10                 |          |         |
| Positive                                      | 13/20          | 20                 |          |         |
| CSF protein quantification                    |                |                    | 0.948    | 0.330   |
| Normal (150-450 mg/l)                         | 5/8            | 10                 |          |         |
| Abnormally increasing                         | 12/18          | 20                 |          |         |
| CSF glucose level                             |                |                    | 0.160    | 0.689   |
| Normal (2.5-4.4 mmol/l)                       | 12/20          | 20                 |          |         |
| Abnormal                                      | 5/6            | 20                 |          |         |
| CSF chloride                                  |                |                    | 0.915    | 0.339   |
| Normal (2.7-3.9 mmol/l)                       | 12/14          | 20                 |          |         |
| Abnormal                                      | 5/12           | 22                 |          |         |
| CSF white blood cell count                    |                |                    | 2.296    | 0.130   |
| Normal (0-5/mm$^3$)                           | 12/19          | 24                 |          |         |
| Abnormally increasing                         | 5/7            | 20                 |          |         |
| Lesion resection by surgery                   |                |                    | 0.562    | 0.454   |
| Yes                                           | 12/31          | 20                 |          |         |
| No                                            | 18/25          | 22                 |          |         |
| Received CR                                   |                |                    | 3.519    | 0.061   |
| No                                            | 22/37          | 20                 |          |         |
| Yes                                           | 8/19           | 69                 |          |         |
| Chemotherapy regiments                        |                |                    | 4.557    | 0.033   |
| HD-MTX                                        | 16/28          | 12                 |          |         |
| HD-MTX + Vm26                                 | 14/28          | 22                 |          |         |
| Therapeutic schedule                          |                |                    | 0.920    | 0.631   |
| Chemotherapy                                  | 16/29          | 18                 |          |         |
| Chemotherapy + radiotherapy                   | 4/7            | 21                 |          |         |
| Radiotherapy + chemotherapy                   | 10/20          | 22                 |          |         |

ECOG, Eastern Cooperative Oncology Group; PFS, progression-free survival; LDH, lactate dehydrogenase; $\beta_2$-MG, $\beta_2$-microglobulin; PD, progressive disease; CSF, cerebrospinal fluid; HD-MTX, high-dose methotrexate; Vm26, Teniposide
increased serum β2-MG and 37.5% of patients exhibited increased urinary β2-MG.

The feasibility and effectiveness of HD-MTX in the treatment of PCNSL have been recognized. It may be used for induction therapy and salvage therapy (1). Results of a study conducted by Prica et al (42) revealed that compared with chemotherapy alone, chemotherapy combined with radiotherapy was an improved induction regimen in younger patients (aged <60 years). However, due to toxic side effects, it is recommended that elderly patients (>60 years old) should receive chemotherapy alone (43,44). A study conducted by Muirhead et al (45) revealed that for patients who were not able to tolerate large-dose chemotherapy, 3-year survival rates of salvage radiotherapy were increased compared with that of the second-line chemotherapy (60 vs. 33%). Salvage radiotherapy may be a choice for young patients with recurrent/refractory PCNSL (7). The use of chemotherapy or radiotherapy should be determined on an individual patient basis. The 56 patients in the present study all received chemotherapy regimens with HD-MTX. Among these patients, 51.8% patients were treated with chemotherapy alone, and the remaining 48.2% patients were treated with chemotherapy and radiotherapy. Between patients who received chemotherapy alone and the patients who received chemotherapy combined with radiotherapy, the difference in median OS was notable, but it was not statistically significant (P=0.24). This may be associated with not enough sample size between chemotherapy alone group and chemotherapy combined with radiotherapy group.

Ferreri et al (33) conducted a multicenter retrospective study. Their results revealed 5 factors that were associated with poor prognosis: i) Age >60 years old; ii) ECOG grades 2-4;
have demonstrated that surgical resection could be established in patients with PCNSL were few, and little recent publications recommended in the majority of cases (27). Cases of surgical operation of patients. revealed that this had no significant association with the PFS that affected the deep structures. Furthermore, statistical data patients who demonstrated deep structure involvement was 28%. In the present study, 60.7% of patients exhibited lesions that affected the deep structures. Furthermore, statistical data revealed that this had no significant association with the PFS of patients.

A previous study suggested that surgery did not confer significant benefits to patients with PCNSL, and had risk of damage to neural functions (28). Surgical removal is not recommended in the majority of cases (27). Cases of surgical operation in patients with PCNSL were few, and little recent publications have demonstrated that surgical resection could be established as a prognostic factor in PCNSL (32,37). In 2012, the results of a German PCNSL SG-1 trial revealed that surgical removal may extend the PFS of patients (50-52). Fraser et al (40) considered that surgical removal was advantageous to rapidly relieving symptoms and decreasing glucocorticoid dose, which eliminated the production of drug-resistant tumor clones. In addition, due to modern surgical positioning technology, in 2012 a German primary central nervous system lymphoma (PCNSL) SG-1 test revealed that, surgical resection of PCNSL tumor lesions can prolong the progression free survival time of patients (53,54). The results of the present study revealed that median PFS was 20 months in patients who underwent the surgical removal of lesions, while PFS was 22 months in patients who underwent directional puncture biopsy. The difference was not statistically significant between these two surgical methods.

A previous study identified that patients with multiple lesions exhibited shorter survival times (24), but this was not recognized. An early study revealed that the number of lesions would affect the survival time of patients, and the survival time of patients with a large number of lesions is relatively shorter (42). However, this research result has not been recognized. In the present study, univariate analysis of the number of lesions also revealed that the difference in PFS between patients with multiple lesions and patients with single lesion was not statistically significant.

In the present study, the univariate analysis on the number of lesions revealed that the difference in PFS between patients with multiple lesions and patients with a single lesion was not statistically significant.

In summary, the results of the present study revealed that age and the treatment regimen were independent prognostic factors of PCNSL. Compared with patients >60 years old, patients who were ≥60 years old exhibited a significantly prolonged PFS. At the same time, the PFS of patients who received the combined treatment of Vm25+HD-MTX was significantly prolonged when comparing with HD-MTX alone treatment. The difference in the toxic reaction of chemotherapy was not statistically significant. However, the sample size of the present study was small, and this single-center retrospective study has limitations. Therefore, the conclusions of the present study should be confirmed through prospective multicenter clinical trials in the future.

The clinical data of the 56 patients with PCNSL were retrospectively analyzed. It was revealed that: i) Age is an independent prognostic factor in patients with PCNSL, in which patients who were ≥60 years old exhibited longer PFS compared with patients who were >60 year old; ii) the median PFS time was 69 months in patients who achieved CR following three courses of treatment and 20 months in patients who did not achieve remission, and the difference between these two was marked but not statistically significant (P=0.061); iii) PFS time was 22 months in the HD-MTX+Vm26 group and was 12 months in the MTX group, and median OS time was 57 months in the HD-MTX+Vm26 group and was 28 months in the MTX group. Both PFS time and median OS time were significantly different between these two groups. Therefore, compared with HD-MTX alone, the combined treatment of HD-MTX+Vm26 exhibited a significantly improved long-term curative effect in the treatment of PCNSL.
The authors declare that they have no competing interests.

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Availability of data and materials

The datasets used and/or analyzed during the current study are available from the corresponding author on reasonable request.

Authors' contributions

YXW contributed to the conception and design of the study and drafted the manuscript. YXW, YH, XPX, BBC, ZGL, YM, TLD and QW acquired, analyzed and interpreted the data and revised the manuscript critically for important intellectual content. All authors read and approved the final version of the manuscript.

Ethics approval and consent to participate

This study was conducted in accordance with the declaration of Helsinki. This study was conducted with approval from the Ethics Committee of Huashan Hospital Affiliated to Fudan University. Written informed consent was obtained from the Ethics Committee of Huashan Hospital Affiliated to Fudan University. This study was conducted with approval from the Ethics Committee of Huashan Hospital Affiliated to Fudan University. Written informed consent was obtained from the participants.

Patient consent for publication

Not applicable.

Competing interests

The authors declare that they have no competing interests.

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