

Significance of Bcl-2 and Bcl-6 immunostaining in B-Non Hodgkin's lymphoma

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Abstract

The determination of prognosis for B-Non-Hodgkin's lymphoma (NHL) is known to be related to the multiple differences in tumor cell biology. Bcl-2 and Bcl-6 are two markers linked to germinal center B cells. Both markers are thought to have an effect on prognosis of mature B-cell neoplasms. Forty-four patients with chronic B-cell neoplasm were included; Bcl-2 and Bcl-6 expression by immunohistochemistry was examined. Bcl-2 protein was positive in 36.4% (16 of 44) of cases (62.5% of follicular lymphoma, 16.7% of mantle cell lymphoma and 30% of diffuse large B-cell lymphoma); the positive group implying a bad prognostic effect of the marker in NHL. Bcl-6 was positive in 13.6% (6 of 44) of cases (11.1% of mantle cell lymphoma and 40% of diffuse large B-cell lymphoma) and its positivity implies a better disease course. Bcl-2 and Bcl-6 can be used as prognostic marker in NHL.

Introduction

The determination of prognosis for each of the Non-Hodgkin's lymphoma (NHL) variants is known to be related to the multiple differences in tumor cell biology (cytogenetics, immunophenotype, growth fraction, cytokine production) found within each of the specific disease variants.^{1.3}

Bcl-2 and Bcl-6 are two markers linked to germinal center B cells. The Bcl-2 gene, located at chromosome 18q21, encodes for a 25kd protein located mainly in the mitochondrial membrane. This Bcl-2 protein is an anti-apoptosis factor that is important in normal B-cell development and differentiation. Bcl-2 overexpression provides a survival advantage for malignant B cells and is thought to play a critical role in resistance to chemotherapy.⁴

The Bcl-6 gene, located at chromosome 3q27, encodes a 79kd DNA binding protein and transcriptional repressor that appears to be involved in mediating growth suppression.⁵ In lymphoid tissues, the Bcl-6 protein is expressed in germinal center B cells in secondary follicles in both the dark (centroblast-rich)

and light (centrocyte-rich) zones.6

Immunohistochemical studies of the prognostic value of Bcl-6 expression led to discordant results. Some supported its ability to predict for a better overall survival rate, whereas others found no difference in overall survival rates.⁷

Materials and Methods

Subjects

Forty-four newly diagnosed B-NHL adult patients attending the hematology unit of the clinical pathology department, Ain Shams University took part in the study. Patients were recruited to the study on the basis of clinical and laboratory criteria of B-NHL. All patients were subjected to clinical sheet details, complete blood count with examination of PB smears stained with Leishman stain, bone marrow aspiration, bone marrow trephine biopsy, immunophenotyping using lymphoproliferative disorder panel (CD19, CD5, CD10, CD11c, CD20, CD22, CD23, CD38, CD79b, FMC7, CD103, CD25, kappa and lambda light chains). Bcl-2 and Bcl-6 immunostaining was performed on fixed bone marrow trephine biopsy specimen.

The diagnosis of NHL was confirmed by tissue biopsy in 40 patients (lymph node biopsy in 38 patients and liver biopsy in 2 patients). In the remaining 4 patients, bone marrow was the first diagnostic material. The 44 NHL cases comprised 18 mantle cell lymphomas (MCL), 16 follicular lymphomas (FL) and 10 diffuse large B-cell lymphomas (DLBCL).

Methods

Two milliliter peripheral blood samples were collected in a sterile EDTA containing vaccutainers for CBC and immunophenotyping. BM aspiration was withdrawn, a few drops were put on glass slides for morphological examination and 1 mL into sterile EDTA containing vacutainers for immunophenotyping. BM trephine core biopsy was obtained and transferred immediately in a sterile plastic cup containing aldehyde solution fixative.

Immunohistochemistry

Fixation was performed for 24 h; decalcification of the core was performed using disodium ethylene-diamine-tetra acetic acid for 48 h. This was followed by passing the core in serial concentrations of ethyl alcohol (50%, 70%, 85%, 90%, and 100%) ending with xyelene then wax followed by paraffin embedding.

Serial $3-\mu m$ sections were cut from the paraffin block, mounted on positively charged

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slides and dried overnight in a 60° C oven. Sections were then deparaffinized in xylene for 24 h and hydrated in a descending grades of alcohol; 100%, 90%, 85% and 70%. Antigen unmasking was performed by heat induced an epitope retrieval method by placing the slides in a plastic Coplin jar filled with citric acid buffer so that the solution covers the slides, then placing the jar in a microwave at 800Watt for 20 min (divided into 4 cycles 5 min each). The Coplin jar was then removed from the oven and allowed to cool for 15 min.

Slides were placed in a humidified chamber and rinsed three times in phosphate buffer saline (PBS). Endogenous peroxidase activity was blocked by incubation of the tissue section with 3% hydrogen peroxide in water for 30 min. After washing, the tissue sections were then incubated with the primary monoclonal antibody, ready to use Bcl-2 and 1:10 dilution in Bcl-6. Two hundred μ l of the monoclonals were added for 1 h at room temperature with Bcl-2 and overnight at 4°C with Bcl-6.

The Streptavidin biotin method was used as a detection kit (LSAB2, Dako, Denmark). Tissue sections were incubated with biotinylated secondary antibody for 40 min, then with Streptavidin conjugated enzyme for 30 min during which the 3,3' diaminobenzidine (DAB) substrate chromogen was freshly prepared then added onto the tissue section for 10 min in the dark.

The sections were counterstained in Meyer hematotoxylin, dehydrated, then put in xyelene and cover-slipped by DPX mount media and examined under a light microscope.

The positivity of the immunostaining was detected by percentage of positive cells and intensity of staining. For Bcl-2 the positivity



cut off was considered when the reactivity of lymphoma cells with antibodies was more than $20\%^8$ and was more than 10% for Bcl-6.⁹

The pattern of staining was also considered. For Bcl-2, membranous and cytoplasmic patterns were considered positive. For Bcl-6, positivity was detected if the pattern of staining was unclear.

Treatment regimen

Patients with NHL were treated with chemotherapy protocols with or without regional radiation therapy. The used therapeutic protocol was CHOP regimen; cyclophosphamide 650 mg/m² day, vincristine 1.4 mg/m² up to 2 mg/m² and doxorubicin 40 mg/m² were given in day one whereas prednisone in a dose of 40 mg/m² was administrated from day 1 till day 5. Cycles were repeated at intervals of 3-4 weeks according to the tumor volume and performance status of patients. Rituximab was added to CHOP (R-CHOP) for B-cell lymphomas with aggressive histology.^{10,11}

Follow up

The follow-up strategy used to assess remission included history, physical examination, serum LDH and CBC after 2-4 cycles according to the International Workshop Group (IWG) criteria.^{12,13} Follow-up ranged from 12 to 30 months with a mean of 19.5 ± 0.7 months. Response to therapy was assessed at the end of induction and complete remission (CR) was defined as the disappearance of all physical and radiographic evidence of lymphoma for at least four weeks after systemic chemotherapy and/ or radiation.¹⁴ In addition, patients with initial BM involvement were required to have clearing (as determined by repeated aspiration and biopsy) to be categorized as complete responders. Nonresponders included patients in partial remission where a reduction in the size of measurable lesions by at least 50% occurred with no new lesions appearing and resistant cases who had no response to the used chemotherapeutic regimen. Relapse was determined as the recurrence of lymphoma in patients who had been in CR for at least four weeks.¹²

Statistical analysis

Data were analyzed using a social science statistical package (SPSS version 15.0.1). The following tests were performed: descriptive statistics including quantitative data (mean \pm standard deviation (SD) for parametric, and in the form of median and range in non-parametric results), qualitative data (number and percentage), analytical statistics including χ^2 , Fisher's exact test, Wilcoxon Rank-sum test (Z-value) and quantitative parametric data was analyzed using Student's t-test (t-value). The overall survival (OS) and disease-free survival

(DFS) rates along with standard error (SE) were estimated using the Kaplan-Meier method and comparison between groups was made using a log rank test. P values ≤ 0.05 and ≤ 0.01 were considered significant and highly significant, respectively, in all analyses.

Results

There were 26 males and 18 females with a male to female ratio 1.4:1.0. Ages ranged from 20 to 70 years old with a mean of 45 ± 15.2 years. The results of this study are summarized in Tables 1-3 and Figures 1-5.

Bcl-2 protein was positive in 36.4% (16 of 44) of cases with the most frequent expression in the FL group (P=0.019); 62.5% of FL, 16.7% of MCL and 30% of DLBCL. Bcl-6 was positive in 13.6% (6 of 44) of cases with the most fre-

quent expression in the DLBCL group (P=0.014) (11.1% of MCL and 40% of DLBCL) (Table 1).

Relation of Bcl-2 protein expression to clinical and laboratory prognostic factors

On comparing Bcl-2 expression with various prognostic parameters, a statistically significant difference was obtained in platelets count in which 75% of Bcl-2 positive cases were thrombocytopenic (P=0.001). A higher serum LDH serum level was detected in Bcl-2 positive cases compared to Bcl-2 negative group and the difference was statistically significant (P=0.004). On the other hand, no statistical significance was found as regards age, staging, total leukocytic count, hemoglobin level and pattern of BM infiltration (Table 2 and Figure1).

Table 1. Details of Bcl-2 and Bcl-6 expression in different groups of chronic B cell neoplasms.

	N. of cases	Bcl-2 positive n (%)	Bcl-6 positive n (%)		
FL	16	10 (62.5)	0 (0)		
MCL	18	3 (16.7)	2 (11.1)		
DLBCL	10	3 (30)	4 (40)		
Total	44	16 (36.4)	6 (13.6)		

Table 2. Comparing Bcl-2 expression and prognostic parameters in NHL group

Parameter		Bcl-2 positive		Bcl-2 negative		Р	Significance
	1	N. cases	% cases	N. cases	% cases		
Age (mean)	≥45 yrs	12	75	16	57.1	0.236	NS
	<45 yrs	4	25	12	42.9		
Sex (mean)	М	10	62.5	16	57.1	0.728	NS
	r	0	31.5	12	42.9	-	
Hb g/dL (mean	l) ≥l0	8	50	14	50	1	NS
D	<10	0	50	14	20		
Plt ×10%L (mea	$an) \geq 150$	4 19	25 75	22	78.6	0.001	HS
TTLO 109/L (<130	14	10	0	01.4	0.050	NO
ILC ×10%L (m	ean) ≥8	8	50 50	6 99	21.4 78.6	0.052	INS
IDIIII/I (maa	<0 m) . E00	14	07 E	10	10.0	0.004	110
LDH IU/L (IIIea	$(11) \geq 500$	14	07.0 12.5	12	42.9 57 1	0.004	пэ
Stago	I	0	0	10 C	91.4	0.920	NC
Stage	I	0	12.5	0 4	21.4 14 3	0.239	IND
	III	6	37.5	8	28.6		
	IV	8	50	10	35.7		
Pattern	Diffuse	4	25	2	7.2	0.36	NS
	Patchy	4	25	10	35.7		
	Interstitial	2	12.5	6	21.4		
	Mixed	6	37.5	10	35.7		
Therapeutic	Responders	2	12.5	18	64.3	0.001	HS
response	Non-responders	14	87.5	10	35.7		
Clinical C	Complete remissio	on 2	12.5	22	78.6	0.001	HS
outcome	Relapse	14	87.5	6	21.4		
Fate	Alive	7	43.8	24	85.7	0.003	HS
	Death	9	56.2	4	14.3		

S, significant; NS, non-significant; TLC, total leukocytic count; Hb, hemoglobin; Plts, platelets; BML, bone marrow lymphocytes; LDH, lactate dehydrogenase.



This study showed that 87.5% of patients with positive Bcl-2 protein expression significantly developed resistance to the therapeutic regimens adopted at the end of induction (P=0.001). Interestingly, these patients also displayed a poor clinical outcome experiencing higher relapse and mortality rates than their negative counterparts (P=0.001 and P=0.003, respectively).

Bcl-2 expression in relation to patients' survival

Kaplan-Meier survival analysis revealed significantly shorter OS and DFS times in B-NHL patients with Bcl-2 protein expression (P<0.05). The estimated 30-month OS survival



Figure 1. Bcl-2 positivity in a case of follicular lymphoma (A): x40.



	< 10 J10	0	100	10	10.0		
Sex (mean)	M F	2 4	33.5	24 66.5	63.2 14	0.167	NS 36.8
Hb g/dL (mean)	≥10 <10	2 4	33.5 66.5	20 18	52.6 47.4	0.38	NS
Plt ×10 ⁹ /L (mean)	≥150 <150	6 0	100 0	20 18	52.6 47.4	0.028	S
TLC ×10 ⁹ /L (mean	ı) ≥8 <8	2 4	33.5 66.5	12 26	31.6 68.4	0.932	NS
LDH IU/L (mean)	≥500 <500	4 2	66.5 33.5	22 16	57.9 42.1	0.685	NS
Stage	I II III IV	2 0 2 2	33.5 33.5 33.5	4 0 12 16	10.5 6 31.6 42.1	0.388	NS 15.8
Pattern Ir	Diffuse Patchy iterstitial Mixed	0 2 0 4	0 33.5 0 66.5	6 12 8 12	15.8 31.6 21.0 31.6	0.26	NS
Therapeutic Re response Non-	esponders -responders	4 2	66.5 33.5	16 22	42.1 57.9	0.26	NS
Clinical Compl outcome	ete remission Relapse	6 0	100	18 20	47.4 52.6	0.016	S
Fate	Alive Death	6 0	100	25 13	78.9 21.1	0.088	NS

S, significant, NS, non-significant. TLC, total leukocytic count; Hb, hemoglobin; Plts, platelets; BML, bone marrow lymphocytes; LDH, lactate dehydrogenase; Unfavorable = survival > 12 months.



Figure 2. Bcl-2 positivity in a case of follicular lymphoma (B): x10



Figure 3. Bcl-6 positivity in a case of DLBCL (x40).



Figure 4. Kaplan-Meier overall survival and disease-free survival times in relation to Bcl-2 protein expression.



S





rate was 44% in Bcl-2 positive patients compared to 85% in negative cases (log rank 8.7; P=0.003) and DFS rate was 12% versus 64% (log rank, 19.13; P=0.001) (Figure 3).

Relation of Bcl-6 protein expression to clinical and laboratory prognostic factors

A statistically significant difference (P=0.001) was detected among the studied patients as regards age in which all the Bcl-6 positive cases were below the age of 45 years. In addition, a statistically significant difference was also obtained (P=0.028) in platelet count in which all Bcl-6 positive cases had a normal platelet count. No statistically significant difference was seen as regards other prognostic factors (P>0.05) (Table 3 and Figure 2).

Impact of BcI-6 protein expression on response to therapy and clinical outcome

No statistically significant difference (P>0.05) was detected among patients with positive Bcl-6 protein expression as regards response to the therapeutic regimens adopted and mortality rates. However, a significantly lower relapse rate was detected in patients with positive Bcl-6 protein than in patients with negative expression (P=0.016).

Bcl-6 expression in relation to patients' survival

As regards Bcl-6 expression, results of survival analysis were consistent among Bcl-6 positive and negative patients (P>0.05). Nevertheless, OS and DFS were higher in patients exhibiting this marker (OS 100% versus 79% and DFS 66% versus 45%) (Figure 4).

Discussion

Several prognostic parameters were used in B-NHL to detect patients' outcome. To detect the value of Bcl-2 and Bcl-6 in diagnosis and prognosis of such diseases, immunostaining of Bcl-2 and Bcl-6 was performed for 44 patients with B-NHL. In this study, using a cut off 20% or more, Bcl-2 protein was positive in 36.4% (16 of 44) of NHL cases (62.5% of FL, 16.7% of MCL and 30% of DLBCL cases).

Studying Bcl-2 in lymphoma cases, Papakonstantinous *et al.*¹⁵ proved that the expression of Bcl-2 protein is not restricted to B-cell lymphomas bearing the t(14; 18)translocation and they showed the complete absence of any correlation between BCL-2 gene rearrangements and Bcl-2 protein expression in NHL. Ben-Ezra and King¹⁶ have suggested that Bcl-2 alone is useful in discriminating FL and benign lymphoid aggregates. In



Figure 5. Kaplan-Meier overall survival and disease-free survival times in relation to Bcl-6 protein expression.

spite of the fact that the absence of Bcl-2 was highly specific for benign lymphoid aggregates, the expression of Bcl-2 in some benign and atypical lymphoid aggregates in the study by West *et al.*¹⁷ did not make Bcl-2 a specific marker for the detection of FL in bone marrow biopsy specimens. In addition, Llanos *et al.*¹⁸ stated that Bcl-2 expression is related to the grade of FL being highest in grade I and lowest in grade III, which matches with our negative cases where 5 of 6 of our negative Bcl-2 FL cases were of grade III.

The percentage of positive Bcl-2 in MCL cases was surprisingly low (16.7%). In fact, our negative MCL cases gave minimal positivity ranging from 5 to 20%; yet we considered them negative results according to the previously agreed cut off. Navratile *et al.*¹⁹ also found that Bcl-2 is expressed in low grade but not high grade lymphoma, and although MCL looks like a slow growing, low-grade tumor under the microscope, it grows fast and behaves like a high-grade lymphoma.

Bcl-2 protein expression was studied in relation to treatment response and clinical outcome, revealing a statistical significance, where a high tendency for developing resistance and a poor clinical outcome with significantly shorter OS and DFS were shown in Bcl-2 positive compared to negative patients. Comparing Bcl-2 to prognostic parameters in the NHL group in the current study, there was a significantly higher LDH level and a significantly lower platelet count in Bcl-2 positive NHL cases.

Hadzi-Pecova and his co-workers²⁰ confirmed that the expression of Bcl-2 protein is significantly present in advanced compared with early stages of B-cell lymphoma. Similarly, high Bcl-2 expression by Hermine *et al.*²¹ was more frequently associated with stages III and IV. However, according to Hadzi-Pecova *et al.*,²⁰ Bcl-2 expression did not show any significant influence either on total survival or on diseasefree survival.

Although some reports failed to show any impact of Bcl-2 protein expression on survival of DLBCL cases, ^{22,23} subsequent studies have reported that Bcl-2-positive cases have a worse disease-free survival.²⁴ Recently, Hallack Neto *et al.*²⁵ found a prognostic significance for Bcl-2 protein regarding overall survival.

Llanos *et al.*¹⁸ found no significant differences regarding the prognostic effect of Bcl-2 expression between FL patient groups. On the contrary, they observed a trend toward poorer survival among DLBCL patients with high Bcl-2 expression. Moreover, Iqbal *et al.*⁴ specified that Bcl-2 expression is associated with poor survival in the activated B cell-like subgroup of



All Bcl-6 positive cases were below the age of 45 years and had a normal platelet count. The current study found no relation between therapeutic response, OS and DFS on the one hand, and Bcl-6 expression on the other hand. However, a significantly lower relapse rate was found in patients with positive Bcl-6 protein.

Some authors suggested expression of Bcl-6 to be a prognostic marker in DLBCL²⁶⁻²⁸ but others have contradicted this.^{7,29,30} These differences may partly be explained by the differences in the cut off value (10-30%) for a positive staining in these studies. Also, differences in staining techniques may be of importance, for instance, in one study using the EnVision method and a low cut off of 10%, as much as 97% of cases were found positive for Bcl-6.⁹

Expression of Bcl-6 by Holler et al.8 had identified prognostically relevant favorable subgroups. Also Lossos et al.27 observed a strong predictive value for survival. In a study by Colomo et al.,30 Bcl-6 was expressed in 91% of DLBCL cases (23% Bcl-6 alone and 49% with other markers). The group of patients with restricted expression of Bcl-6 presented with earlier stage, low-risk IPI and normal LDH values. However, although this group of patients had the highest overall survival rate, the difference did not reach statistical significance. The reasons for these discordant results in different studies are not clear but they may be related in part to the heterogeneity of the selected patients. The aberrant expression of Bcl-6 in MCL found in the current study was also experienced by some authors. Only one of 20 MCL cases studied by Chuang et al.31 expressed Bcl-6. Similarly, Gualco et al. 32 found Bcl-6 expression in 12% of the MCL cases.

In conclusion, Bcl-2 and Bcl-6 are found frequently in B-NHL and both proteins should be considered as reliable prognostic markers. Bcl-2 positive patients were associated with poor prognosis as well as short OS and DFS times whereas Bcl-6 positive patients were associated with a favorable prognosis. This indicated their potential as promising targets for therapeutic intervention. However, prospective studies with more patients in each group together with a longer follow up are recommended.

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