IMMUNOLOGICAL PHENOTYPING OF ACUTE LYMPHOBLASTIC LEUKEMIA (ALL) IN UNIVERSITY AFFILIATED HOSPITALS: A PRELIMINARY REPORT ON 50 PATIENTS

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ABSTRACT

The peripheral blood and bone marrow samples from 50 patients with ALL were investigated by indirect immunofluorescent technique. The most common type of ALL was common-ALL (82%) and the least common was T-ALL (2%). Other subtypes of ALL were unclassified by our technique (8%), B-ALL (4%), and pre B-ALL (4%). The correlation between immunophenotype, clinical condition and hematological profile is discussed.

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INTRODUCTION

After skin cancer, leukemias and lymphomas are the most common malignancies in the southern parts of Iran. Among the leukemia group, the incidence of the lymphocytic type is slightly higher than the granulocytic type and acute lymphoblastic leukemia (ALL) is the most common type of malignancies in children.

Recent advances in immunology have led to important insights into lymphocyte differentiation and the cellular origin of ALL. Now, it is well recognized that the identification of the lymphoblast's surface markers, or immunological phenotyping of ALL has provided important information in the diagnosis, prognosis and treatment of these patients.

The purpose of this study is to report the immunological subtypes in 50 patients obtained from hospitals in Shiraz. The samples were analysed by indirect immunofluorescent technique, using specific monoclonal antibodies that define cell surface antigens. The monoclonal antibodies used were anti HLA-DR, Anti-B lymphocyte antigen, anti-T lymphocyte antigen, antimyelo/monocytic antigen and anti-cALLA. These were combined with more traditional cell markers such as surface membrane immunoglobulins (SmIg), and sheep erythrocyte receptors on T lymphocytes, and also cytochemical stains such as PAS and Sudan black B.

RESULTS

The data from this study show that common-ALL is the most common phenotype of ALL at the present time in this area. About 82% of all cases and 88.6% of childhood-ALL are classified as common-ALL. 8% of cases were unclassified, 4% were B-ALL, 4% were pre B-ALL and 2% were of T-cell origin. The relative incidence of immunological subtypes of ALL are presented in Table I.

The majority of patients with C-ALL (64.6%) were...
Table I. The relative incidence of immunological subtypes of ALL.

<table>
<thead>
<tr>
<th>Phenotype</th>
<th>Subtype</th>
<th>No. patients</th>
<th>Percent</th>
</tr>
</thead>
<tbody>
<tr>
<td>CALLA+, Dr+, B.Ag+, SmI.g-, T.Ag-, ER-, myelo/monocytic Ag+</td>
<td>Common-ALL</td>
<td>41</td>
<td>82</td>
</tr>
<tr>
<td>CALLA+, Dr+, B.Ag+, SmI.g-, T.Ag-, ER-, myelo/monocytic Ag-</td>
<td>Pre B-ALL</td>
<td>2</td>
<td>4</td>
</tr>
<tr>
<td>CALLA-, Dr+, B.Ag+, SmI.g-, T.Ag-, ER+, myelo/monocytic Ag-</td>
<td>B-ALL</td>
<td>2</td>
<td>4</td>
</tr>
<tr>
<td>CALLA-, Dr-, B.Ag-, SmI.g-, T.Ag+, ER-, myelo/monocytic Ag-</td>
<td>T-ALL</td>
<td>1</td>
<td>2</td>
</tr>
<tr>
<td>CALLA-, Dr-, B.Ag-, SmI.g-, ER-, myelo/monocytic Ag-</td>
<td>Unclassified</td>
<td>4</td>
<td>8</td>
</tr>
<tr>
<td>Total</td>
<td></td>
<td>50</td>
<td>100</td>
</tr>
</tbody>
</table>

cALLA: Common acute lymphoblastic leukemia antigen; Ag: Antigen; SmI.g: surface membrane immunoglobulin; ER: E-Rosette;

reported to have the L2 morphologic form, whereas only 10.2% had the L1 type, and 2.5% the L3 type. One case of C-ALL was morphologically L3 type.

The majority of patients with C-ALL (88.6%) were aged 1-12 years. 66% of all cases and 65.8% of patients with C-ALL were male. With cytochemical staining all cases were Sudan black-B negative, and about 51% of all cases and 51.2% of C-ALL were PAS-positive. All cases of B-ALL were PAS negative.

As summarized in Table II, an analysis of clinical data recorded in 50 patients shows that fever and pallor are the most common symptoms; and splenomegaly (70.8%), hepatomegaly (66%) and lymphadenopathy (60.4%), are the most common clinical findings.

About 7.1% of all patients and 5.8% of C-ALL cases had mediastinal mass, and none of the patients whose charts were available had a testicular mass. A study of the patients’ charts showed that CNS involvement was found in 40% of all cases and in 40% of C-ALL cases. The range of WBC count was from 1100/mm³ to 370,000/mm³, and about 18.7% of cases had WBC counts above 50,000/mm³. WBC counts in all cases of the unclassified group were above 80,000/mm³.

DISCUSSION

The common classification of ALL is based on the FAB system, which defines three categories of ALL with regard to the morphology of the blast cells. This classification has a limited value for significant diagnos-
C-ALL, B-ALL and T-ALL, which we could recognize easily by monoclonal antibodies against cALLA, B-Ag, and T-Ag. However, we had some difficulties with other groups: 8% of cases were negative for all of the monoclonal antibodies which we used, so we called them an unclassified group. Also, in 4% of cases, cALLA and SmIg were negative but B antigen was positive. We could not classify them as B-ALL (because of absence of SmIg) and C-ALL (because of absence of cALLA), so we classified them as pre-B-ALL. Our study shows that common-ALL is the most common type of ALL, and T-ALL is the least common especially in children. Only one of 50 patients had T-ALL (2%).

In contrast with foreign reports, the majority of patients with C-ALL (84.6%) were reported to have the L2 morphologic form, whereas only 10.2% had the L1 type. Although all cases of ALL were Sudan black-B negative, about half of all the cases and 51.2% of the C-ALL group were PAS-positive. This matter shows that these stainings are not very significant for the diagnosis of ALL. In comparison with foreign reports with a high incidence of mediastinal mass in patients with ALL, only about 7.1% of all patients had mediastinal mass in this study. This may be because of the very low incidence of T-ALL in this area, as mediastinal mass is very common in T-ALL.

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REFERENCES

1. Iranian Cancer Society Reports, Fars Province Cancer Registry. 1978-1981.