CONCISE REPORT

Pre-B Cell Leukemia Associated With Chromosome Translocation 1;19

By Andrew J. Carroll, William M. Crist, Richard T. Parmley, Maryann Roper, Max D. Cooper, and Wayne H. Finley

Chromosome banding studies on 60 children with acute lymphocytic leukemia (ALL), including "null," pre-B, B, and T cell phenotypes, were performed. In 4 of 17 patients with pre-B cell ALL, we noted a previously undescribed chromosome translocation, t(1;19)(q23;q13). This translocation was not found in patients with "null" cell, B cell, or T cell ALL. Since each patient with the 1;19 translocation experienced early treatment failure, t(1;19)(q23;q13) may mark a subgroup of patients with pre-B cell ALL who have an especially poor prognosis.

Immunologic classification of leukemia cells from each patient was performed as follows. Pre-B cells were identified by an immunofluorescent staining technique that permits the differential detection of surface and cytoplasmic immunoglobulin components on the same cell. If greater than 10% of marrow blasts contained μ heavy chains without detectable slg, a diagnosis of pre-B cell ALL was made. Spontaneous rosette formation with neuraminidase-treated sheep erythrocytes was used as a marker for T lymphoblasts.

Cytogenetic Studies

Bone marrow aspirates were cultured in RPMI 1640 supplemented with 10% fetal calf serum for 24 hr at 37°C, then exposed to colcemid (0.06 μg/ml) for 3.5 hr at 4°C. Routine methods were employed for culture harvest, slide preparation, and GTG-banding. In addition, direct chromosome preparations were examined.

RESULTS

Chromosomal banding studies were performed on the leukemic cells from 60 children with ALL of the following phenotypes: "null" cell, 34; pre-B, 17; B cell, 2; and T cell, 7. A unique chromosomal translocation, t(1;19)(q23;q13) was noted in leukemic cells of 4 patients with the pre-B cell ALL phenotype (Table 1) and was not seen in leukemic cells from the other 56 patients. Three distinct cytogenetic patterns were noted, each of which involved translocation of the distal one-half of the long arm (q23—qter) of chromosome 1 to the long arm of chromosome 19 in the q13 region. (This is a tentative designation because of the similar banded appearance of the short and long arms of chromosome 19.) Illustrative partial karyotypes from cases 1, 2, and 4 are shown in Fig. 1. Case 2

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Supported in part by Grants CA-13148 and CA-16673 awarded by the National Cancer Institute, DHEW, and by Bureau of Community Health Services Grant MCI-905.

Submitted August 22, 1983; accepted October 18, 1983.

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0006-4971/84/0303-0034$01.00/0

Table 1. Cytogenetic Findings for Four Patients With Pre-B Cell ALL and Chromosome 1:19 Translocation

<table>
<thead>
<tr>
<th>Patient</th>
<th>Clinical Status</th>
<th>Karyotype</th>
<th>No. of Cells Studied</th>
<th>Percent Abnormal</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>Diagnosis</td>
<td>46,XY,t(1;19)(q23;q13), -19, +der(19),t(1;19)(q23;q13)</td>
<td>22</td>
<td>100</td>
</tr>
<tr>
<td></td>
<td>2nd Remission</td>
<td>46,XY</td>
<td>15</td>
<td>0</td>
</tr>
<tr>
<td>2</td>
<td>Diagnosis</td>
<td>47,XX,t(1;19)(q23;q13), +8</td>
<td>21</td>
<td>100</td>
</tr>
<tr>
<td></td>
<td>1st Relapse</td>
<td>47,XX,t(1;19)(q23;q13), +8</td>
<td>18</td>
<td>33</td>
</tr>
<tr>
<td>3</td>
<td>1st Relapse</td>
<td>46,XX,-19,+der(19),t(1;19)(q23;q13)</td>
<td>30</td>
<td>100</td>
</tr>
<tr>
<td></td>
<td>2nd Remission</td>
<td>46,XX</td>
<td>8</td>
<td>0</td>
</tr>
<tr>
<td>4</td>
<td>1st Relapse</td>
<td>47,XY,-19,+der(19),t(1;19)(q23;q13), +6</td>
<td>26</td>
<td>61</td>
</tr>
<tr>
<td></td>
<td>2nd Remission</td>
<td>46,XY</td>
<td>15</td>
<td>0</td>
</tr>
</tbody>
</table>

displayed a form in which the translocation between chromosomes 1 and 19 appeared to be reciprocal, i.e., a normal chromosome 1 and 19 were present with a derived (der) no. 1 and a der(19). In cases 3 and 4, a normal chromosome 19 and a der(19) were present; unlike case 2, however, both no. 1 chromosomes were normal. In case 1, a normal no. 1 and a der(1) were noted along with 2 der(19) chromosomes.

Among the 13 pre-B cell patients without the t(1;19), 6 were cytogenetically normal, 5 were hyperdiploid, and 2 were pseudodiploid. Structural abnormalities of a diverse nature were identified, including t(9;22)(q34;q11), t(8;14)(q11;q32), inv(11)(p15;q23), and t(1;4)(q21;q35). No rearrangements involving chromosome 19 were noted within this group.

Clinical and immunologic features of the 4 patients with and the 13 pre-B cell leukemias without the 1;19 translocation are compared in Table 2. The patients with pre-B leukemias and a 1;19 translocation had noticeably lower white blood cell counts and, more importantly, experienced leukemic relapses much sooner than in pre-B leukemias not associated with the 1;19 translocation. The limited number of patients within each category did not permit meaningful statistical analyses of these trends.

**DISCUSSION**

We have prospectively performed immunologic typing and banded chromosome analysis on the leukemic cells of 60 children with ALL and have shown that 4 of 17 children (24%) with pre-B cell ALL had a unique chromosome anomaly, t(1;19)(q23;q13). This translocation was not found in any of the 43 other cases of "null" cell, B cell, or T cell ALL, nor have we observed this translocation in any of 50 cases of chronic myelogenous leukemia or 75 cases of acute nonlymphocytic leukemia among adults or children examined during the same period of time.

Previous cytogenetic studies of patients with pre-B cell ALL have consisted largely of isolated case reports, and a number of chromosomal anomalies have been described in their leukemia cells. Two cases of pre-B cell ALL with a 14q+ chromosome anomaly, similar to that associated with Burkitt's lymphoma, have been reported. A poor outcome was noted in each of these cases. However, in a recent cytogenetic survey of 49 patients with pre-B cell ALL, none had the 14q+ anomaly, suggesting that this must be a relatively rare occurrence in patients with pre-B cell ALL; other cytogenetic abnormalities were not detailed. Several individuals with Philadelphia chromosome (Ph+) positive chronic myelogenous leukemia have subsequently undergone a pre-B cell lymphoblastic crisis.

Abnormalities of chromosome 1, particularly trisomy for all or part of the long arm, as in patients 1, 3, and 4, and trisomy for chromosome 8, as in patient 2, are relatively common in hematologic disorders. Chromosome 19 anomalies are far less common. In some patients with non-Hodgkin's lymphomas, 19p+ or 19q+ markers have been reported; however, we are unaware of any previous report describing a t(1;19)(q23;q13) associated with chronic or acute leu-
kemia. The report\(^1\) of a translocation 7;19 in a child whose leukemia cells were transitional between pre-B and mature B cells is of special interest in view of the \(t(1;19)\) seen in our 4 patients. These findings may suggest a role for chromosome 19 rearrangements in some malignant disorders of B cell precursors.

Patients with Burkitt’s lymphoma-leukemia, a clinically aggressive malignancy of B cell phenotype, have been shown to have a translocation involving chromosomes 8 and 14 that results in the apposition of the cellular myc oncogene (on chromosome 8) to that coding for \(\mu\) chain synthesis (on chromosome 14).\(^1\)\(^-\)\(^3\) Promoter or enhancer sequences near the latter gene lead to the increased expression of the myc oncogene product\(^4\) and possibly to the malignant cell transformation. It will be of interest to determine if cellular oncogenes are located near the chromosome breakpoints (1q23) and 19q13) noted in our patients’ leu-

### REFERENCES


### Table 2. Clinical and Laboratory Features of Patients With Pre-B Cell ALL With or Without a 1;19 Translocation

<table>
<thead>
<tr>
<th>Patient Group</th>
<th>Age (yr) Mean±SE</th>
<th>Sex* (M/F) 1/2</th>
<th>Race* (White/Black) 3/1</th>
<th>WBC (x 10(^3)/cu mm) 12±5</th>
<th>Hemoglobin (g/dl) 10±2</th>
<th>Platelets (x 10(^3)/mm) 82±29</th>
<th>Percent Cells Positive for slg 2±1</th>
<th>slg 1±0</th>
<th>CALLA 81±5</th>
<th>Duration of First Remission (wk) 74±8</th>
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</thead>
<tbody>
<tr>
<td>4 Patients with t(1;19) (Mean±SE)</td>
<td>11±3</td>
<td>2/2</td>
<td>3/1</td>
<td>12±5</td>
<td>10±2</td>
<td>82±29</td>
<td>2±1</td>
<td>81±5</td>
<td>74±8</td>
<td>34</td>
</tr>
<tr>
<td>13 Patients without t(1;19) (Mean±SE)</td>
<td>6±1</td>
<td>6/7</td>
<td>10/3</td>
<td>88±46</td>
<td>9±1</td>
<td>115±32</td>
<td>3±1</td>
<td>64±8</td>
<td>82±5</td>
<td>66</td>
</tr>
</tbody>
</table>

*Expressed as ratio.
†Projected median.
murine plasmacytoma cells. Proc Natl Acad Sci USA 79:7837, 1982


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