A Rare Case of Pediatric T Lymphoblastic Leukemia With t(11;17)(q23;q21) Involving Mixed-Lineage Leukemia Gene Rearrangement

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Dear Editor,

T lymphoblastic leukemia (T-ALL) comprises 10-15% of pediatric ALL cases and is known to be a clinically and genetically heterogeneous disease [1]. Recent studies have presented various cytogenetic abnormalities in T-ALL. The alterations often present as reciprocal translocations implicated in several transcriptional factor genes and lead to arrest during T-cell differentiation [1, 2]. Although cytogenetic aberrations are known to be critical in leukemogenesis of many hematologic malignancies, the relationships between these aberrations and the prognosis are not well understood in T-ALL except for t(11;19), a mixed-lineage leukemia gene (MLL)-ENL fusion related to favorable prognosis [3]. Here, we report a very rare case of pediatric T-ALL with t(11;17)(q23;q21) involving MLL rearrangement, which has not been previously reported in Korea.

A 16-month-old boy was admitted to our hospital because of fever, cough, rhinorrhea, and poor oral intake. Physical examination revealed anemic conjunctiva, cervical lymph node enlargement, and splenomegaly. Chest radiography showed no abnormal findings. The complete blood count (CBC) results were as follows: hemoglobin, 8.2 g/dL; leukocytes, 41.3×10⁹/L; platelets, 177×10⁹/L; and reticulocyte count, 3.4%. The blood smear showed increased blasts (66%). The bone marrow (BM) specimen contained 65% blasts and showed a hypercellular pattern. Small-to-medium sized blasts were observed with condensed nuclear chromatin and an irregular nuclei margin. Immunophenotyping using the BM aspirate was brightly positive for CD2 (86%), cCD3 (91%), CD5 (91%), CD7 (92%), and CD34 (76%). Serum lactate dehydrogenase was increased to 638 IU/L. Cytogenetic studies on the BM revealed the karyotype of 46,XY,t(11;17)(q23;q21)[17]/46,XY[3] (Fig. 1A). The patient was diagnosed with T-ALL according to the 2008 WHO classification [4].

The patient began receiving induction chemotherapy comprised of prednisolone, vincristine, L-asparaginase, intrathecal cytosine arabinoside, and methotrexate. He seemed to respond to chemotherapy, with decreasing leukocyte counts and spleen size. However, the BM aspirate performed on the 7th day showed increased blasts to 69%. Moreover, interphase FISH analysis using an 11q23 break-apart probe revealed MLL rearrangement (158 of 200, 79%) (Fig. 1B). He was switched to more intensive induction chemotherapy including doxorubicin...
and cyclophosphamide. The patient achieved complete remission after 28 days of re-induction chemotherapy. Because of his high-risk features, the patient received cord blood stem cell transplantation. At the time of writing, he is alive and has been in a continuous complete remission state for 26 months.

**Table 1.** Reported cases of T-ALL with t(11;17)(q23;q21)

| Case No. | Age/sex | Leukocytes (×10^9/L) | Hb (g/dL) | Platelets (×10^9/L) | Blasts in PB (%) | Karyotypes | Immunophenotypes | OS (mo) | Country | Reference |
|----------|---------|----------------------|-----------|---------------------|-----------------|------------|------------------|---------|----------|-----------|
| 1        | 52 yr/F | 0.05                 | 8.4       | 0.08                | 68              | 46,XX,t(11;17) (q23-25;q21) | CD7(+), OKT10(+), CD1(-), CD3(-), CD4(-), CD8(-), CD10(-), CD19(-), anti-HLA-DR(-), 63D3(-), and TdT(-)* | > 14.5 | the Netherlands | 9 |
| 2        | 49 yr/M | ND                   | ND        | ND                  | ND              | 46,X,Y11;17 (q23;q21) | ND | 11.6† | EU‡ | 10 |
| 3        | 16 mo/M | 41.3                 | 8.2       | 177                 | 66              | 46,X,Y11;17 (q23;q21) / 46,X,Y1[3] | CD2 (86%), cCD3 (91%), CD5 (91%), CD7 (92%), and CD34 (76%) | > 26.4† | Korea | Present case |

*The results were performed with immunohistochemical stain; † The patients received bone marrow transplantation; ‡ The European 11q23 Workshop participants.

Abbreviations: T-ALL, T lymphoblastic leukemia; PB, peripheral blood; OS, overall survival; F, female; CD, cluster of differentiation; TdT, terminal deoxynucleotidyl transferase; M, male; ND, no data; EU, European Union; mo, months.

**Fig. 1.** Bone marrow karyogram at diagnosis and interphase FISH analysis after induction chemotherapy. (A) G-banded karyogram showing t(11;17)(q23;q21). The arrows denote the rearranged chromosomes. (B) FISH analysis using 11q23 break-apart probe revealed MLL rearrangement (158/200, 79%). Separation of a green and a red signal was observed.
ACACA, and LASP1 at 17q21 [8]. Thus far, MLL with each of the other three fusion partner genes at 17q21 has been identified mainly in AML and rarely in ALL [8]. Unfortunately, the fusion partner gene in this case could not be confirmed by using LDI-PCR because of specimen problems. A literature search revealed only two cases regarding t(11;17)(q23;q21) in adults with T-ALL [9, 10] (Table 1).

Prior studies have suggested that immunophenotypic markers are correlated with genetic alterations in pediatric acute leukemia [2]. The immunophenotype of leukemic cells in our patient could be classified into the pre-T stage and immature stage according to the European Group for the Immunological Characterization of Leukaemias and T-cell receptor classification system [2]. The immature stage of T-ALL was reported with respect to poor outcome and the lack of a common genetic lesion [1, 2]. In several studies, MLL-rearranged T-ALL overexpressed multiple HOX, which are transcription factors that initiate T-cell differentiation [5].

Additional cases involving genetic profiling of t(11;17)(q23;q21) involving MLL rearrangement will help assess whether this rare fusion impacts the disease progression and reveals prognostic information.

Authors’ Disclosures of Potential Conflicts of Interest

No potential conflicts of interest relevant to this article were reported.

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