

# Leukaemia Section

## Mini Review

### **i(5)(p10) in acute myeloid leukemia**

**Claudia Schoch**

MLL Münchner Leukämielabor GmbH, Max-Lebsche-Platz 31, 81377 München, Germany (CS)

Published in Atlas Database: February 2005

Online updated version: <http://AtlasGeneticsOncology.org/Anomalies/i5p1D1376.html>

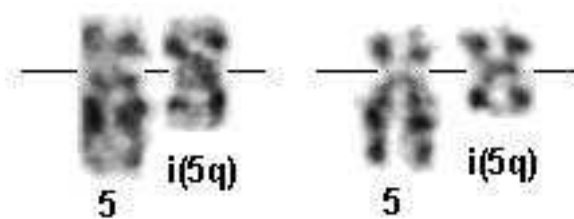
DOI: 10.4267/2042/38189

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#### Identity

##### Note

The isochromosome of the short arm of chromosome 5 - i(5)(p10) - has only been described in a few cases of myeloid leukemia. So far it has not been described as the sole abnormality. In four cases the i(5)(p10) was accompanied by trisomy 8, in three cases the i(5)(p10) occurred in addition to two normal chromosomes 5. An i(5)(p10) was also described in cases with a complex aberrant karyotype.



i(5)(p10) G-banding - Claudia Schoch.

#### Clinics and pathology

##### Phenotype/cell stem origin

Classified as AML, predominantly AML M5a.

##### Etiology

Unclear.

##### Epidemiology

Mean age 40-50 yrs.

##### Clinics

Blood data WBC  $8-40 \times 10^9/l$ , platelet counts  $15-114 \times 10^9/l$ .

##### Cytology

Typical cytomorphological features of AML M5a with

more than 80% of bone marrow cells being monoblasts showing strong cytochemical reaction with nonspecific esterase. Expression of CD33 and CD65.

##### Treatment

According to AML protocols.

##### Prognosis

Unclear due to low number of cases, seems to be poor.

#### Cytogenetics

##### Cytogenetics morphological

Isochromosome of the short arm of chromosome 5.

##### Additional anomalies

Trisomy 8, gain of chromosome 5.

#### Genes involved and proteins

##### Note

Gene dosage effect of genes located on the short arm of chromosome 5?

#### References

- El-Rifai W, Elonen E, Larramendy M, Ruutu T, Knuutila S. Chromosomal breakpoints and changes in DNA copy number in refractory acute myeloid leukemia. *Leukemia*. 1997 Jul;11(7):958-63
- Tamura S, Takemoto Y, Hashimoto-Tamaoki T, Mimura K, Sugahara Y, Senoh J, Furuyama JI, Kakishita E. Cytogenetic analysis of de novo acute myeloid leukemia with trilineage myelodysplasia in comparison with myelodysplastic syndrome evolving to acute myeloid leukemia. *Int J Oncol*. 1998 Jun;12(6):1259-62
- Markovic VD, Bouman D, Bayani J, Al-Maghrabi J, Kamel-Reid S, Squire JA. Lack of BCR/ABL reciprocal fusion in variant Philadelphia chromosome translocations: a use of double fusion signal FISH and spectral karyotyping. *Leukemia*. 2000 Jun;14(6):1157-60

Schoch C, Bursch S, Kern W, Schnittger S, Hiddemann W, Haferlach T. Gain of an isochromosome 5p: a new recurrent chromosome abnormality in acute monoblastic leukemia. *Cancer Genet Cytogenet.* 2001 May;127(1):85-8

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*This article should be referenced as such:*

Schoch C. *i(5)(p10) in acute myeloid leukemia.* *Atlas Genet Cytogenet Oncol Haematol.* 2005; 9(2):150-151.

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