Is DNA Index Measurement A Helpful Tool To Evaluate Iranian Children With Acute Lymphoblastic Leukemia (ALL)?

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Abstract

Aims and Background: The measurement of DNA content or ploidy has become a useful tool for diagnosis and treatment of malignancies particularly in patients with Acute Lymphoblastic Leukemia (ALL) as it has been associated with the most favorable prognosis in these patients.

The purpose of this study was to assess the incidence of DNA aneuploidy in Iranian children with ALL and the relationship between aneuploidy and morphological subclass, immunological phenotype, leukocyte count, age and sex in Tehran’s hospitals.

Methods: DNA content and laboratory features were determined in 70 ALL patients from Tehran’s hospitals, all of whom were under 15 years old and have not received any treatment. The morphological phenotype of the blasts was analyzed by microscopic examination. DNA content, the phenotype of the lymphoblast and WBC count were detected in the blood using flow cytometry.

Results: Diploidy, hyperdiploidy and hypodiploidy were reported in 50, 45 and 5% among the cases, respectively. The mean DNA index in hyperdiploid patients was 1.21 and hyperdiploidy was found to be associated with lower leukocyte count (p<0.001), lower age, L1 morphology and early pre-B immunophenotype.

Conclusions: This study showed that detected DNA index in Iranian children with ALL is associated with some favorable prognostic factor such as WBC count, age, immunological phenotype and FAB classification but that there is a need for further validation using extra molecular methods.

Keywords: DNA content; Immunophenotyping; FAB classification; Acute Lymphoblastic Leukemia

Introduction

Acute Lymphoblastic Leukaemia (ALL) is the most common malignancy of childhood which is associated with a clonal proliferation of lymphoblast cells in bone marrow and a sharp peak in incidence among children aged 2–5 years[1,2].

Various genetic alterations have been discovered in ALL, but hyperdiploidy has been known to be a favorable prognostic factor in childhood acute lymphoblastic leukemia. In contrast hypodiploidy has been introduced with a poor clinical outcome[3]. However high-hyperdiploidy and near haploid ALL are also reported to be associated with a unfavourable prognosis[4-9]. In addition it is reported that the existence of i(17q) exerts an adverse influence on treatment outcome in ALL, even in hyperdiploid cases[10].

Prognostic factors in children with ALL include age of child at diagnosis, race, gender, white blood cell counts, immunophenotyping, genetic abnormalities and response to early treatment. However there is clear evidence that DNA analysis of leukemic
blast cells contributes important prognostic information in these patients[11-14].

The purpose of this study was to assess the incidence of DNA aneuploidy using DNA index in Iranian children with ALL and its relationship with immunological phenotype, FAB classification, leukocyte count Age and sex.

Methods

Ethylenediaminetetraacetic acid (EDTA) anti coagulat-ed peripheral blood samples obtained from 70 patients with lymphoblastic leukemia and 70 normal, healthy cases from hospitals of Tehran. All of the patients and the controls were between 1-15 years of old and patients tested before taking any treatment to prevent drug interaction.

All specimens were obtained and prepared for morpho-logical examination using standard techniques. Peripheral blood smears were stained with Wright- Giemsa and examined under light microscopy and a scoring system for types L1 and L2 was carried out based on French-American-British Criteria (FAB).

Immunophenotyping

Mononuclear cells were isolated and stained with var-ious combinations of Fluorescein Isothiocyanate (FITC) and Phycoerythrin (PE - Labeled Monoclonal Antibodies, Sigma).

The panel of antibodies (all from Sigma) was used in distinct myeloid series (CD13, CD33 and CD14) from lymphoid and B cells from T cells (CD19, CD3 and CD7). CD10 (CAL-LA) antibody was also used to show the early pre-B phenotype. The last antibody was against Terminal Deoxynucleotidyl Trans-ferase (TdT) which is a useful marker in the diagnosis of ALL, L1 and L2 and mature B-lymphoid neoplasms or L3 phenotype.

DNA Analysis

To investigate DNA content all samples were treated with 1% Triton-100X to increase the permeability of the cell membrane. Treating the cells with ribonuclease (50 µl of a 100 µg/ml sock of RNase) was performed to remove RNA. Propidi-um Iodide (200 µl PI from 50 µg/ml stock solution) was used as a DNA-binding flurochrome and the samples were analyzed by flow cytometry. The DNA index was distinct as the modal DNA content of blasts compared to that of normal lymphocytes.

Results

Our data showed the percentage of female and male 37% and 63% respectively in hyperdiploid group, whereas in diploid group it was 30% and 70% respectively (Figure 1) The results of DNA analysis showed the diploidy, hyperdiploidy and hypodiploidy 50%, 45%, and 5% among the patients respectively. A mean DNA Index (DI) of 1.21 was detected among the hypodiploid patients (Figure 2). All the controls showed a single diploid pick in their DNA content histograms with a DI=1.
In hyperdiploid group L1 and L2 morphology were 63% and 37% whereas diploid group showed a percentage of 46.7 and 53.3 respectively (Figure 4). When the flow cytometric immunophenotyping data were correlated with the DNA contents, the cases with hyperdiploid showed a phenotype of early pre-B (40.7%), pre-B (48.1%), T (7.4%) and unclassified (3.7%), whereas diploid group showed an early pre-B (6.7%), pre-B (50%), T (40%) and unclassified (3.3%). Although just a few presents of our patients had been hypodiploid, they all demonstrated a pre-B immunophenotype (Figure 4 &5).

A positive correlation was found between L1 morphology and hyperdiploidy, as patients with a hyperdiploid peak in their histogram showed L1 morphology, indicating a possible favorable prognosis in this group. However, a better outcome for L1 and a higher relapse time for L2 have been established previously.[21-23]

Since cases with L3 morphology are uncommon, we could not find any cases with this morphology in our study. This type is uncommon and associated with a poorer prognosis.[24]

The phenotype is very imperative to determine the treatment protocol that the patient may receive. Acute lymphoblastic leukemias fall into one of the known immunophenotypes including early pre-B, pre-B, B-cell and T-cell. It is stablished that the prognosis for T- and B-cell ALL used to be poorer than for the pre-B ALLs. In the present study, early pre-B immunophenotype (the presence of CD10 or CALLA positive and the lack of cytoplasmic or surface immunoglobulins), which is associated with a favorable prognosis were found more in the hyperdiploid cases indicating a possible better outcome in these patients. This was similar to another study in the middle east.[25]. The number of patients with T cell phenotype was significantly higher in diploied group when compared with hyperdiploid indicating a poorer prognosis in the first group. Unexpectedly we found a number of patients with T phenotype, which are thought to be linked with a poor prognosis[25], but with a hyperdiploid phenotype. However The same result has been reported previously.[27]. This indicates that molecular cytogenetic studies must be associated with DNA index measurement to identify and characterize chromosomal rearrangements. Further investigations, therefore, are needed for our T-ALL patients, to be able to compare with the results of studies in other areas.[28-30].

A lower incidence of hyperdiploidy was found in elder patients compared with younger, which was statistically significant when compared with diploid and hyperdiploid group (p<0.05 and p<0.0001 respectively). This was in agreement with the previous reports[2,15,31], indicating the highest frequency of hyperdiploidy between ages of 2 and 7, which could be associated with a good outcome. In addition, we found a group of patients with T cell phenotype, who showed a positivity for CALLA antigen, suggesting further molecular studies such as cytogenetic in this group. However, it was previously reported that within T-ALL the expression of CALLA might be prognostically important.[32].

Conclusions

Taken together, this study showed that hyperdiploidy in Iranian children with ALL is associated with some favorable prognostic factor such as WBC count age, immunological phenotype and FAB classification, but there is a need for further validation via measuring additional prognostic markers.
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References