

Leukaemia Section

Short Communication

t(11;14)(q13;q32) in multiple myeloma

Jean-Loup Huret, Jean-Luc Laiï

Genetics, Dept Medical Information, University of Poitiers, CHU Poitiers Hospital, F-86021 Poitiers, France (JLH); INSERM Unité 524, Institut de Recherche sur le Cancer de Lille, Lille, France (JLL)

Published in Atlas Database: January 1998

Online version is available at: <http://AtlasGeneticsOncology.org/Anomalies/t1114MM.html>

DOI: 10.4267/2042/32110

This work is licensed under a Creative Commons Attribution-Non commercial-No Derivative Works 2.0 France Licence.

© 1998 Atlas of Genetics and Cytogenetics in Oncology and Haematology

Clinics and pathology

Disease

Multiple myeloma (MM) is a malignant plasma cell proliferation.

Phenotype / cell stem origin

Phenotype of mature differentiated B-cell, but also with CD56 expression, which is not found in normal plasma cells.

Epidemiology

Multiple myeloma's annual incidence: 30/10⁶; mean age: 62 yrs; t(11;14) is found in 10-20% of cases of MM with an abnormal karyotype; t(11;14) is not found associated with particular sex or age group; found mostly in stage III MM.

Clinics

Bone pain; susceptibility to infections; renal failure; neurologic dysfunctions.

Pathology

MM staging:

- Stage I: low tumour cell mass; normal Hb; low serum calcium; no bone lesion; low monoclonal Ig rate;
- Stage II: fitting neither stage I nor stage II;
- Stage III: high tumour cell mass; low Hb and/or high serum calcium and/or advanced lytic bone lesions and/or high monoclonal Ig rate.

Prognosis

Evolution: multiple myeloma can evolve towards plasma cell leukaemia;

Prognosis (highly variable) is according to the staging and other parameters, of which are now the karyotypic findings.

Cytogenetics

Cytogenetics, morphological

t(11;14) is balanced in most cases; some cases are: -14, +der(14)t(11;14); t(11;14) may well be a secondary

event in MM, as it has been found occurring during course of the disease.

Cytogenetics, molecular

FISH is indicated, as metaphases are arduous to obtain in such a disease implicating mature cells.

Additional anomalies

t(11;14) is part of a complex karyotype; accompanied with -13 or del(13q) in 'only' 1/4 of cases while -13/del(13q) is found in about 40% of MM cases with an abnormal karyotype; structural (and variable) anomalies of chromosome 1 are found in 1/3 of cases with t(11;14).

Variants

Complex three way translocations t(11;Var;14) have been described.

Genes involved and Proteins

BCL1

Location: 11q13

IgH

Location: 14q32

References

Feinman R, Sawyer J, Hardin J, Tricot G. Cytogenetics and molecular genetics in multiple myeloma. *Hematol Oncol Clin North Am* 1997 Feb;11(1):1-25. (Review).

Laiï JL, Michaux L, Dastugue N, Vasseur F, Daudignon A, Facon T, Bauters F, Zandecki M. Cytogenetics in multiple myeloma: a multicenter study of 24 patients with t(11;14)(q13;q32) or its variant. *Cancer Genet Cytogenet* 1998 Jul 15;104(2):133-8.

This article should be referenced as such:

Huret JL, Laiï JL. t(11;14)(q13;q32) in multiple myeloma. *Atlas Genet Cytogenet Oncol Haematol*.1998;2(1):34.