A Novel Translocation: t(2;14)(p12;q32) in a Case of Precursor B-acute Lymphoblastic Leukemia

Sir,
Precursor B-cell acute lymphoblastic leukemia (B-ALL) is a disease of cytogenetic heterogeneity characterized by aneuploidy and/or structural alterations in pediatric ALL.[1] Rearrangement of IgH gene is seen nearly in all cases of B-ALL, namely, t(8;14)(q24;q32) and its variants t(2;8)(p12;q24) and t(8;22)(q24;q11) with less frequency in precursor B-ALL.[2] Recent developments include identification of several novel IgH gene translocations and submicroscopic deletions,[3] the significance of which requires further study. We report here a novel translocation t(2;14)(p12;q32) in a case of pre-B-ALL with IgH gene rearrangement.

An 8-year-old male child presented with fever and body ache, persisting for a month; there was no history of bleeding manifestation or any other symptoms. On clinical examination, the child had cervical lymphadenopathy and mild hepatosplenomegaly. Hemogram revealed hemoglobin level of 11.9 g, a total count of $51.4 \times 10^9/L$, and a platelet count of $18 \times 10^9/L$. The peripheral blood film showed 85% blasts which were medium to large with scant cytoplasm, irregular nuclei, coarse chromatin, and 1–2 nucleoli. A bone marrow study revealed 80% blasts with similar morphology as in peripheral blood film [Figure 1]. Other marrow elements were suppressed. Bone marrow study suggested ALL of L2 morphology.

In flow cytometry study, CD45 gating was done and the CD45 dim positive blasts were found to be positive for terminal deoxynucleotidyl transferase, CD34, CD19, cCD79a, and CD10, with aberrant CD15 expression. Other myeloid and monocytic markers were negative. With bone marrow and flow cytometry studies, a diagnosis of precursor B-ALL was made, in accordance with the WHO 2008 classification.

Cytogenetic analysis was done on G-banded metaphases from unstimulated bone marrow short-term cultures. The karyotype was made in accordance with ISCN 2013.[4] Two abnormal clones were identified, one with the karyotype 47, XY, t(2;14)(p12;q32) +del (12)(p13) and another with 46, XY, t(2;14)(p12;q32) [Figure 2]. Here, the breakpoints noted were at 2p12 and 14q32, which are classically involved in Burkitt lymphoma ALL-L3. Hence, cytogenetics is in favor of ALL-L3. Due to disparity between hematological investigation and cytogenetics, fluorescence in situ hybridization (FISH) analysis was done using locus specific identifier IgH dual color break-apart rearrangement probe. 74.5% of cells showed the presence of rearrangement of IgH gene at 14q32 [Figure 3]. This finding confirms the involvement of 14q32 in t(2;14)(p12;q32) detected by routine cytogenetics. By considering all the findings, precursor B-ALL with IgH gene rearrangement was the final diagnosis made. The patient has been started on MCP-842 protocol including methotrexate, cyclophosphamide, and prednisolone and is currently in induction phase. There was no evidence of central nervous system involvement at the time of diagnosis.

Discussion
Precursor B-ALL is a neoplasm of precursor cells (lymphoblasts) committed to B-cell lineage. The 2008 WHO classification recognizes specific entities with unique genetic lesions and prognostic features. In addition,
Letters to Editor

Prasannakumari Sampathkumar, Shanthi Velusamy1, Namrata Rajkumar2, M Padma3

Department of Pathology, Cytogenetics Unit, Kidwai Memorial Institute of Oncology, 1Department of Pathology and 2Paediatric Oncology, Kidwai Memorial Institute of Oncology, 3Department of Pathology, Hematology Unit, Kidwai Memorial Institute of Oncology, Bengaluru, Karnataka, India

Address for correspondence:
Dr. Shanthi Velusamy, Department of Pathology, Kidwai Memorial Institute of Oncology, Bengaluru, Karnataka, India.
E-mail: shanz84@gmail.com

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Conflicts of interest
There are no conflicts of interest.

Figure 3: Interphase fluorescence in situ hybridization using dual color break apart rearrangement probes showing rearrangement of IgH gene

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