

## Case Report Section

Paper co-edited with the European LeukemiaNet

# A new case of t(8;14)(q11;q32) in an acute lymphoblastic leukemia

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Published in Atlas Database: May 2003

Online updated version: <http://AtlasGeneticsOncology.org/Reports/814QuilichiniID100007.html>

DOI: 10.4267/2042/38002

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  2003 Atlas of Genetics and Cytogenetics in Oncology and Haematology

### Clinics

#### Age and sex

13 years old male patient.

#### Previous history

No preleukemia ; No previous malignancy ; No inborn condition of note

#### Organomegaly

No hepatomegaly, no splenomegaly, no enlarged lymph nodes, no central nervous system involvement

### Blood

**WBC:** 21X 10<sup>9</sup>/l

**HB:** 9.1g/dl

**Platelets:** 32X 10<sup>9</sup>/l

Blasts : 79%

Bone marrow: 97%

### Cyto-Pathology Classification

#### Cytology

Neutrophils: 11 %; eosinophils: 0 %; basophils : 0%; lymphocytes : 9 %; monocytes : 1%

#### Immunophenotype

HLA DR+ (high), CD34+ (high), CD10+ (low), CD19+ (high), c mu -, CD79a+ (high). Blasts were myeloperoxidase negative.

#### Diagnosis

Diagnosis of precursor B-cell acute lymphoblastic leukemia (WHO classification), LAL-B II (EGIL classification).

### Survival

**Date of diagnosis:** 04-2003, complete remission.

### Karyotype

**Sample:** Blood and bone marrow.

**Culture time:** Overnight unstimulated culture.

**Banding:** R-

**Results:**

45, X, - Y, t(8;14)(q11;q32) [ 16 ] / 46, XY [ 4 ]

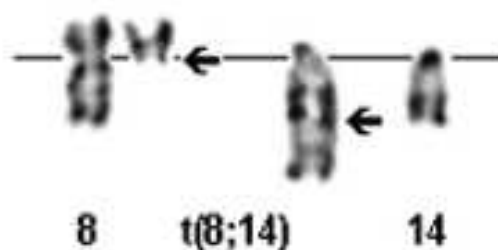


Figure 2: Karyotype (R-bands) : 45, X, -Y, t(8;14)(q11;q32).

#### Other molecular cytogenetics technics:

1- Fluorescence in situ hybridization (FISH) was performed using a probe specific for the IgH locus (14q32) and a chromosome 8 specific labelled Spectrum Orange painting probe (Abott) according to the manufacturer's instructions. FISH confirmed translocation t (8;14)(q11;q32) and reveals an IgH rearrangement (Figure 3).

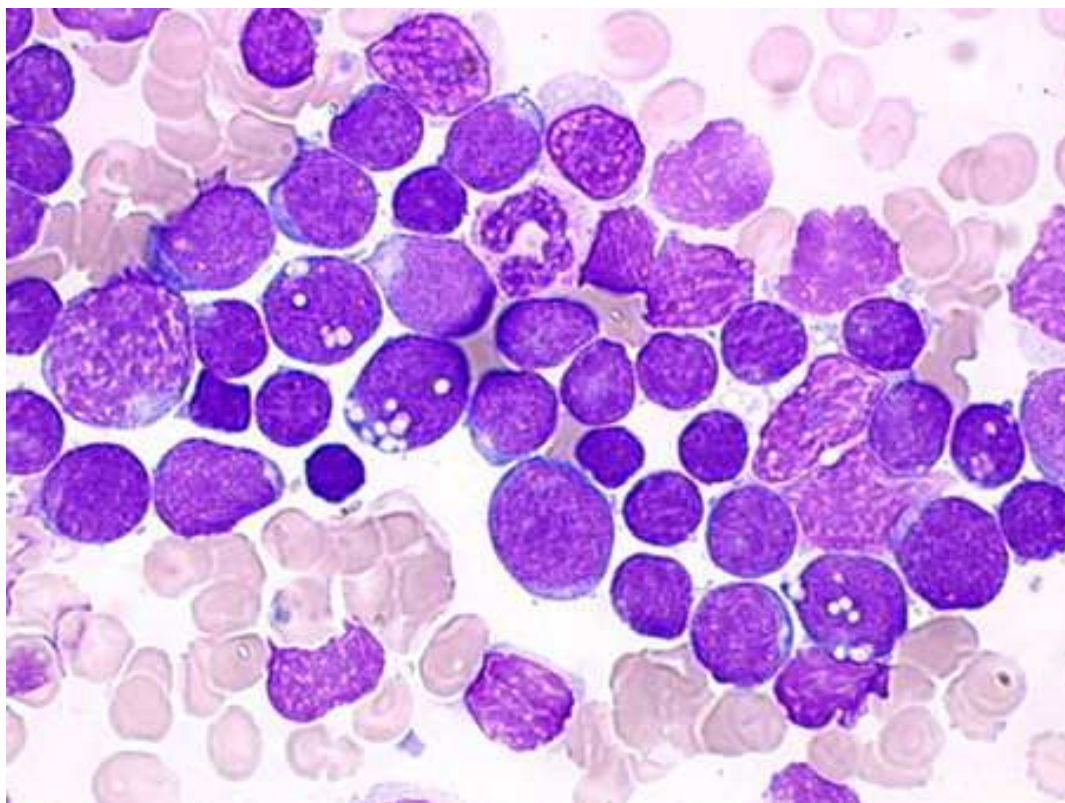


Figure 1: Bone marrow (MGG-stained) x 50: Increased cellularity, no megakaryocyte. Blasts : 97 % ; Neutrophils : 3%.

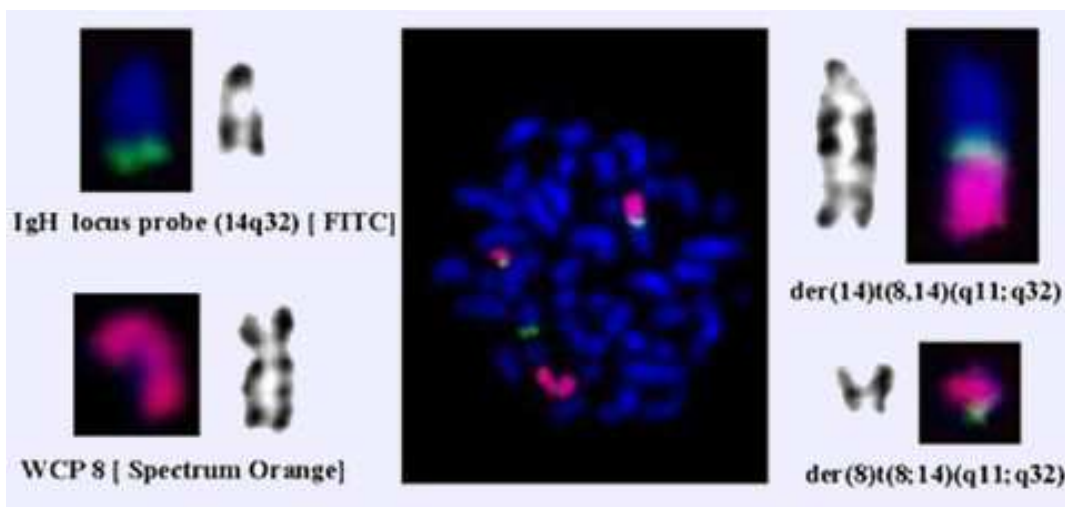


Figure 3: Fluorescence in situ hybridization using WCP 8 probe (Spectrum Orange) and LSI-IgH locus probe (FITC): partial karyotype results (R-banding and FISH).

2- FISH was performed using a chimeric X/Y direct labelled cocktail probe (Qbiogene) according to the manufacturer's instructions. FISH confirmed the loss of chromosome Y in 90% on interphase cells (data not shown).

### Other Molecular Studies

**Results:**  
 BCR/ABL: negative  
 TEL /AML1: negative

MLL gene : no rearrangement

### Comments

To our knowledge, the number of t(8;14)(q11;q32) cases reported in the literature is 44. In 10 cases, it was the sole acquired abnormality. In 34 cases, the t(8;14)(q11;q32) is associated with others numerical or structural abnormalities. 12 patients showed a constitutional trisomy 21. This case, an eleven year old boy represents an additional case. He does not present

Down syndrome (this association is further strengthened). Using EGIL classification, bone marrow immunophenotype classified this case as a common ALL (negative c mu and CD10 positive expression though a low level). It is the first report showing association of a t(8;14)(q11;q32) and loss of chromosome Y.

Splitting of the IgH gene was demonstrated in our case by FISH technique. As in translocations described in leukemias and lymphomas implicating the locus 14q32, the rearrangement could be crucial in this leukemic event.

The breakpoint at 8q11 has not yet been cloned.

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*This article should be referenced as such:*

Quilichini B, Zattara H, Casalonga F, Bastide-Alliez LA, Curtillet C, Fossat C, Michel G. A new case of t(8;14)(q11;q32) in an acute lymphoblastic leukemia. *Atlas Genet Cytogenet Oncol Haematol*. 2003; 7(3):212-214.

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