

Primary central nervous system lymphoma: initial features, outcome, and late effects in 75 children and adolescents

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Key Points

- Children with PCNSL and no immunodeficiency have a good outcome when treated by a histological subtype-driven and radiation-free protocol.
- New treatment guidelines are needed for PCNSL in children and adolescents with an underlying immunodeficiency.

Introduction

Primary central nervous system lymphoma (PCNSL) is a rare subtype of non-Hodgkin lymphoma (NHL) in childhood and adolescence, with an increased risk among patients with immunodeficiency.¹⁻⁵ Given the rarity and lack of any prospective trials, reliable data on PCNSL are lacking, with most data coming from case reports or small series. Thus, 2 of the largest childhood NHL consortia, the European Intergroup for Childhood Non-Hodgkin Lymphoma (EICNHL) and the international Berlin-Frankfurt-Münster (i-BFM) Group, as well as selected North American centers, designed a retrospective multinational study on pediatric PCNSL. Here, we present data from the largest series of pediatric patients with PCNSL reported to date.

Methods

We conducted a retrospective survey of children and adolescents up to 19 years of age who were diagnosed with PCNSL between 1991 and 2019. Patient data were retrieved from 21 EICNHL and/or i-BFM Group members or selected North American institutions. The survey included questions about demographics, disease, treatment, outcome, and late effects. PCNSL was defined by intracerebral/intraspinal mass(es) and/or cranial nerve palsies not caused by an extradural tumor and/or tumor cells in the cerebrospinal fluid and/or involvement of the eye(s). Although the 2016 World Health Organization Classification now recognizes diffuse large B-cell lymphoma (DLBCL) of the central nervous system (CNS)

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Table 1. Initial characteristics and therapy of the 75 PCNSL study patients

Variable	Patients
Patients, n	75
Sex	
Female	29 (39)
Male	46 (61)
Age, y	
Median (range)	12.5 (1.25-18.87)
<10	26 (35)
≥10 to <15	31 (41)
≥15	18 (24)
Preexisting disorder	
Yes	14 (19)
No	56 (75)
Unknown	5 (6)
Lansky score	
<40%	14 (19)
≥40-79%	27 (36)
≥80%	19 (25)
Unknown/not applicable	15 (20)
B symptoms	
Yes	9 (12)
No	59 (79)
Unknown	7 (9)
Cranial nerve palsies	
Yes	33 (44)
No	39 (52)
Unknown	3 (4)
Histopathologic subtype	
DLBCL	37 (49)
Burkitt lymphoma	9 (12)
Mature B-NHL NOS	5 (7)
ALCL	17 (23)
Other NHL	7 (9)
BCP LBL, n	4
EN MZL, n	1
PTCL, n	2
Localization	
Intracranial	70 (93)
Intraspinal only	2 (3)
Leptomeningeal only	3 (4)
No. of lesions	
1	36 (48)
≥2	32 (43)
Unknown/not applicable	7 (9)
Initial LDH level, U/L	
<500	53 (71)
≥500	10 (13)
Unknown	12 (16)

Table 1. (continued)

Variable	Patients
Initial therapy	
Chemo only	31 (41)
Chemo and rituximab	15 (20)
Chemo and RT	24 (32)
Chemo and rituximab and RT	2 (3)
Other	3 (4)
Therapy	
Pediatric NHL-type*	57 (76)
Adult NHL-type†	1 (1)
Miscellaneous‡	17 (23)
Chemotherapeutic drugs, n§	
High-dose methotrexate	68
High-dose cytarabine	55
Anthracyclines	59
Alkylating agents	64
Intrathecal therapies	69
RT	
Yes	26 (35)
No	49 (65)
Stem cell therapy in CR1	
Autologous	3 (4)
Allogeneic	1 (1)
Rituximab in mature B-NHL	17 (32)

Unless otherwise noted, data are shown as n (%).
 BCP, B-cell precursor; Chemo, chemotherapy; CR1, first complete remission; EN MZL, extranodal marginal zone lymphoma; LDH, lactate dehydrogenase; NOS, not otherwise specified; RT, radiotherapy.
 *NHL Berlin-Frankfurt-Münster therapy, n = 30; Lymphomes Malins B therapy, n = 14; Inter-B-NHL therapy, n = 5; Associazione Italiana Ematologia Oncologia Pediatrica linfoma non-Hodgkin (AIEOP LNH) therapy, n = 2; EICNHL therapy, n = 2; Sociedade de Hematologia e Oncologia Pediatrica LNH therapy, n = 1; Japanese Pediatric Leukemia/Lymphoma Study Group B-NHL therapy, n = 2; Children's Oncology Group AALL1131 trial, n = 1, including 17 of the 26 (65%) patients with RT.
 †Cancer and Leukemia Group B 50202 trial, n = 1.
 ‡Including 9 of the 26 (35%) patients with RT.
 §High-dose-methotrexate (n = 68): <5 g/m², n = 5; 5 g/m², n = 38; 8 g/m², n = 25. High-dose-cytarabine (n = 55): ≤6 g/m², n = 13; 8 g/m², n = 1; 12 to 12.5 g/m², n = 41. Anthracyclines (n = 59) with doxorubicin in 57 patients and a median cumulative dose of 100 mg/m² (range: 25-240 mg/m²).

as a distinct lymphoma, PCNSL is an accepted clinical term that is independent from the histopathology.^{1,2,5-7} For this reason, we included all histologic subtypes. For patients enrolled in national trials/registries, data collection was approved by the respective ethics committees.⁸⁻²³ In nontrial/nonregistry patients, the retrospective data collection was performed with local Institutional Review Board approval.

Seventy-five patients with PCNSL were identified. Diagnosis of the NHL subtype was based on contemporaneous pathological criteria.^{6,24-27} Event-free survival (EFS) was calculated from date of diagnosis to date of first event. Events considered were relapse, progression, and death. Overall survival (OS) was defined as the time from diagnosis to death from any cause or date of last follow-up. EFS and OS curves were estimated according to the Kaplan-Meier method. The Cox proportional-hazard model was used for multivariate analyses of EFS and OS.

Table 2. Disease outcome and long-term side effects of the 75 PCNSL study patients

Variable	Patients
Treatment failure	
Median (range) time to, y	0.77 (0.27-6.78)
Relapse*	12 (16)
Progression	2 (3)
Localization of failure, n	
CNS only	8
CNS and other	2
Outside CNS only	2
Unknown	1
Multiple sites n.f.s.	1
Death	
	12 (16)
Relapse/progression, n	6
Without any therapy, n	1
Therapy-related toxicity, n	4
Unknown, n	1
Second malignancy, n	0
Continuous CR	58 (75)
Follow-up	
Median (range), y	5.22 (0.19-25.29)
In CR1	53 (91)
In CR2	5 (9)
CR not yet achieved at LFU, n	5
Long-term side effects	
Neurological	22 (29)
Endocrine	3 (4)
Cardiac	0 (0)
Other	7 (9)
Osteonecrosis, n	5
Psychosocial, n	1
GVHD, CMV-associated retinitis, n	1

Unless otherwise noted, data are shown as n (%).
 CMV, cytomegalovirus; CR, complete remission; CR1, first complete remission; CR2, second complete remission; GVHD, graft-versus-host disease; LFU, last follow-up; n.f.s., not further specified.
 *ALCL, n = 6; DLBCL, n = 5; Burkitt lymphoma, n = 1; extranodal marginal zone lymphoma, n = 1; LBL, n = 1.

Results and discussion

The study cohort is shown in Tables 1-3 and supplemental Tables 1-3. The male/female ratio was 1.7:1, and median age was 12.5 years. Three histopathological subgroups were identified: (1) mature B-cell NHL (B-NHL; n = 51; 68%, including DLBCL [n = 37], Burkitt lymphoma [n = 9], and B-NHL, not otherwise specified [n = 5]), (2) anaplastic large cell lymphoma (ALCL; n = 17; 23%), and (3) other NHL (n = 7; 9%, including B-cell precursor lymphoblastic lymphoma [LBL; n = 4], peripheral T-cell lymphoma [PTCL; n = 2], and extranodal marginal zone lymphoma [n = 1]). Fourteen patients (19%) had preexisting disorders, including immunodeficiency (n = 11), hematologic malignancy (n = 2), and factor VII deficiency (n = 1). Epstein-Barr

Table 3. Characteristics of the 14 PCNSL patients with preexisting disorders

Type of preexisting disorder	Patients, n	Histology	Event
Immunodeficiency			
HIV infection	2	Burkitt lymphoma/ DLBCL	No/dead: TRM (sepsis)
St. p. heart transplantation	1	DLBCL	Dead: TRM (graft failure)
St. p. kidney transplantation	1	DLBCL	No
St. p. liver transplantation	1	DLBCL	No
SLE	1	DLBCL	Dead (cause unknown)
CLIPPERS	1	DLBCL	No
Colitis ulcerosa	1	DLBCL	Dead: TRM (details NA)
PID n.f.s.	2	DLBCL/DLBCL	No/dead of disease
CVID	1	ALCL	Dead: TRM (pneumonia)
Other disorders			
B-cell precursor LBL	1	DLBCL	No
JMML	1	DLBCL	Relapse
Factor VIII deficiency	1	ALCL	No

CLIPPERS, chronic lymphocytic inflammation with pontine perivascular enhancement responsive to steroids; CVID, common variable immunodeficiency; JMML, juvenile myelomonocytic leukemia; NA, not available; PID n.f.s., primary immunodeficiency not further specified; SLE, systemic lupus erythematosus; St. p., status post; TRM, treatment-related mortality.

virus (EBV) status was documented in 47 patients, among whom 9 (19%) were EBV+.

Median time to diagnosis was 6.4 weeks, with 31 patients (41%) receiving a diagnosis within 4 weeks of symptom onset. Thirty-three patients (44%) presented with cranial nerve palsies. Seventy patients (93%) had solid intracranial masses, 2 patients (3%) had intraspinal mass(es) only, and 3 patients (4%) had leptomeningeal disease with involvement of the cerebrospinal fluid only. The Lansky score was available in 60 patients: 19 had a score \geq 80%, and 14 had a score $<$ 40%.²⁸

Fifty-eight patients (77%) were treated according to NHL-subtype-directed protocols (pediatric type, n = 57; adult type, n = 1; B-NHL-block therapy: B-NHLs, n = 38, ALCL, n = 14; leukemia-type therapy: LBL, n = 4, PTCL, n = 2), and 17 patients (23%) received miscellaneous therapies.⁸⁻²³ Thirty-one patients (41%) were treated with chemotherapy only, 15 (20%) were treated with chemotherapy + rituximab, 24 (32%) were treated with chemotherapy + radiotherapy, and 2 (3%) were treated with chemotherapy + rituximab + radiotherapy. One patient underwent observation alone after total resection, 1 received no therapy for 3 months for logistic reasons, and 1 patient remained without therapy.

The chemotherapeutic cycles included high-dose methotrexate in 68 patients (91%), high-dose cytarabine in 55 patients (73%), and anthracyclines in 59 patients (79%). Alkylating agents were used in 64 patients (85%), with cyclophosphamide used in 61 patients. Intrathecal therapies were used in 69 patients (92%) including triple therapies in 56 patients. Twenty-six patients underwent irradiation (35%; whole brain, n = 22) with a dose range of 18 to 45 Gy (ALCL, n = 13; DLBCL, n = 9; other NHL, n = 4).

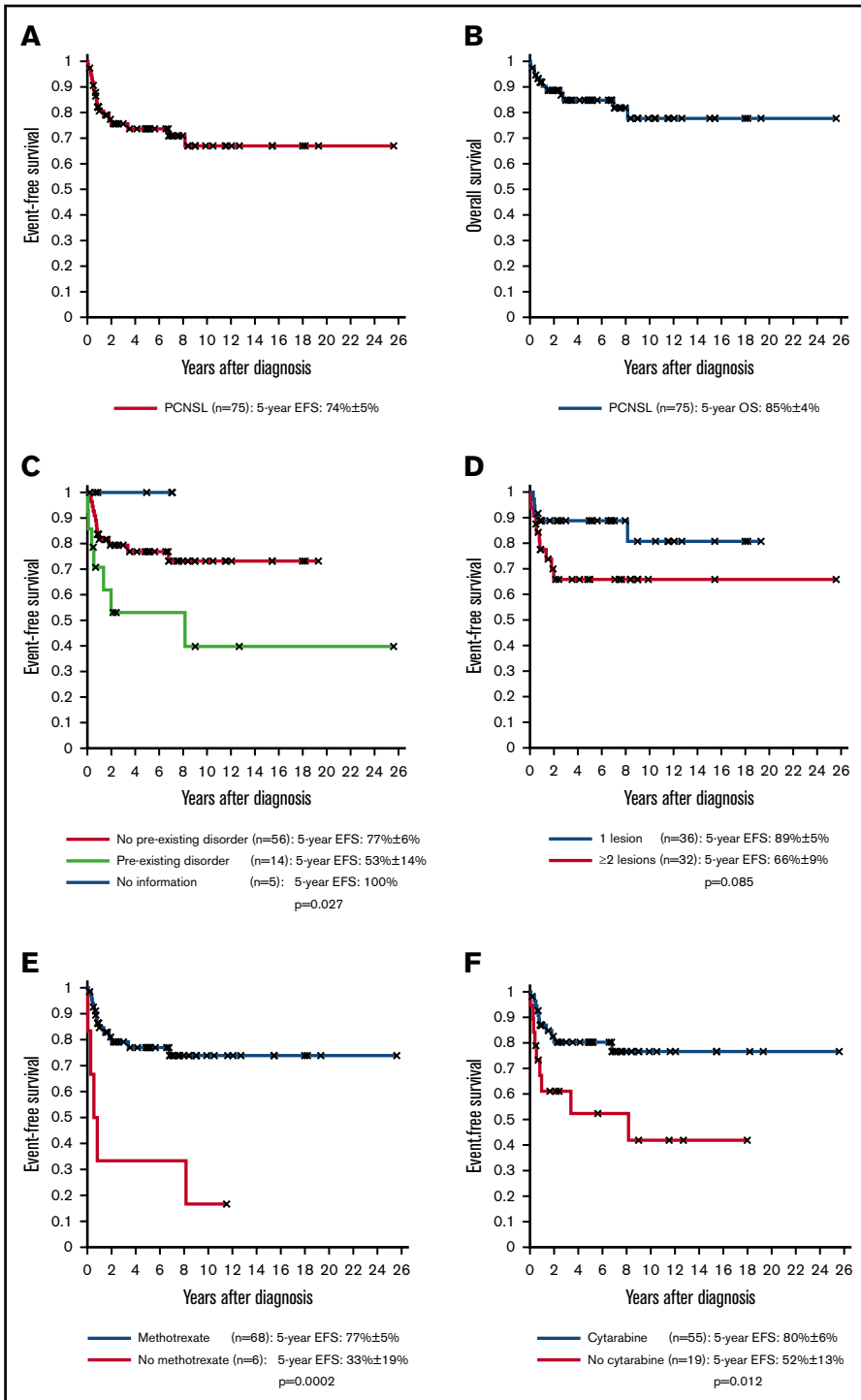


Figure 1. EFS and OS rates of the 75 PCNSL patients according to risk factors. EFS (A) and OS (B) of the 75 PCNSL patients. (C) EFS of the 56 PCNSL patients without a preexisting disorder vs 14 PCNSL patients with a preexisting disorder. (D) EFS of the 36 PCNSL patients with 1 lesion vs 32 PCNSL patients with ≥ 2 lesions. (E) EFS of the 68 PCNSL patients who received therapies that included high-dose methotrexate vs 6 PCNSL patients who did not receive high-dose methotrexate. (F) EFS of the 55 PCNSL patients who received therapies with high-dose cytarabine vs 19 PCNSL patients who did not receive high-dose cytarabine.

Four patients (5%) completed therapy with a stem cell transplantation (autologous, $n = 3$; allogeneic, $n = 1$). After a median time of 5.22 years, 5-year EFS and OS were $74\% \pm 5\%$ and $85\% \pm 4\%$, respectively (Figure 1A-B). Of the 75 patients, 14 (19%) had primary treatment failure (relapse; $n = 12$; progression, $n = 2$) after a median time of 0.77 years, including only 1 patient with a preexisting disorder. Twelve patients (16%) died, including 6 after relapse, 1 from PCNSL without any therapy, 4 from

treatment-related toxicity as a first event, and 1 from unknown causes. Of the latter 6 patients, all had preexisting disorders.

Univariate analysis showed that preexisting disorders (EFS, $77\% \pm 6\%$ vs $53\% \pm 14\%$, $P = .027$; OS, $89\% \pm 5\%$ vs $62\% \pm 14\%$, $P = .001$) and number of lesions (1 vs ≥ 2 lesions: EFS, $89\% \pm 5\%$ vs $66\% \pm 9\%$, $P = .085$; OS, $97\% \pm 3\%$ vs $76\% \pm 8\%$, $P = .043$) were associated with inferior outcome (Figure 1C-D). In contrast,

high-dose methotrexate ($P = .0002$), high-dose cytarabine ($P = .012$), and alkylating agents ($P = .033$) were associated with improved EFS (Figure 1E-F). Other factors, including age, sex, Lansky score, B symptoms, histopathology, type of therapy, rituximab in B-NHLs, or radiotherapy, had no significant impact on EFS or OS (supplemental Table 4). In multivariate analysis, high-dose methotrexate remained significant for EFS, and preexisting disorders remained significant for OS. Analysis of 60 patients without preexisting disorders showed an EFS of $83\% \pm 5\%$ for pediatric NHL therapy vs $57\% \pm 19\%$ for miscellaneous therapy ($P = .044$; supplemental Figure 1), whereas, among 14 patients with a preexisting disorder, 5 of 6 patients with NHL therapy had an event but only 2 of 8 patients with miscellaneous therapy had an event.

Long-term sequelae were primarily neurological ($n = 22$, 29%; including 8/26 with radiotherapy) and rarely affected endocrine organs ($n = 3$, 4%), heart ($n = 0$), or other organs/systems ($n = 7$, 9%).

This report on 75 patients with PCNSL represents the largest series of PCNSL in childhood and adolescence reported to date.^{1-3,29-31} Our results show that PCNSL is associated with male sex, age ≥ 10 years, solid intracerebral disease, and immunodeficiency. Among adults with PCNSL, the majority of cases are DLBCL (90%).⁷ In our cohort, DLBCL was the most common histology, but it was observed in only 49% of cases, suggesting that histopathology of PCNSL in pediatrics might be distinct.

EFS of $74\% \pm 5\%$ for the entire cohort is comparable to that of children with CNS⁺ systemic NHL and similar to the 2-year progression-free survival of 61% that was reported by the International PCNSL Collaborative Group in 29 2- to 21-year-old PCNSL patients.^{11-13,15,16,23,29} Our favorable outcome has been achieved by histological subtype-directed polychemotherapy in most patients, suggesting that pediatric PCNSL can be successfully managed by protocols designed for children with CNS⁺ systemic NHL, usually including high-dose methotrexate and high-dose cytarabine, steroids, and no irradiation (except for ALCL).^{11-13,15,16,23,29} Our evaluation of prognostic factors found preexisting disorders to be associated with inferior EFS and OS. In contrast, usage of high-dose methotrexate, high-dose cytarabine, and alkylators, which are fundamental drugs of all pediatric NHL-subtype protocols/trials, were associated with improved prognosis.^{11-13,15,16,21,23,29}

Our study has limitations, including its retrospective nature, with potential for reporting bias. For subanalyses, patient numbers were small and included different histologic entities. Nevertheless, given the rarity of this entity, these data are likely to be the best available to determine important factors in the treatment and outcome of pediatric PCNSL. This international collaboration allowed us to conclude that treatment recommendations are especially needed for PCNSL patients with immunodeficiencies, who are at risk for treatment-related toxicity/mortality, whereas immunocompetent PCNSL patients receiving pediatric histological subtype-directed polychemotherapy and no irradiation do well.

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Authorship

Contribution: A.A., O.A., and L.G.-R. designed and planned the study and wrote the manuscript; A.A., S. Bansil, S.T., and L.R. pooled data, checked data, and performed statistical analyses; S. Bomken, B.B., F.C., A.K.S.C., H.D., A.F., M.H., J.J., E.K., R.S.K., J. Lazic, J. Loeffen, N.M., T.O., M.P., A.P., C.R., G.T., A.U., J.V.-A., W.W., G.W., A.A., O.A., and L.G.-R. were principal investigators or coinvestigators in their study groups and institutions, coordinated the national trials in their countries, provided study materials, and recruited patients; and all authors read and approved the final version of the manuscript.

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