A Basic Approach to Lymph Node and Flow Cytometry Fine-Needle Cytology

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**Key Words**
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**Abstract**
According to the World Health Organization (WHO), the new classification of lymphomas is mainly based on morphological, immunophenotypical, and molecular criteria. Consequently, this new approach has led from the substantial role that architecture played in the past to a secondary panel highlighting the role of fine-needle biopsy (FNB). Applied together with other ancillary techniques, such as flow cytometry (FC), FNB is a potential tool for the diagnosis of lymphomas, and enlarged lymph nodes represent an excellent target for the implementation of this technique. Despite the difficulties inherent in this technology, which might pose problems in differential diagnosis, in the majority of cases this joint work allows an accurate diagnosis of malignancy and even correct subcharacterization in routine lymphomas. Additionally, in selected cases, other molecular techniques like FISH and PCR can also be performed on FNB specimens, helping in the characterization and diagnosis of lymphomas. In this review, we discuss the basic aspects of the combination of FNB cytology and FC in the diagnosis and subclassification of lymphomas. The preanalytical phase is extensively discussed. The advantages, disadvantages, and technical limitations of this joint work are addressed in general and in terms of the accurate subclassification of lymphomas.

**Introduction**

The new classification of lymphomas of the World Health Organization (WHO) classification of tumors of hematopoietic and lymphoid tissue is based on morphological criteria, immunophenotype, and molecular characteristics of lymphoid cells in the appropriate clinical setting [1]. As a consequence of this, the substantial role that architecture played in the past in the classification of lymphomas has become, following the above mentioned approach, secondary in the majority of routine lymphoma diagnoses.

This new approach to the diagnosis of lymphomas highlights the enormous diagnostic potential that fine-needle biopsy (FNB) cytology has when applied together with other ancillary diagnostic techniques, such as immunocytochemistry, flow cytometry (FC), cytogentic, and molecular techniques (FISH, CISH, and PCR), which
nowadays are accessible to most laboratories [2, 3]. Furthermore, in the majority of cases, lymphadenopathies are benign/reactive and, among those that have a diagnosis of lymphoma, B-cell lymphomas represent 90% of the non-Hodgkin lymphomas. Of the latter, more than half are diffuse large B-cell lymphomas (DLBCL) or follicular lymphomas (FL) (50 and 20%, respectively) [4]. All of these facts highlight the huge potential of FNB and FC together as diagnostic tools for the primary assessment of lymph nodes.

FNB is a simple procedure in which a needle is inserted into a site and a small amount of tissue is sucked out to provide a sample of cells for laboratory analysis. When performed by trained operators and for the correct indications, this is one of the safest and most minimally invasive, cost-effective, and accurate diagnostic techniques.

FNB has the following advantages over other techniques, histology: (1) it achieves quicker results, (2) it is better tolerated by the patient, (3) through rapid on-site evaluation (ROSE) it allows triaging and selection of material for other ancillary tests and ensures that the needle is inserted into a lymph node, (4) it provides a better cytology and better rinsing for FC and molecular biology, and (5) it is less expensive.

Enlarged lymph nodes represent an excellent target for the implementation of this technique for the diagnosis of both benign and malignant lymphadenopathies. Despite the numerous publications on this subject stressing the enhancement of FNB and FC sensibility and specificity as a first-approach technique for the diagnosis of enlarged lymph nodes and lymphoma [5–7], the prevailing assumption is still that FNB should only be used in cases in which an excisional biopsy is medically contraindicated, in follow-up, in relapses of previously characterized lymphomas, or to rule out metastatic and reactive causes. In most guidelines of lymphoma protocols, histology is still considered the gold standard for the morphological diagnosis of lymphoma [5, 8, 9].

The main arguments against the use of FNB for lymphoma diagnosis insist on the fact that FNB does not provide an architecture assessment, precluding accurate subclassification or grading in some lymphoma subtypes. As an example, there is often a reference to the impossibility of properly grading FL by FNB. FL is graded from 1 to 3 based on the number of centroblasts per high-power field (×40) in 10 representative neoplastic follicles. Differentiation between low grade (grades 1–2) and high grade (grade 3) is important not only for the prediction of tumor behavior but also for therapeutic decisions [1]. The aforementioned criteria do not prevent the diagnosis of FL by FNB but can in some cases limit accurate characterization and determine, in a second step, the need for histology. The loss of architecture is also pointed out as an obstacle in the identification of the more recently emerging diagnosis of in situ follicular neoplasia and in situ mantle cell lymphoma (MCL). Another argument against FNB is the need to store material for upcoming molecular studies with the aim of identifying new biological markers for diagnosis or for new therapeutic options.

FC is an ancillary technique that can be easily coupled with the FNB technique, allowing a more accurate and reliable determination of the immunophenotype of lymphoid populations. Immunophenotyping by FC is nowadays recognized for its capacity to distinguish abnormal populations from reactive ones, allowing objective classification and quantification of the cells. The ability to perform a complete basic phenotype in limited cytology specimens makes it the ideal technique for combined application with FNB in the diagnosis of lymphoma.

FC has the ability to be operational in small samples, and it has some important advantages over immunohistochemical stains. Multiple phenotypic and 2 physical (size and complexity) parameters of individual cells can be simultaneously assessed to identify both major and minor abnormal populations. FC has the capacity to evaluate multiple parameters in a single cell, such as co-expression of CD5 and CD23 in chronic lymphocytic lymphoma (CLL). FC remains the best way to identify surface immunoglobulins and evaluate monoclonality in lymphoid B-cell populations by analyzing the immunoglobulin κ and λ chain ratio. FC is also able to evaluate the mean fluorescence intensity (MFI) of antigen markers, which can have diagnostic implications, as is the case of CD20 and surface immunoglobulin low-intensity, which characterizes CLL. FC also facilitates the identification of aberrant T-cell phenotypes, helping in the cytological diagnosis of T-cell lymphomas. FC is also crucial for the identification of minor populations of neoplastic cells on a background of reactive lymphocyte populations, in a partly involved lymph node by lymphoma, that otherwise would go undetected by immunohistochemistry (IHC). Recently, the possibility of detecting T-cell clonality with the aid of a broad panel of antibodies against the variable region of the TCRβ gene was reported [10, 11].

Beyond the pros and cons previously mentioned regarding FNB/FC, this tandem approach has limitations that have been pointed out and should always be taken into account. FNB requires an experienced operator and...
in many hospitals there is a limited availability of cytopathologists. Another problem is the availability of ancillary tests in many departments. FNB of lymph nodes should be carried out with ROSE in order to evaluate the adequacy of the collected material and to perform a proper specimen triage. This procedure is time consuming and requires commitment of the professionals.

Conversely, false-negative results can be due to error or inadequate sampling, i.e. in lesions with marked fibrosis, extensive necrosis, or partial involvement of the lymph nodes. These limitations can be avoided by doing multiple passes and ensuring sampling of different areas of the node. Errors can occur in the interpretation of the smears, as well as in the interpretation of FC results. Errors can, furthermore, be due to sample representation or paucity of diagnostic cells such as Reed-Sternberg cells in Hodgkin lymphoma (HL). Attempts to diagnose HL in FC have long been unsuccessful, and there is still no consensus essentially due to the lack of specific markers. Recently, some authors have demonstrated that Reed-Sternberg cells can be detected by FC and that classical HL (cHL) can be diagnosed by FC with a high sensitivity (88.7%) and specificity (100%), and it can be distinguished from other putative differential diagnosis like T-cell-rich large B-cell lymphoma and anaplastic lymphoma [12, 13].

Difficulties in diagnosis also emerge in distinguishing some exceptional sets, i.e. HL versus mediastinal primary B-cell lymphoma (MPBCL), or pleomorphic T-cell lymphomas versus T-cell-rich large B-cell lymphoma or anaplastic large-cell lymphoma (ALCL), and also due to the fact that the lymphoma immunophenotypic profile can occasionally be atypical, hampering recognition [14]. In some lymphomas, as in the case of HL, some DLBCL, and ALCL, FC has had a limited role in the diagnosis. False-negative results (reporting only to subsidiary reactive populations) have been assigned to the shortage of neoplastic cells present in these lymphomas, to cellular lysis during preparation, and to immunophenotypic background noise, conditioned by the subsidiary cell population (T-cell rosettes surrounding Reed-Sternberg cells), or due to the fact that neoplastic cells are too large to be detected by the flow cytometer [13, 15]. The limitations of FC can be reduced in some situations by the complementary role of IHC as an alternative way to characterize cell populations. Some antigens, such as cyclin D1, are better evaluated by IHC. IHC also has the main advantage of providing simultaneous morphological control. In summary, FC and IHC are techniques that complement each other in the immunohistocharacterization of FNB lymphoid populations, reducing the risk of a misdiagnosis.

Histology and immunohistochemical stains are not devoid of interpretation problems and pitfalls. Evaluation of immunostains is a major challenge, with a moderate number of uninterpretable cases. Some antibodies, like CD103 and CD11c, useful in the diagnosis of hairy cell leukemia and nodal marginal zone lymphoma (MZL), cannot be used in paraffin sections.

Besides FC, other molecular techniques like FISH and PCR can also be performed with good results in FNB specimens as an alternative way to detect clonal populations by detecting IGH locus translocation, in the evaluation of T-cell receptor rearrangements, or to detect specific translocations or deletions that help characterize a high proportion of lymphomas (table 1) [2]. Some of these molecular alterations also have important implications for prognostic evaluation and for the patient’s selection of treatment, as in the case of 13q14, 11q, and 17q deletion, associated with disease progression in CLL. The detection of specific cytogenetic aberrations is cardinal for the diagnosis of MCL, Burkitt lymphoma (BL), and FL [16–18].

As mentioned before, the ancillary techniques discussed in this review, which can contribute to the diagnosis and subclassification of lymphomas by FNB, are numerous and currently accessible to all of the main lymphoma diagnostic reference centers. However, they are costly and therefore should be used sparingly. In routine clinical practice, it is neither necessary nor cost effective to perform multiple tests in every specimen. Common sense and the cytopathologist’s experience should determine how exhaustive the cytological diagnosis should be. It is also up to the cytopathologist to make a decision regarding the need to provide a comprehensive diagnosis or just an operative diagnosis, pushing a more accurate diagnosis for the histology. The effectiveness of FNB in the diagnosis of lymphomas depends on multidisciplinary team work to ensure an adequate sample and correlation between morphology and other ancillary techniques. The final report of an FNB lymphoid neoplasm should include the results of the several ancillary techniques that have been used to achieve the diagnosis, and a final interpretative conclusion should be provided [19].

**Indications for FNB of Lymph Nodes**

Although FNB may be used in a persistent reactive lymphadenopathy just to reassure the patient, this is not an optimal indication. FNB should preferably be confined to cases with a strong clinical suspicion of a specific infection such as tuberculosis or neoplasia.
Enlarged lymph nodes are more likely to be malignant in adults than in children. Furthermore, a solitary enlarged lymph node is more likely to be malignant. Enlarged lymph nodes in the posterior cervical triangle and in the supraclavicular area are more prone to being malignant than enlarged lymph nodes in other areas.

There are few, if any, true contraindications for the performance of FNB of lymph nodes if one excludes the uncooperative-patient situation. In patients with a bleeding diathesis, FNB is the safest modality to achieve a diagnosis. Pressure to the puncture site will reduce, if not prevent, hematoma formation.

FNB: Technical Aspects

FNB and Sample Preparation

FNB applies to palpable, superficial lesions as well as to deep lesions. Deep lesions can be dealt with the assistance of ultrasound and image-based guidance for placement of the needle. In most situations, FNB is performed by a radiologist or a pathologist. In our opinion, the best results are obtained when this technique is performed by a pathologist. FNB of lymph nodes requires more commitment than any other FNB done at any other location, and nothing is better than having the same expert responsible for the collection of the sample and for its assessment. FNB should be a bloodless technique. To perform FNB, any technique is suitable, provided that the operator obtains satisfactory results. In our experience, this task is facilitated through performance of a nonaspirative FNB technique, allowing, even for an unskilled operator, the achievement of good results and minimizing blood contamination. This nonaspirate technique, also called the Zajdela or fine-needle capillary method, relies on capillary action to draw the sheared cells into the small-caliber needle [20]. This technique gives the operator greater control over the target and reduces bleeding.

Adequate evaluation of the lymphoid cytomorphology relies on high cellularity samples and excellent technical preparation. Preparations can be air dried or sub-

Table 1. Immunophenotype and genetic diagnostic/prognostic criteria for lymph node, current chronic B-cell lymphomas

<table>
<thead>
<tr>
<th>Immunophenotype</th>
<th>Cytogenetic abnormalities</th>
<th>Clinical implications</th>
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<tbody>
<tr>
<td>B-cell LL</td>
<td>CD45+, TDT+, CD20+/−, CD79b+, CD43+, CD34+, CD10+/−</td>
<td>t(8;13); t(1;14) t(10;14); t(5;14)</td>
</tr>
<tr>
<td>CLL</td>
<td>Igw, CD79b+, CD45+, CD19+, CD20+, CD5+, CD23+, CD200+, CD43+, CD10−</td>
<td>t(14;19); del; 11q23 10−20%, del 13q14.3 50−60%, del 17p10%; del 6q Prognostic markers</td>
</tr>
<tr>
<td>LPL</td>
<td>CD45+, CD19+, CD20+, CD5−</td>
<td>t(9;14) 50%</td>
</tr>
<tr>
<td></td>
<td>CD23−, CD10−, CD103−</td>
<td>MYD88 LP265P</td>
</tr>
<tr>
<td>MCL</td>
<td>CD45+, CD79b+, CD19+, CD20+, CD5+, CD23−, CD43+, CD200−</td>
<td>t(11;14) 90% Aggressive behavior</td>
</tr>
<tr>
<td>MZL nodal</td>
<td>CD45+, CD19+, CD20+, CD5−, CD10−, CD23−, CD103−, CD11c+, CD43 50%</td>
<td>Trisomy 18; 3 1q21; 1q34</td>
</tr>
<tr>
<td>FL</td>
<td>CD45+, CD19+ w, CD20+b, CD10+, CD38+, CD43−</td>
<td>t(14;18) 90%</td>
</tr>
<tr>
<td>BL</td>
<td>CD45+, CD19+, CD20+, CD10+, CD43+, CD38++, Bcl2−</td>
<td>t(8;14); t(2;8); t(8;22)</td>
</tr>
<tr>
<td>DLBCL</td>
<td>CD45+, CD19+, CD20+, CD5+ 10%, CD10 30−60%</td>
<td>t(14;18)-myc rearrangement 30% 3q and 18 abnormalities, del 6, t(14;18) 1−5%, t(3;14) 30%, t(9;14), t(2;17) 1−1%, t(2;5)-rare Aggressive behavior</td>
</tr>
<tr>
<td>HCL</td>
<td>CD45+, CD19+, CD20+b, CD103+, CD22++, CD11c+, CD25+</td>
<td></td>
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</table>

B-cell LL = B-cell lymphoblastic lymphoma; W = weak immunoexpression; B = bright expression.
merged in alcohol at 95°C for subsequent staining with Giemsa or Papanicolaou and hematoxylin-eosin (H&E), respectively. A wide cuvette with 95°C alcohol should be prepared in advance so that the smears can enter readily and without tilting into the alcohol to avoid cell drying. FNB should be performed using a 25 or higher gauge needle, with or without a mandrel. The choice of needle depends on the dimension of the lymph node and its depth. Shorter needles are easy to handle; however, they are devoid of a mandrel, hampering the delivery of the sample for other purposes. Longer needles, in general, have a mandrel, allowing a slow delivery and spread of the material, as the mandrel is pushed along the needle, for smearing and collection in Eppendorf tubes. The material collected in Eppendorf tubes can be further used in the preparation of cytospins and cell blocks or for ancillary techniques.

In palpable lesions, the lymph node should be blocked between 2 fingers (i.e., index and middle); in the case of axillary lymph nodes, the lymph node can be immobilized while pressing it against the thoracic wall with the fingers. The distance between the skin and the target should be shortened. The needle should slowly enter the lymph node, and small and slow movements (about 5–10) should be done back and forth, changing direction and avoiding bleeding. The needle direction can be changed by moving the nodule while pressing one or the other finger more intensely.

Material should not be allowed to flood into the needle cannon but only to occlude the hub of the needle. Whenever blood reaches the hub of the needle, the pass should be stopped immediately; blood damages the smear and compromises a second pass.

When dealing with a hard, sclerotic lymph node, the needle movements should be more vigorous. Small lymph nodes (1 cm or less) may be easier to approach with a shorter needle.

In deep lymph nodes, FNB must be performed under ultrasound control. The shortest and most direct path for the needle should be chosen, obviating the risk of deviation. Attention should be paid regarding the excessive use of ultrasound gel. Gel should be applied to the ultrasound probe and this should be then covered with film to avoid contamination of the sample by the gel and to prevent observation in Giemsa-stained smears. Alcohol sprayed abundantly onto the skin can be used to permit the probe to slide.

ROSE for adequacy of the material, preliminary diagnosis, and triage for further ancillary techniques should be done (fig. 1). This observation can be accomplished using a rapid Wright-Giemsa type stain (e.g., Diff-Quik) in air-dried smears. Additional material should always be collected for different purposes (cytospins/cell blocks, FC, and FISH/PCR) in case it is needed for a differential diagnosis. Material for further ancillary techniques, i.e., other than FC, should be collected in an Eppendorf tube (table 2).

![Fig. 1. FNB and sample management. PAS = Periodic acid-Schiff.](image-url)
Cell blocks or cytospins should be always prepared for immunostains, useful in some differential diagnoses (e.g. cyclin D1, CD30, CD15, and PAX 5, among others) or in the evaluation of Ki 67. Material should be collected in PBS-buffered solution at pH 7.4. Both immunoalkaline phosphatase and immunoperoxidase immune techniques can be used on cytospins [21]. Material for cytogenetics or molecular analysis will also allow the search of specific translocations that characterize some lymphomas or evaluation of the molecular alterations that condition the selection of therapy for evaluation of the prognosis (table 2).

### Smear Staining and Evaluation

Diff-Quik/Giemsa and H&E/Papanicolaou are complementary stains that should be used whenever the amount of collected material allows. Prompt Diff-Quik/Giemsa staining permits ROSE to be done in a shorter time. This stain highlights background details and emphasizes the heterogeneity and polymorphism of the cell population. It highlights the presence of fibrous connective tissue, cellular aggregates, eosinophilic cytoplasmic granulation, and other cytoplasm characteristics. H&E/Papanicolaou staining provides detailed information about the characteristics of the nucleus.

Most lymphadenopathies are mainly due to metastasis, inflammatory/infectious sets, or nonspecific hyperplasia. These clinicopathological sets are generally easily diagnosed with the joint work of FNB plus FC and immunostains. Zardawi et al. [22] reported a 99% sensitivity in malignancy diagnosis, a 0% false negative rate, and a 100% positive predictive value for malignancy diagnoses using the combination of FNB and FC [22, 23].

Two main sets should be considered in FNB diagnosis of a lymph node: nonneoplastic and neoplastic. The first set includes nonspecific reactive changes, inflammatory, specific infectious lesions, suppurative or granulomatous lymphadenitis, and, less frequently, lymphadenopathies of an unknown etiology like Kikuchi or Rosai-Dorfman disease. Histochemical stains like Gram, Grocott, and Ziehl-Neelsen can be performed to identify a putative infectious agent. Material can also be sent for microbiology tests or for PCR-based DNA methods to identify microorganisms.

In the neoplastic setting, the distinction between metastatic and malignant lymphoproliferative diseases should be made. Metastatic tumors, mainly isolated cell carcinomas, small-cell tumors, small-cell carcinoma/neuroendocrine tumors in adults, and small-cell sarcomas in children, can pose difficult differential diagnostic problems which can be solved with the help of immunocytochemistry. In these contexts FC has little to add, and it is not useful for diagnosis. Nevertheless, some information can be retrieved if, while doing ROSE, misinterpretation of the smear determines that a sample has been collected for FC. Many of these small-cell tumors/carcinomas have neuroendocrine/neuroepithelial differentiation and are CD56 positive and CD45 negative and easily detected in FC.

The interpretation of FNB smears in lymph nodes suspicious for lymphoproliferative disorders will be discussed subsequently.

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**Table 2. Ancillary techniques that can be associated with FNB: aim, type of collection medium and preanalytical details**

<table>
<thead>
<tr>
<th>Technique</th>
<th>Aim</th>
<th>Material collection</th>
</tr>
</thead>
<tbody>
<tr>
<td>FC</td>
<td>Immunophenotyping</td>
<td>PBS¹, TransFix², RPMI¹, EDTA¹</td>
</tr>
<tr>
<td>Cytogenetics</td>
<td>Karyotyping</td>
<td>Saline solution, RPMI¹, ³</td>
</tr>
<tr>
<td>FISH</td>
<td>Evaluation of molecular alterations, translocation, deletions</td>
<td>Cytospins, cytoblocks, needle wash in saline solution, RPMI, EDTA</td>
</tr>
<tr>
<td>CISH</td>
<td>Evaluation of molecular alterations</td>
<td>Cytospins, cytoblocks</td>
</tr>
</tbody>
</table>

¹ The material must be processed during the first 6 h. ² The material can be processed in 3 or 4 days. ³ Collection in sterile conditions.
Evaluation of a Smear Suspicious for Lymphoma

Despite the impossibility of having a preserved architecture in a cytological smear, a good sample and a perfect smear will permit determination of whether lymphocytes lay in a diffuse towel or there are different morphological areas with cellular aggregates (representative of a germinal center), fibrotic fragments, or histiocytic aggregates (representative of a sinus histiocytosis). These details are, in some sets, a clue to a differential diagnosis; numerous histiocytes with epithelioid features, single or in small granulomas/aggregates, can point to an HL (fig. 2a), a T-cell lymphoma (fig. 2b), or even a seminoma (fig. 2c). A hard lymph node with fibrosis will provide a hypocellular smear which might indicate, in the proper clinical context, the possibility of HL (nodular sclerosis) or a primary mediastinal large B-cell lymphoma. Subsidiary cells dispersed in the background, like eosinophils or mast cells, can point to specific diagnoses such as HL, T-cell lymphoma, and lymphoplasmacytic lymphoma (LPL). Lymphohistiocytic aggregates with or without associated macrophages with tingible bodies are observed in follicular reactive hyperplasia, but they can also be present in some FL, in MCL, and in MZL. Otherwise, these aggregates of follicular dendritic cells should not be visible in CLL; instead, more clarified areas can be identified in the smear of a CLL, composed of prolymphocytes and paraimmunoblasts (fig. 3a).

Lymphoid lineage cell populations identifiable on an FNB smear of a lymph node are composed of small round lymphocytes (naïve or memory cells), cleaved centrocytes, centroblasts, immunoblasts, prolymphocytes, paraimmunoblasts, plasmablasts, plasmacytoid lymphocytes, plasma cells, and monocytoid lymphocytes. When evaluating lymphocytic populations, two characteristics should be assessed: the homogeneity of the cell population and whether the lymphocytic population is monomorphous with a prevailing cellular type or there is a polymorphous population with a mixture of cells (small lymphocytes, centrocytes, centroblasts, immunoblasts, or plasma cells). A heterogeneous distribution and a polymorphic lymphocytic population, with cellular aggregates representative of germinal centers, usually point to nonspecific reactive follicular hyperplasia. Nevertheless, reactive hyperplasia may be quite varied, and it may be composed of monotonous smears with a predominance of small lymphocytes or else a predominance of large cells and immunoblasts, sometimes even with atypical forms (common in clinical conditions associated with viral infections, HIV, or the Epstein-Barr virus) corresponding to para-cortical hyperplasia (fig. 4). In this situation, the differential diagnosis of HL or T-cell lymphoma should be raised. Immunostains and FC may be employed in the characterization of these cases.

Some lymphomas, mainly FL and MZL, can also share a polymorphous and heterogeneous cytomorphology that simulates a reactive pattern (fig. 5a). In contrast, a monotonous and monomorphous lymphocytic population is highly suspicious for malignancy.

Another detail that should be evaluated in lymphoid populations is cell size. Lymphomas can be classified depending on whether they are composed of small (equal to or smaller than a resting lymphocyte or double the size of an erythrocyte – small naïve lymphocytes, centrocytes, and prolymphocytes), medium-sized (centroblasts and plasmablasts, or large cells (2 or 3 times the size

Fig. 2. Histiocytes with epithelioid features, isolated or in small granulomas/aggregates, can point to an HL (Giemsa, ×400; a), a T-cell lymphoma (Giemsa, ×200; b), or even a seminoma (H&E, ×400; c).
Fig. 3. a CLL composed of small CLL typical lymphocytes with coarse chromatin, prolymphocytes, and paraimmunoblasts. H&E. ×600. b Immunophenotyping showing an abnormal B-cell (light blue) population expressing CD20\(^{\text{W}}\), CD5\(^{+}\), CD23\(^{+}\), CD43\(^{+}\), CD79b\(^{\text{W}}\), CD200\(^{+}\), with light chain \(\lambda\) restriction. Color refers to the online version only.
of a resting lymphocyte-immunoblast) (fig. 6). Nuclei should also be evaluated according to whether they are round, cleaved, or polilobulated and whether the chromatin is fine like in lymphoblasts or coarsely condensed as in CLL or BL. The nuclear membrane is also useful for characterization of lymphomas – MCL and FL are constituted by cleaved small centrocytes.

It is also essential to have knowledge of the lymphocytic evolution and transformation throughout their several sequential biological compartments. Each of these compartments is characterized by a specific morphology and immunophenotype and is the cell of origin of specific subtypes of lymphomas [24].

Small/Intermediate-Sized Cell Lymphomas

Small and intermediate-sized B-cell lymphomas encompass several differential diagnoses such as reactive lymphoid hyperplasia, as previously addressed, CLL, FL (grade 1–2), MCL, LPL, and MZL. All of these pathological entities share, at a low magnification, a similar morphological pattern which is characterized by small/intermediate lymphoid cells. However, at a high magnification, distinct morphological features can be perceived, providing helpful clues for determination of the proper diagnosis and guiding the selection of ancillary studies.

CLL is characterized by a monomorphic small/intermediate B-cell population composed of small lymphocytes with regular nuclei and coarse clumped chromatin. In some cases, and at a low amplification, clarified areas in the smear correspond to a variable percentage of prolymphocytes and paraimmunoblasts (fig. 3a). In general, patients with CLL present with a wide range of symptoms and multiple enlarged lymph nodes and the diagnosis is frequently made through peripheral blood analysis. Therefore, it is rather unusual that a first diagnosis of CLL be carried out in an FNB of an isolated lymph node. In most CLL patients, FNB is useful to exclude transformation in a high-grade lymphoma (Richter’s transformation).

MCL (classic) is also characterized by a homogeneous monotonous small cell population of cleaved centrocyte-like B cells. The blastoid and pleomorphic variant will be addressed in the differential diagnosis of large-sized B-cell lymphomas. Although most cases have a characteristic immunophenotype (CD5+, CD200–, CD10–, CD23–), it is advisable to perform immunostaining with cyclin D1 or SOX11 in cytopsins or cell blocks to confirm the diagnosis. FL, LPL, and MZL are composed of a more or less polymorphous lymphoid small/intermediate cell population. FL cytomorphology (grades 1–2) is in most situations indistinguishable from a reactive follicular hyperplasia. At a high magnification, attention should be paid to the scarceness of small naive lymphocytes and to the high percentage of centrocytes and centroblasts. FL also frequently displays lymphohistiocytic aggregates and macrophages with tingible bodies as seen in reactive lymphoid hyperplasia. Grading of FL can be performed in cytology, but this is not universally accepted [25, 26]. The distinction between grades 1 and 2 is poorly reproducible and has no clinical/therapeutic implications. Grade 3 FL are considered in the differential diagnosis with large-sized B-cell lymphomas.

LPL is characterized by a homogeneous polymorphic population with a mixture of small lymphocytes and a variable number of plasmacytoid lymphocytes and mature plasma cells. Mast cells are commonly identified in the subsidiary population. Differential diagnosis with MZL can be morphologically difficult, and ancillary testing as well as correlation with the clinical presentation are necessary. LPL patients have, in most cases, monoclonal IgM in the serum (Waldenström macroglobulinemia). MZL can present as a monotonous intermediate-sized cell smear, but more frequently it consists of a polymorphic cell population with small lymphocytes, centrocyte-like cells, centroblasts, plasmacytoid lymphocytes, plasma cells, and monocytoid cells. This type of lymphoma is rare and represents a challenge for cytological diagnosis; it has no characteristic immunophenotype and its cyto-
morphological characteristics overlap with those of other small-cell lymphomas and even reactive processes. Unlike extranodal MZL, nodal MZL lacks specific molecular markers like MALT1 and it is frequently a diagnosis of exclusion [27].

Lymphoblastic lymphomas and BL are sometimes considered medium-sized lymphomas, but in general they pose differential diagnostic problems mainly with large-sized lymphomas and will be addressed later [28].
Large-Sized Lymphomas

Large-sized B-cell lymphomas, whose paradigm is DLBCL, include other differential diagnoses such as reactive lymphoid hyperplasia, FL (grade 3), BL, blastoid MCL, HL, lymphoblastic lymphoma, anaplastic T-cell lymphoma, and grey-zone lymphomas. DLBCL account for the great majority of non-Hodgkin lymphomas and they encompass different variants, some of which should be highlighted. Although it is not crucial to individualize subtypes like centroblastic versus immunoblastic DLBCL, other subtypes like primary mediastinal B-cell lymphoma and T-cell/histiocyte-rich large B-cell lymphoma should be identified and represent a major challenge for FNB diagnosis.

Morphologically, DLBCL displays variable cytomorphological characteristics, reflecting the variability of its biology and pathogenicity. Smears are in general highly cellular and they display a more or less heterogeneous and polymorphic large cell population on a necrotic background with apoptotic bodies and macrophages with tingible bodies (fig. 7a). Nuclear features are variable, including round or irregular nuclei and a variable number of nucleoli.
To better understand this group of lymphomas, a huge effort has been made to correlate the cell of origin to distinct molecular subgroups with different immunocytochemical features, outcomes, and responses to target therapy. Based on the cell of origin, at least 3 subgroups of DLBCL were identified, i.e. germinal center B-cell-like (GCB), activated B-cell-like (ABC), and unclassifiable DLBCL [29]. Patients with ABC and unclassifiable DLBCL seem to respond less effectively to therapeutic regimens and have worse survival [30]. Immunohistochemi-
cal algorithms, which can be performed in surgically resected specimens, FNB cell blocks, or cytospins, have been developed to identify these subgroups [31, 32]. The most frequent algorithm for subtyping of the cell of origin subgroups includes antibodies like CD10, BCL6, and MUM1 and it shows about 80% concordance with the gene expression profile molecular subgroups (fig. 8) [32, 33]. More recently other algorithms were proposed with CD10, FOXP1, GCET-1, MUM1, and BCL6, gaining 92.6% concordance with the gene expression profile molecular subgroups [34], yet their use has not become widespread since they involve antibodies not used routinely in most laboratories. However, with restricted use FC can, through CD10 expression, guide in GCB or ABC DLBCL subtyping.

Most DLBCL NOS lymphomas are easily diagnosed in cytological preparations, but in some cases differential diagnostic problems arise with carcinomas, melanomas, and even seminomas. This is easily solved by using FC or immunostains in cytospins or cell blocks.

BL (fig. 9a) is characterized by a monotonous monomorphic large cell population. Characteristically, BL cells display typically coarse chromatin and cytoplasm featuring small lipidic vacuoles (fig. 9a; Giemsa).

The blastoid variant of MCL should also be included in the differential diagnosis of monotonous large B-cell lymphomas. Characteristically, BL cells display typically coarse chromatin and cytoplasm featuring small lipidic vacuoles (fig. 9a; Giemsa).

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Flow Cytometry

FC is a technology composed of a refined array of lasers, optics, fluidics, and electronic detectors capable of measuring light scatter and/or fluorescence emission from cells. This technique enables therefore the analysis and quantification of multiple cellular characteristics from a purified cellular suspension that is hydrodynamically focused on a single cell-wide laminar flow column.

Immunophenotyping by FC has become a standard routine in the diagnosis and monitoring of hematological malignancies. The principle of this technique is based on the conjugation of fluorochrome-labeled antibodies that bind to specific antigens.

Flow cytometers involve 3 major components: a fluidic, an optical, and an electronic system. The fluidic system carries particles (one by one) in a fluid stream across a laser beam for interception; the optical system includes lasers that provide a light source and optical filters to direct the light signals to each detector, and the electronics system converts light signals into electronic signals which in turn are converted to channel numbers by the analog-to-digital converter for further computer processing. This technique can provide information based on the physical properties of each individual cell, including relative size (based on forward-scattered light), relative granularity or
Fig. 9. a BL is characterized by a monotonous monomorphic large cell population. Characteristically, BL cells display a typical coarse chromatin (right; H&E, ×400), and cytoplasm featuring small lipidic vacuoles (left; Giemsa, ×600). b Immunophenotyping of BL. Abnormal B-cell population (red): CD10+, CD38+, LAIR1−, with light chain κ restriction. Color refers to the online version only.
internal complexity (based on side-scattered light – the higher the complexity the greater the amount of side-scattered light), and the expression of surface and cytoplasmic antigens. For the evaluation of intracellular markers, the cells must be fixed to ensure the stability of soluble antigens, and a cell permeabilization step prior to staining is required. The analyzed parameters are displayed in several graphic representations like histograms and dot plots. The term 'gating strategy' is used to define a restricted population that we want to study or characterize.

Immunophenotyping by FC applied to the study of lymphadenopathies and to the diagnosis of lymphoma has become routine practice, and it is nowadays recognized for its effectiveness. This technique allows objective observation of lymphocytes, with quantification and characterization, to distinguish abnormal from reactive populations.

The recent development of a great number of flow cytometers, software acquisition/analysis programs, different combinations of monoclonal antibodies, the use of a growing number of fluorochromes, the complexity of instrument settings, calibration, and novel protocols adapted to each work group have led to a complex process which is highly dependent on the expertise of the operator and needs to be accurate. It is essential to implement standardization in all steps of the pre- and postanalytic phases. Standardization involves numerous steps that include the use of monoclonal antibodies versus fluorochrome panels, the implementation of protocols for laboratory procedures, and definition of the appropriate software tools to recognize immunophenotypic patterns for multivariate analysis of the data.

Therefore, prompted by the need to optimize and standardize flow cytometric tests for diagnosis, prognosis, and evaluation of the treatment effectiveness of hematological malignancies, an important group, i.e. the European Union EuroFlow Consortium, emerged in 2005. At the time of reporting, EuroFlow was an independent Scientific Working Group of the European Hematology Association, and it encompassed a total of 20 diagnostic research groups and 11 countries in Europe and America [37]. With the creation of the EuroFlow Group, numerous protocols have been developed with the aim of standardizing the several proceedings involved in the entire pre- and postanalytical processes. The use of these protocols in laboratories has brought about a great evolution in the establishment of accurate diagnoses.

Protocols for the diagnosis and classification of hematological malignancies consist of a sequential combination of screening tubes that address a succession of specific questions.

Included in this analytic protocol are at least 2 types of cellular markers which were carefully chosen, i.e. backbone markers, which are used to identify distinct cell populations, and characterization markers, which are used to characterize target populations. Backbone markers expectantly identify both normal and malignant cells, and they are usually placed in all of the analytic panels in the same fluorochrome position, conditioning a constant and identical location of the target population and allowing automated gate setting of the target population. In contrast to backbone markers, which are present in the first screening tube as well as in the multitube panel, characterization markers are present only once in one tube of the panel, and they are chosen in accordance with the target population previously identified. In the EuroFlow-chosen protocols, the best possible combinations of backbone markers and characterization markers were designed. Each marker is selected based on its contributing information combined with all of the other markers of the panel.

In a first approach and as a critical prerequisite, an appropriate algorithm should be designed for sequential use of the different markers. For most lymphoma diagnoses, a rapid screening tube (lymphoid screening tube – backbone markers) is used to identify all relevant cell subcategories present in the sample (B, T, and NK cells) and to detect the presence of abnormal light chain-restricted populations (table 3). In a second step, several appropriate characterizing markers are selected to classify B, T, or NK cell chronic lymphoproliferative disorders according to WHO-defined disease categories (table 4). Labeling for the different antibodies is assessed using positivity and negativity but also MFI. CD45 is an important marker to discriminate, based on its MFI, mature from immature lymphocytes. CD19 and CD20, together with CD45, allow subtyping into mature B lymphocytes (CD19+, CD20+bright, and CD45+bright) and B-cell precursors (CD19+, CD20–/low intensity, and CD45+low intensity). Pan B-cell markers like CD19, CD20, and CD22, as well as surface membrane and cytoplasmic immunoglobulins, are used to demonstrate B-cell nature and light chain restriction. CD2, CD3, CD4, CD5, and CD7 are used to characterize T cells. NK cells should express antibodies of mature lymphocytes (CD45+bright) and CD56 in the absence of CD19; plasma cell backbone markers are CD45, CD19, and CD38. Additional markers are included to complement the subsets of B cells, T cells, and plasma cells (table 4).

MFI expression of some antibodies, is also used to characterize some specific subtypes of lymphomas. Un-
There is a consensus that the current diagnosis of lymphoma by FNB is usually based on the combined work of clinicians, morphology, immunophenotyping, and cytogenetics. In the majority of cases, this joint work allows achieving an accurate diagnosis and subcharacterization of some lymphomas, such as in CLL.

**Conclusions**

There is a consensus that the current diagnosis of lymphoma by FNB is usually based on the combined work of clinicians, morphology, immunophenotyping, and cytogenetics. In the majority of cases, this joint work allows achieving an accurate diagnosis and subcharacterization of some lymphomas, such as in CLL.
lapping with other entities. New immunocytochemical markers, such as SOX11, HGAL, LMO2, and LEF1, among others, have emerged and are contributing to a more accurate diagnosis [29]. Additionally, in selected cases, the use of molecular techniques may be helpful, since many lymphomas have genetic features that, in the appropriate clinical, morphological, and immunophenotypical context, can help in the diagnosis.

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