

Original Article

Immunophenotyping and gene rearrangement analysis in lymphoid/lymphoproliferative disorders of the lungs*

Imunofenotipagem e rearranjo gênico em doenças pulmonares linfocíticas e linfoproliferativas

Camila Cristina Ishikawa¹, Alexandre Muxfeldt Ab'Saber², Edwin Roger Parra², Chin Jia Lin², Carmen Silvia Valente Barbas³, Vera Luiza Capelozzi⁴

Abstract

Objective: To determine the usefulness, in routine practice, of using polymerase chain reaction to analyze B and T lymphocyte clonality in pulmonary tissue as a tool for the diagnosis of pulmonary lymphoproliferative disorders. **Methods:** Immunohistochemistry and molecular gene rearrangement analysis were performed in order to assess 8 cases of lymphoid interstitial pneumonia (LIP) and 7 cases of pulmonary lymphoproliferative disorders. **Results:** All 8 cases of LIP presented moderate to strong immunostaining for CD3, compared with only 2 cases of lymphoma and 1 case of pseudolymphoma ($p = 0.02$). Gene rearrangement was detected in 4 of the 8 cases, which changed the diagnosis from LIP to lymphoma, showing the importance of gene rearrangement detection in cases of LIP. In this situation, gene rearrangement using the VH/JH and V γ 11/J γ 12 primer pairs was detected in 3 cases and 1 case, respectively, and no gene abnormalities were found using the D β 1/J β 2 and V γ 101/J γ 12 primer pairs in any of the cases. A significant positive association was found between the intensity of CD20 and CD68 expression and gene rearrangement using the VH/JH primer pair. Prior to the gene rearrangement, 4 patients with LIP died quickly, whereas only one patient with LIP died after the gene rearrangement. **Conclusions:** Detection of monoclonal B and T cells by immunophenotyping and polymerase chain reaction had an impact on the diagnosis of pulmonary lymphomas in patients previously diagnosed with LIP. Therefore, immunophenotyping and polymerase chain reaction should be used as 'gold standard' techniques in routine practice.

Keywords: Lymphoma; Lung diseases, interstitial; Molecular biology; Polymerase chain reaction.

Resumo

Objetivo: Determinar a utilidade, na prática rotineira, da análise da clonalidade dos linfócitos T e B nos tecidos pulmonares por reação em cadeia da polimerase no diagnóstico das doenças linfoproliferativas pulmonares. **Métodos:** Avaliaram-se, mediante análise imunohistoquímica e rearranjo molecular dos genes, 8 casos de pneumonia intersticial linfocítica (PIL) e 7 casos de doenças linfoproliferativas pulmonares. **Resultados:** Todos os 8 casos de PIL expressaram imunocoloração moderada a forte para CD3, em contraste com apenas 2 casos de linfoma e 1 caso de pseudolinfoma. Rearranjo gênico foi detectado em 4 de 8 casos de PIL, o que mudou o diagnóstico de PIL para linfoma, indicando, assim, a importância da detecção de rearranjo gênico em casos de PIL. Nesta situação, rearranjo gênico usando-se os pares de primers VH/JH e V γ 11/J γ 12 foi detectado em 3 e 1 casos de PIL, respectivamente, e não foram detectadas anormalidades gênicas usando-se as pares D β 1/J β 2 e V γ 101/J γ 12. Uma associação positiva foi detectada entre a intensidade de imunoexpressão CD20 e CD68 e rearranjo gênico usando-se o par de primers VH/JH. Antes do rearranjo gênico, 4 pacientes com PIL morreram rapidamente, enquanto que, após o rearranjo gênico, apenas 1 paciente com PIL morreu. **Conclusões:** A detecção de células B e T monoclonais por imunofenotipagem e reação em cadeia da polimerase mostrou impacto no diagnóstico de linfomas pulmonares em pacientes previamente diagnosticados com PIL. Portanto, imunofenotipagem e reação em cadeia da polimerase devem ser incluídas como métodos de 'padrão ouro' na rotina diagnóstica.

Descritores: Linfoma; Doenças pulmonares intersticiais; Biologia molecular; Reação em cadeia da polimerase.

*Study carried out at the Department of Pathology and in the Division of Pulmonology. Instituto do Coração – InCor, Heart Institute – Faculdade de Medicina da Universidade de São Paulo – FMUSP, University of São Paulo School of Medicine – São Paulo, Brazil.

1. Physician. Faculdade de Medicina da Universidade de São Paulo – FMUSP, University of São Paulo School of Medicine – São Paulo, Brazil.

2. Professor. Faculdade de Medicina da Universidade de São Paulo – FMUSP, University of São Paulo School of Medicine – São Paulo, Brazil.

3. PhD. Faculdade de Medicina da Universidade de São Paulo – FMUSP, University of São Paulo School of Medicine – São Paulo, Brazil.

4. Associate Professor in the Department of Pathology. Faculdade de Medicina da Universidade de São Paulo – FMUSP, University of São Paulo School of Medicine – São Paulo, Brazil.

Correspondence to: Vera Luiza Capelozzi or Edwin Roger Parra. Departamento de Patologia, Faculdade de Medicina da Universidade de São Paulo, Av. Dr. Arnaldo, 455, CEP 01246-903, São Paulo, SP, Brasil.

Tel 55 11 3066-7427. Fax 55 11 3064-2744. E-mail: vcapelozzi@lim05.fm.usp.br / erparra20003@yahoo.com.br

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Introduction

Lymphoid interstitial pneumonia (LIP) is a clinicopathologic term that relates histologically to a dense interstitial infiltrate, composed primarily of T cells, plasma cells, and histiocytes, germinal centers often being identified.⁽¹⁻³⁾ This disease is included in the spectrum of pulmonary lymphoid proliferations, ranging from follicular bronchitis/bronchiolitis and pulmonary lymphoid hyperplasia (proliferations largely limited to the airways) to low-grade malignant lymphoma.^(4,5) It appears that LIP sometimes evolves to lymphoma. Although the frequency of such evolution is probably low, it is difficult to assess, since low-grade lymphomas can mimic LIP.^(3,4)

The major differential diagnosis of LIP is low-grade lymphoma,⁽³⁾ which is typically a well-differentiated B-cell tumor that appears to arise from mucosa-associated lymphoid tissue and is the most common histological subtype of pulmonary non-Hodgkin's lymphoma (NHL).^(6,7) Drawing the distinction between LIP and low-grade lymphoma can be difficult in routine hematoxylin and eosin (H&E)-stained sections and can require immunohistochemical analyses including CD3, CD20, CD4, CD45RO, CD8, CD15, CD30, and CD68 antibodies,^(8,9) as well as molecular gene rearrangement studies using techniques such as polymerase chain reaction (PCR).^(10,11)

The development of molecular biology techniques, such as Southern blotting⁽¹²⁾ and PCR,⁽¹³⁻¹⁵⁾ has provided the ability to detect monoclonal populations of B and T lymphocytes through the detection of rearrangements of the genes that encode B-cell immunoglobulin and T-cell receptor (TCR) proteins. This detection has been applied to support the diagnosis of nodal lymphomas,^(16,17) as well as extranodal lymphomas such as gastric⁽¹⁸⁾ and pulmonary⁽¹⁹⁻²³⁾ lymphomas.

The aim of this study was to determine the feasibility of detecting monoclonal populations of B and T lymphocytes in routine practice. We conducted a retrospective study to assess immunoglobulin heavy chain (IgH) and TCR gene rearrangements using PCR in the diagnosis of LIP. In order to validate the use of this procedure, we also assessed IgH and TCR gene rearrangements in control cases of pulmonary lymphoproliferative disorders.

Methods

This study was approved by the Ethics Committee of the University of São Paulo.

Paraffin blocks with sufficient amounts of tissue were collected from our archives in order to cut serial sections for histochemical preparations. All biopsies had been obtained during the routine clinical care of these patients. The study population consisted of 9 cases (6 females and 3 males; median age of 41 years). Experienced pulmonary pathologists evaluated the samples and determined that 8 of the cases presented a histological pattern consistent with LIP, and that the remaining case presented a histological pattern consistent with pseudolymphoma (Table 1). The diagnosis was based on clinical and histological findings in biopsies (7 surgical lung biopsies and 2 transbronchial biopsies) obtained between 1982 and 2002 and unanimously reclassified according to the American Thoracic Society/European Respiratory Society International Multidisciplinary Consensus Classification of Idiopathic Interstitial Pneumonias.⁽²⁴⁾ Cases of LIP secondary to Sjögren's syndrome or acquired immunodeficiency syndrome were excluded from the study.

As a control, 6 cases of pulmonary lymphoma were included (median age, 43 years): 1 case of low-grade B-cell NHL, 1 case of low-grade B-cell lymphoma with plasmacytic differentiation, 2 cases of high-grade T-cell NHL, 1 case of lymphomatoid granulomatosis, and 1 case of Hodgkin's disease. Tissue specimens had been obtained from autopsy (1 case) or open lung biopsy (5 cases).

Formalin-fixed, paraffin-embedded sections of 3 μ m in thickness were deparaffinized with xylene and rehydrated using a graded alcohol series. Endogenous peroxidase was blocked with seven 5-min washes in 3% hydrogen peroxide. Better amplification was achieved in high temperature masking solution (10 mM citrate buffer, pH 6.0). The sections were washed in Tris-buffered saline and incubated with 2% blocking serum for 20 min. They were subsequently incubated for 12-18 h at 4 °C with primary antibodies at the appropriate dilutions and then incubated with secondary antibodies (LSAB kit peroxidase; Dako, Carpinteria, CA, USA) for 45 min at 37 °C. The peroxidase reaction was developed using 3,3'-diaminobenzidine tetrahydrochloride.⁽²⁰⁾ The sections were counterstained with H&E.

Table 1 – Clinical data of the patients.

Case	Age	Gender	Biopsy	Histology	Survival (months)	Status
1	57	F	TBB	LIP	25	1
2	2	M	OLB	LIP	2	0
3	40	F	OLB	PSLY	17	1
4	17	F	OLB	LIP	65	1
5	52	F	TBB	LIP	2	0
6	20	M	OLB	LYG	20	0
7	38	F	OLB	LGBCL	46	0
8	28	M	OLB	LIP	30	0
9	73	F	OLB	LIP	48	1
10	31	M	OLB	LIP	69	1
11	71	F	OLB	LIP	24	1
12	24	F	Mediast/ OLB	Hodgkin's disease	29	1
13	71	F	OLB	NHL (B-cell low-grade)	17	1
14	81	F	Lymph node	NHL (T-cell high-grade)	NE	0
15	67	M	Lymph node	NHL (T-cell high-grade)	NE	0

LIP: lymphoid interstitial pneumonia; LGBCL: low-grade B-cell lymphoma with plasmacytic differentiation; LYG: lymphomatoid granulomatosis; NHL: non-Hodgkin's lymphoma; NE = not evaluated; OLB: open lung biopsy; PSLY: pseudolymphoma; Status: 0 = dead, 1 = alive; and TBB: transbronchial biopsy.

Any brown cytoplasmic staining of cells characterized positive expression for the following markers: CD3 (dilution 1:200; Dako USA, Santa Barbara, CA, USA); CD20 (dilution 1:800; Dako USA); CD15 (dilution 1:100; Dako USA); CD30 (dilution 1:50; Dako Denmark, Glostrup, Denmark); CD45 (dilution 1:25; Novocastra, Newcastle Upon Tyne, UK); CD68 (dilution 1:100; Dako USA). We also graded the degree of tumor staining for CD3, CD20, CD15, CD30, CD45, and CD68. First, at low magnification, we selected the region of greatest expression. Subsequently, at a magnification of $\times 400$, the degree of tumor staining was graded according to the sum of the intensity of staining and the proportion of cells staining, yielding a potential value of 0 to 4.

DNA was extracted from a 25- μ m section of the paraffin block that best represented each case (previously selected from H&E stained slides). Disposable microtome blades were used, and instruments were cleaned with xylene after each tissue section to avoid cross-contamination.

Paraffin sections were not deparaffinized. DNA was extracted by proteolysis (40 μ L of 10 mg/mL of proteinase K), in the presence of 800 μ L of extraction buffer consisting of 50 mM KCl; 10 mM Tris-HCl (pH 8.3); 2.5 mM MgCl₂; 0.1 mg/mL gelatin; 0.45% Nonidet P40; 0.45% Tween 20. Overnight incubation at 57 °C was followed by 10 min of boiling to deactivate the proteinase K.^(25,26)

A universal VH primer and a consensus JH primer were used to detect complete IgH rearrangements. The analyses of the T-cell receptor (TCR) β - and TCR γ -chain gene rearrangement were performed using two different primer combinations: D β 1/J β 2 and V γ 11/V γ 101/J γ 12. These primers were obtained from Integrated DNA Technologies (Coralville, IA, USA).

Primer combinations were as follows: VH (5'-CTGTCGACACGGCCGTGTATTACTG-3'); JH (5'-AACTGCAGAGGAGACGGTGACC-3'); D β 15'-(CAAAGCTGTAACATTG TGGGGAC-3'); J β 2 (5'-AGCAC(T/G/C)GTGAGCC(T/G)GGTGCC-3'); V γ 115'-(TCT GG(A/G)GTCTATTACTGTGC-3'); V γ 101 (5'-CTCACACTC(C/T)CACTTC-3'); and J γ 12 (5'-CAAGTGTGTGTTCCACTGCC-3').

The reaction mixture consisted of 30 pmol of each oligonucleotide primer; 3 μ L of tissue extract; 200 μ M of each deoxynucleotide triphosphate (dATP, dCTP, dGTP, dTTP); 50 mM KCl; 10 mM Tris-HCl (pH 9.0); 3 mM MgCl₂; 100 μ g/mL bovine serum albumin; and 0.25 μ L of Taq polymerase, in a total volume of 50 μ L with sterile distilled water.

Forty PCR cycles were performed, each cycle consisting of a denaturing step at 93 °C for 1 min, an annealing step for 1 min, and an elongation step at 73 °C for 1 min. In the annealing step, different temperatures were used for each primer pair: 52 °C (for VH/JH, V γ 11/J γ 12, and V γ 101/J γ 12) and 55 °C (for D β 1/J β 2). Following the 40 cycles, there was a

5-min period at 73 °C to ensure complete extension and annealing of the PCR products.

Aliquots of 30 µL were then analyzed by electrophoresis on a 10% polyacrylamide gel stained with ethidium bromide and photographed under ultraviolet light. Bands of relevant size were identified by comparison with a molecular weight marker (1 kb DNA ladder; Invitrogen Carlsbad, CA, USA).⁽¹⁰⁾

All necessary precautions were taken in order to prevent contamination.

In order to identify any correlations among immunohistochemical findings, PCR results, and clinical data, the chi-square test, Fisher's exact test, and the Student's t-test were used. Survival curves were created using the Kaplan-Meier method, and the statistical significance of differences was calculated using the log-rank test, with $p < 0.05$ indicating a significant difference, and the Statistical Package for Social Science (version 10.0 for Windows; SPSS Inc., Chicago, IL, USA).

Results

Figure 1 shows the lymphocytic infiltrates in different histological patterns of lymphoproliferative pulmonary disorders seen after H&E and immunohistochemical staining. In the cases of LIP (Figures 1a to 1d), there was marked lymphocytic infiltration with extensive involvement of the alveolar septa (Figures 1a and 1b). The lymphocytic infiltrate consisted mostly of lymphocytes (B and T cells) with varying numbers of plasma cells. The T lymphocytes (CD3 cells) were seen predominantly in the alveolar septal interstitium (Figure 1c), whereas B lymphocytes (CD20 cells) were primarily found in aggregates within lymphoid follicles and focally in thickening alveolar walls (Figure 1d). In contrast, the cases of low-grade lymphoma (Figures 1e to 1h) presented diffuse, dense, and monomorphous infiltration of small lymphoid cells with irregular nuclear contours (Figures 1e and 1f) remodeling the lung architecture. The neoplastic cells were CD20-positive monoclonal B cells (Figure 1g), with a background population of variable reactive CD3-positive T-cells (Figure 1h).

The molecular biology results are listed in Table 2, which includes the total number of cases and controls, as well as the presence of monoclonality for the genes studied.

The PCR analysis of the IgH gene rearrangement using the VH/JH primer pair revealed a band

of 130 bp (predicted size range, 100-150 bp) indicating the presence of a predominant clonal lymphocyte population in high-grade T lymphoma (Case 13); LIP (Cases 2, 8, and 9); pseudolymphoma (Case 3); lymphomatoid granulomatosis (Case 6); and Hodgkin's disease (Case 12). The cases exhibiting more than one evident band were considered polyclonal, whereas bands lower than 75 bp were assumed to be nonspecific (Figure 2a).

TCR β -chain gene rearrangement analysis using the DB1/J β 2 primer pair showed a band of 60 bp (predicted size range, 55-100 bp) in pseudolymphoma (Case 3) (Figure 2b), probably indicating a monoclonal amplification.

A band of 80 bp (predicted size range, 70-110 bp) was considered as evidence of amplification of a monoclonal population for the V γ 11/J γ 12 primer pair in high-grade T lymphoma (Cases 14 and 15); LIP (Cases 2, 8, and 10); lymphomatoid granulomatosis (Case 6); and Hodgkin's disease (Case 12) (Figure 2c). In contrast, no bands, and therefore no monoclonal amplification, were visualized using the V γ 101/J γ 12 primer pair (Figure 2d).

Table 1 depicts clinical data. The median age of the patients was 40 years (range, 2-81 years). There were 10 females and 5 males. Patients younger than 40 years presented greater CD15 expression in lymphomas ($p = 0.04$). All male patients presented VY11 expression, whereas 8 of the 10 female patients did not express VY11 ($p = 0.007$). No differences were detected in terms of immunophenotyping, gene rearrangement, gender, or age.

Table 2 depicts the results regarding tumor staining intensity for CD3, CD20, CD15, CD30, CD45, and CD68 immunophenotyping stratified by diagnosis (LIP or lymphoma). All 8 cases of LIP presented moderate to strong immunostaining for CD3, whereas only 2 cases of lymphomas and 1 case of pseudolymphoma presented similar immunostaining, and this difference was statistically significant ($p = 0.02$). All 8 cases of LIP also presented CD20 expression, as did 6 cases of lymphoma and 1 case of pseudolymphoma, with no differences in terms of the staining intensity. Similarly, no differences were observed for CD15, CD30, CD45, or CD68.

Table 2 depicts the results regarding gene rearrangement detection using the VH/JH, DB1/J β 2, V γ 11/J γ 12, and V γ 101/J γ 12 primer pairs stratified by diagnosis (LIP or lymphoma). Gene rearrange-

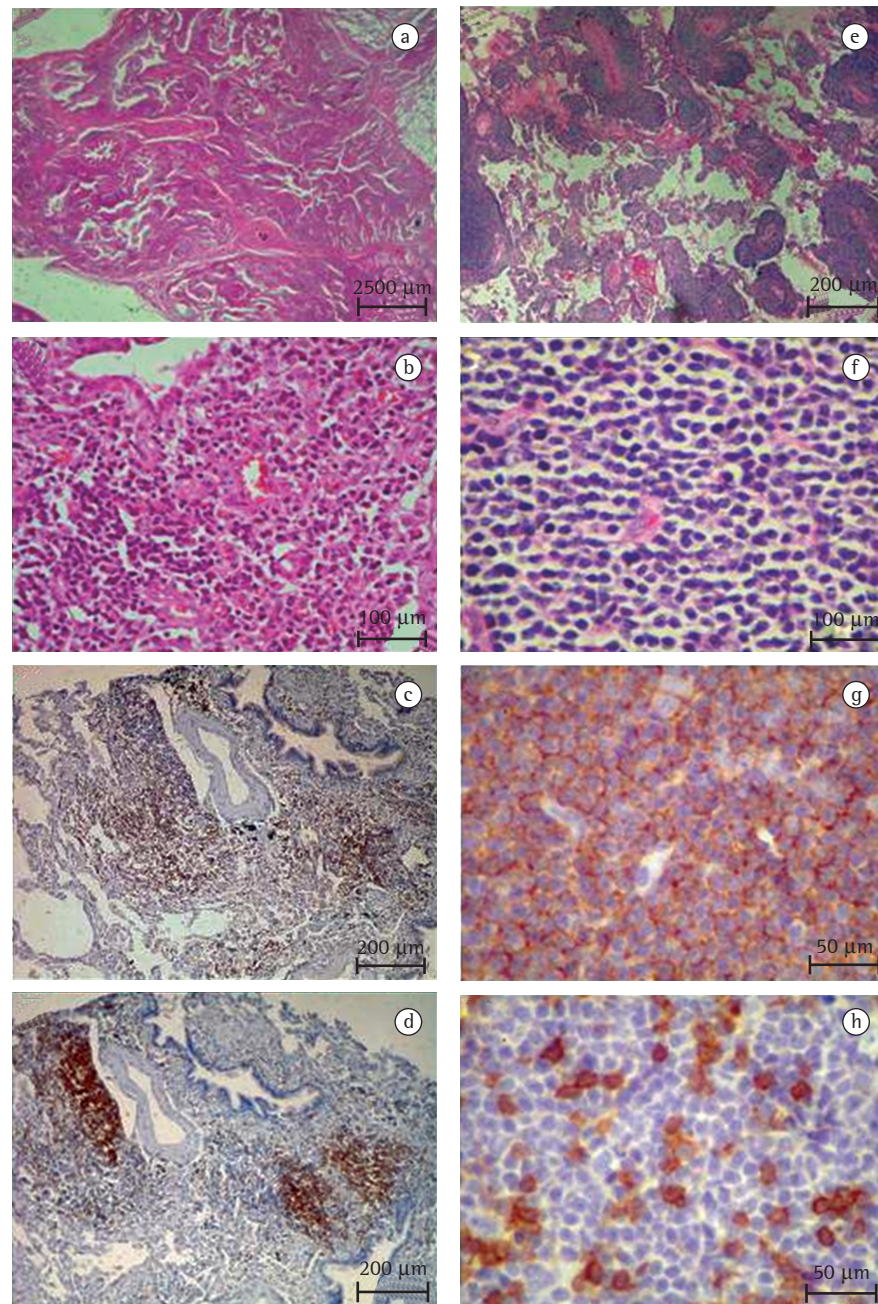


Figure 1 – (Panels a to d). Panoramic view of the histological pattern found in lymphoid interstitial pneumonia. Note the lymphoid infiltrate surrounding the broncovascular axis with extension to the alveolar septa (a), composed mostly of T and B lymphocytes and varying numbers of plasma cells (b). T lymphocytes (CD3 cells) can be seen primarily in the septal interstitium (c), whereas B lymphocytes (CD20 cells) are predominantly found in aggregates within lymphoid follicles and focally in thickening alveolar walls (d). a) H&E $\times 10$; b) H&E $\times 200$; c) CD3 $\times 100$; d) CD20 $\times 100$. (Panels e to h) Panoramic view of the histological pattern found in low-grade lymphoma (Case 13). Note the diffuse, dense, and monomorphous population of small lymphoid cells remodeling and infiltrating the lung (E). These cells present irregular nuclear contours (f). Neoplastic lymphoid cells staining for CD20 (g), indicates the B-cell phenotype with a variable reactive background population of CD3-positive T-cells (h).

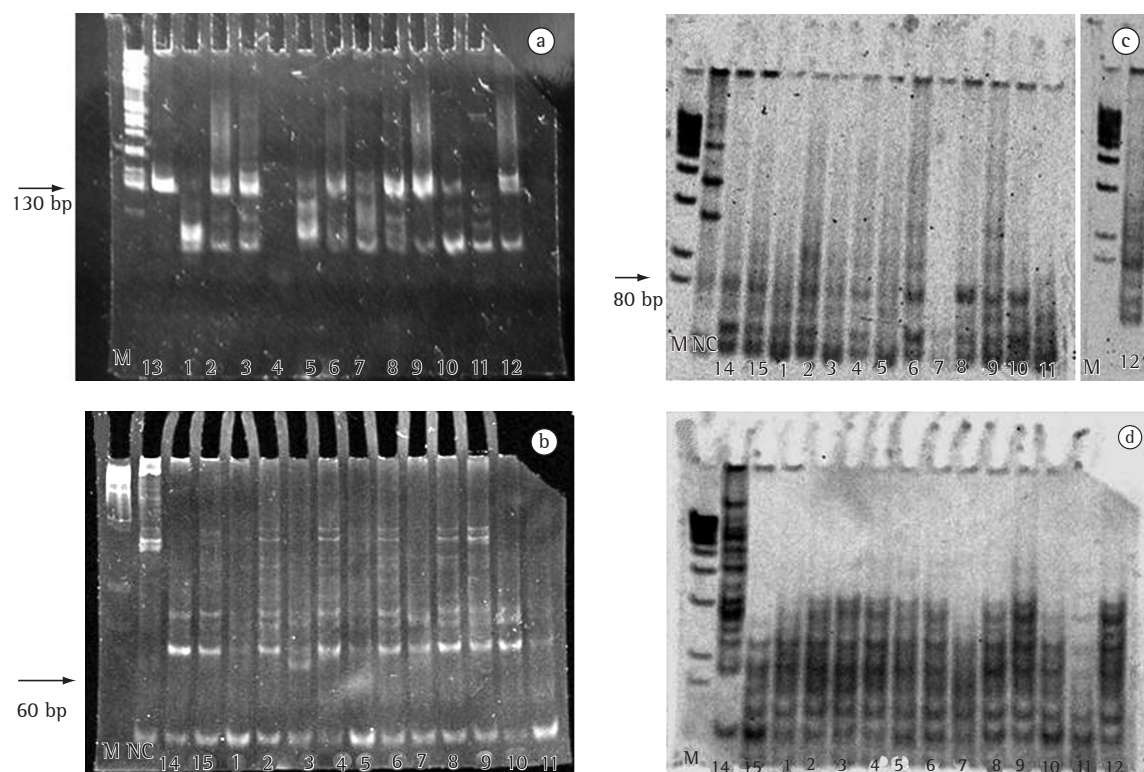


Figure 2 - a) IgH PCR-amplified products analyzed on 10% polyacrylamide gel and stained with ethidium bromide: lane 1 (Case 13); lane 2 (Case 1); lane 3 (Case 2); lane 4 (Case 3); lane 5 (Case 4); lane 6 (Case 5); lane 7 (Case 6); lane 8 (Case 7); lane 9 (Case 8); lane 10 (Case 9); lane 11 (Case 10); lane 12 (Case 11); and lane 13 (Case 12). Cases 13, 2, 3, 8, 9, and 12 show 130-bp bands, indicating a predominant clonal lymphocyte population. b) TCR β -chain PCR-amplified products: lane 1 (Case 14); lane 2 (Case 15); lane 3 (Case 1); lane 4 (Case 2); lane 5 (Case 3); lane 6 (Case 4); lane 7 (Case 5); lane 8 (Case 6); lane 9 (Case 7); lane 10 (Case 8); lane 11 (Case 9); lane 12 (Case 10); and lane 13 (Case 11). A band of ≈ 60 bp was observed in Case 3, probably indicating monoclonal amplification. c) TCR γ -chain PCR-amplified products: lane 1 (Case 14); lane 2 (Case 15); lane 3 (Case 1); lane 4 (Case 2); lane 5 (Case 3); lane 6 (Case 4); lane 7 (Case 5); lane 8 (Case 6); lane 9 (Case 7); lane 10 (Case 8); lane 11 (Case 9); lane 12 (Case 10); lane 13 (Case 11); and lane 14 (Case 12). Amplification of a monoclonal population using the V γ 11/J γ 12 primer pair was observed in cases 14, 15, 2, 6, 8, 10, and 12. d) TCR γ -chain PCR-amplified products: lane 1 (Case 14); lane 2 (Case 15); lane 3 (Case 1); lane 4 (Case 2); lane 5 (Case 3); lane 6 (Case 4); lane 7 (Case 5); lane 8 (Case 6); lane 9 (Case 7); lane 10 (Case 8); lane 11 (Case 9); lane 12 (Case 10); lane 13 (Case 11); and lane 14 (Case 12). No monoclonal amplification was observed using the V γ 101/J γ 12 primer pair.

ment was detected in 4 of the 8 cases of LIP, which changed the diagnosis from LIP to lymphoma, thus showing the importance of gene rearrangement detection in cases of LIP. In this situation, gene rearrangement using the VH/JH and V γ 11/J γ 12 primer pairs was detected in 3 and 1 cases of LIP, respectively, and no gene abnormalities were found using the D β 1/J β 2 and V γ 101/J γ 12 primer pairs in any of the cases.

Table 3 depicts the associations between immunophenotyping and gene rearrangement.

A significant positive association was found between the intensity of CD20 and CD68 expression and VH rearrangement ($p = 0.01$ and $p = 0.002$, respectively).

The relationships among clinical data, lymphoid/lymphoproliferative disorders, and overall survival were examined using the Kaplan-Meier method and the log-rank test prior to and after the gene rearrangement. No age- or gender-related differences were found. Prior to the gene rearrangement, the mean survival was 48 months for the 8 patients with

Table 2 – Lymphocyte immunophenotyping and gene rearrangement by tumor staining degree.

Diagnosis Initial/Final	Imunohistochemistry						Primers			
	CD3	CD15	CD20	CD30	CD45	CD68	VH/JH	Dβ1/Jβ2	Vγ11/Jγ12	Vγ101/Jγ12
LIP/LIP	2	NP	1	NP	NP	NP	0	0	0	0
LIP/LGB	3	0	2	0	2	2	1	0	1	0
PSLY/LGB	4	0	2	0	4	3	1	1	0	0
LIP/LIP	2	3	1	1	2	1	0	0	0	0
LIP/LIP	3	NP	2	NP	NP	NP	0	0	0	0
LYG/LGB	1	2	2	0	3	3	1	0	1	0
LGBCL/LGBCL	1	0	1	0	2	2	0	0	0	0
LIP/LGB	3	0	2	0	3	1	1	0	1	0
LIP/LGB	3	0	2	0	3	1	1	0	0	0
LIP/LGT	2	1	1	1	0	2	0	0	1	0
LIP/LIP	2	0	1	0	0	0	0	0	0	0
HD/HD	0	2	0	3	2	1	1	0	1	0
LGB/LGB	1	0	4	0	4	1	1	0	0	0
HGT/HGT	3	NP	0	1	NP	NP	0	0	1	0
HGT/HGT	4	0	0	1	0	0	0	0	1	0

LIP: lymphoid interstitial pneumonia; LGB: low-grade B-cell lymphoma; LGT= low-grade T-cell lymphoma; LGBCL: low-grade B-cell lymphoma with plasmacytic differentiation; LYG: lymphomatoid granulomatosis; NHL: non-Hodgkin's lymphoma; NP = not performed (scanty material originated from transbronchial biopsy); OLB: open lung biopsy; PSLY: pseudolymphoma; Status: 0 = dead, 1 = alive; and TBB: transbronchial biopsy.

LIP and 46 months for the 7 patients with lymphoma and pseudolymphoma. After the gene rearrangement, although not statistically significant, the mean survival for the patients with LIP was 58 months, compared with 30 months for patients with lymphoma (log-rank = 2.64; $p = 0.10$). Prior to the gene rearrangement, 4 patients with LIP died quickly, whereas only one patient with LIP died after the gene rearrangement. This difference did not achieve statistical significance (log-rank = 0.20; $p = 0.65$).

Discussion

The position of LIP within classification systems has changed with advances in the understanding of the nature of pulmonary lymphocytic infiltrates, and many groups prefer to classify LIP under the heading of pulmonary lymphoproliferative disorders.⁽²⁷⁾ A diagnosis of pulmonary lymphoproliferative disorders can be evoked in many clinical conditions, such as LIP, pseudolymphoma, primary lymphoma, and pulmonary localization of an otherwise extended NHL.

In our study, we found that LIP was characterized by marked lymphocytic infiltration with extensive involvement of the alveolar septa, similar to that described by other authors in 1969,⁽²⁸⁾ contrasting with the diffuse, dense and monomorphous infil-

tration of small lymphoid cells in lymphomas. All 8 cases of LIP presented significant moderate-to-strong CD3 immunostaining, whereas only 2 cases of lymphoma and 1 case of pseudolymphoma had similar immunostaining. In contrast, CD20, CD15, CD30, CD45, and CD68 immunostaining was seen in nearly all cases. Although CD20, CD45, and CD68 antibodies are routinely employed to determine clonality in different types of lymphoproliferative disorders, our findings demonstrated that only CD3 was specifically related to cases of LIP. This finding suggests that the differential diagnosis between LIP and lymphomas is difficult due to the (not always detectable in routine practice) presence of clonal B-cell or T-cell populations in lymphocytes. There have been a few reports in the literature of clonal B-lymphocyte or T-lymphocyte populations in the pulmonary tissue of patients with B-cell or T-cell pulmonary lymphoma.^(11,22,24,29) In this context, the PCR technique is promising.

In fact, by employing the PCR technique to detect complete IgH rearrangements and analyzing TCR β- and TCR γ-chain using two different primer combinations such as Dβ1/Jβ2 and Vγ11/Vγ101/Jγ12, we found that the VH/JH primer pair indicated the presence of a predominant clonal lymphocyte population in high-grade T lymphoma

Table 3 – Correlations between immunophenotyping and gene rearrangement.

Lymphocyte immunophenotyping		Gene rearrangement			
		VH/JH	D β 1/J β 2	V γ 11/J γ 12	V γ 101/J γ 12
CD3	Correlation coefficient	-0.048	0.415	0.064	0.0
	(two-tailed)	0.865	0.124	0.821	0.0
CD15	Correlation coefficient	-0.117	-0.208	0.259	0.0
	(two-tailed)	0.718	0.516	0.416	0.0
CD20	Correlation coefficient	0.605	0.229	-0.311	0.0
	(two-tailed)	0.017 ^a	0.412	0.260	0.0
CD30	Correlation coefficient	-0.431	-0.224	0.431	0.0
	(two-tailed)	0.142	0.462	0.142	0.0
CD45	Correlation coefficient	0.786	0.452	-0.300	0.0
	(two-tailed)	0.002 ^b	0.140	0.344	0.0
CD68	Correlation coefficient	0.334	0.458	0.101	0.0
	(two-tailed)	0.289	0.134	0.754	0.0

^aSignificant at the 0.05 level (two-tailed); and ^bSignificant at the 0.01 level (two-tailed).

(Case 13), LIP (Cases 2, 8 and 9), pseudolymphoma (Case 3), lymphomatoid granulomatosis (Case 6), and Hodgkin's disease (Case 12). The analysis of TCR β -chain gene rearrangement using the D β 1/J β 2 primer pair identified monoclonal amplification in pseudolymphoma (Case 3). Amplification of a monoclonal population using the V γ 11/J γ 12 primer pair was found in high-grade T lymphomas (cases 14 and 15); LIP (Cases 2, 8, and 10); lymphomatoid granulomatosis (Case 6); and Hodgkin's disease (Case 12). These results suggest that VH/JH and V γ 11/J γ 12 are the best combinations for detecting gene rearrangement in cases of lymphoma that present a histological pattern consistent with LIP.

Our findings underscore the suggestion that, since many cases of LIP evolve to lymphoma, LIP should be considered preneoplastic or seen as a true lymphoma if any gene rearrangement is detected. In fact, our study showed that 4 of the 8 cases of LIP were cases of low-grade B-cell or T-cell lymphoma and cases of Hodgkin's disease from the outset.

We also found that it is difficult to draw the distinctions among idiopathic LIP, lymphoma, and Hodgkin's disease based solely on the routine analysis of histological sections,⁽³⁰⁾ which should not be undervalued but rather should be complemented with immunophenotyping and PCR analysis, thus

contributing to better identification/classification of lymphoma cases. In fact, we found a significant positive association between immunohistochemistry and the PCR technique (Table 3). In addition, our study showed that immunohistochemistry and molecular analysis can separate reactive and neoplastic infiltrates, and that 50% of the cases of LIP were actually found to be malignant transformations, including Hodgkin's disease (a variant of lymphocytic predominance). We also confirmed that pseudolymphoma actually represents a low-grade lymphoma. These findings call for modifications in the therapeutic protocol employed for this subgroup of patients. However, this might not occur in all cases of LIP, suggesting that some cases of LIP meet the criteria for benign status and should be treated only with steroids, whereas cases of LIP with gene rearrangement detection should be seen as lymphoma cases and treated with chemotherapy.

In this study, we emphasized the diagnosis of LIP using immunohistochemistry and PCR analysis of the IgH gene rearrangement. This study is very important for oncologists, pulmonologists, and pulmonary pathologists because it allows the establishment of the correct diagnosis of lymphoma and the introduction of the appropriate treatment for

the disease, as well as, perhaps, a better chance of survival for these patients.

Since LIP and primary lymphomas of the lung are rare, accounting for less than 1% of all lung pathologies in most studies, the small size of our study sample was to be expected. However, the clinical impact of our findings was investigated. No differences were detected in terms of immunophenotyping, gene rearrangement, gender, or age. We also examined survival curves prior to and after the gene rearrangement. The differences did not achieve statistical significance, probably due to the limited number of patients and to the fact that the clinicians were unaware of the fact that some of the cases of LIP were lymphomas. The diagnosis of lymphoma in some of the cases of LIP was made only after the gene rearrangement. The analysis of the survival curves prior to and after the gene rearrangement revealed that 4 LIP patients died quickly prior, whereas only 1 LIP patient died after, confirming that the more aggressive stage of the disease occurs prior to the gene rearrangement.

Regardless of the lymphocytic/lymphoproliferative pathogenetic mechanism, detection of monoclonal B and T cells by immunophenotyping and PCR had an impact on the diagnosis of pulmonary lymphomas in patients previously diagnosed with LIP. Therefore, immunophenotyping and PCR should be used as the 'gold standard' techniques in routine practice. Interpretable PCR results were obtained in the majority of the cases analyzed, demonstrating that our PCR analysis could become a routine procedure. The detection of gene rearrangement in lung biopsies, especially in cases diagnosed as LIP, is very important for the establishment of an accurate diagnosis of lymphoma. In order to determine whether or not these cases represent malignant transformation from LIP, randomized, prospective trials involving large patient samples are needed. In addition, we believe that further studies of lymphoid/lymphoproliferative disorders are warranted in order to validate the results of our immunophenotyping and gene rearrangement analysis.

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