

Leukaemia Section

Short Communication

t(12;12)(p13;q13)

Jean-Loup Huret

Genetics, Dept Medical Information, UMR 8125 CNRS, University of Poitiers, CHU Poitiers Hospital, F-86021 Poitiers, France (JLH)

Published in Atlas Database: May 2005

Online updated version: <http://AtlasGeneticsOncology.org/Anomalies/t1212p13q13ID1391.html>

DOI: 10.4267/2042/38226

This work is licensed under a Creative Commons Attribution-Noncommercial-No Derivative Works 2.0 France Licence.
© 2005 Atlas of Genetics and Cytogenetics in Oncology and Haematology

Clinics and pathology

Disease

Myelodysplastic syndrome (MDS)

Epidemiology

Only 1 case to date: a 72 yr old female patient with RAEB1 which progressed to acute non lymphocytic leukemia (ANLL).

Cytogenetics

Sole anomaly in this patient.

Prognosis

The patient died 3 mths after diagnosis.

Genes involved and proteins

HGMA2

Location

12q15

Protein

Probable role in regulation of cell proliferation (transcriptional regulation of cell cycle and DNA repair genes).

Result of the chromosomal anomaly

Fusion protein

Description

Ectopic expression of HMGA2.

References

Odero MD, Grand FH, Iqbal S, Ross F, Roman JP, Vizmanos JL, Andrieux J, Lai JL, Calasanz MJ, Cross NC. Disruption and aberrant expression of HMGA2 as a consequence of diverse chromosomal translocations in myeloid malignancies. *Leukemia*. 2005 Feb;19(2):245-52

This article should be referenced as such:

Huret JL. t(12;12)(p13;q13). *Atlas Genet Cytogenet Oncol Haematol*. 2005; 9(3):247.