

Case Report Section

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Translocation (X; 20)(q13;q13.3): a nonrandom abnormality in four patients with myeloid disorders: case 1

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Clinics

Age and sex

57 years old female patient.

Previous history

No preleukemia. Previous malignancy: Breast infiltrating ductal carcinoma treated with radiation and tamoxifen.

Organomegaly

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

Blood

WBC: $3.8 \times 10^9/l$

HB: 11.4g/dl

Platelets: $160 \times 10^9/l$

Cyto-Pathology Classification

Diagnosis

Myelodysplastic syndrome (MDS): refractory anaemia.

Survival

Treatment

Maintenance therapy for MDS patient declined bone marrow transplantation (BMT).

Relapse: no

Status: Alive

Survival: 42 months

Karyotype

Sample: BM

Culture time: 24/48h unstimulated cultures.

Banding: G-banding.

Results: $46,X,t(X;20)(q13;q13.3)[5]/46,XX[15]$.

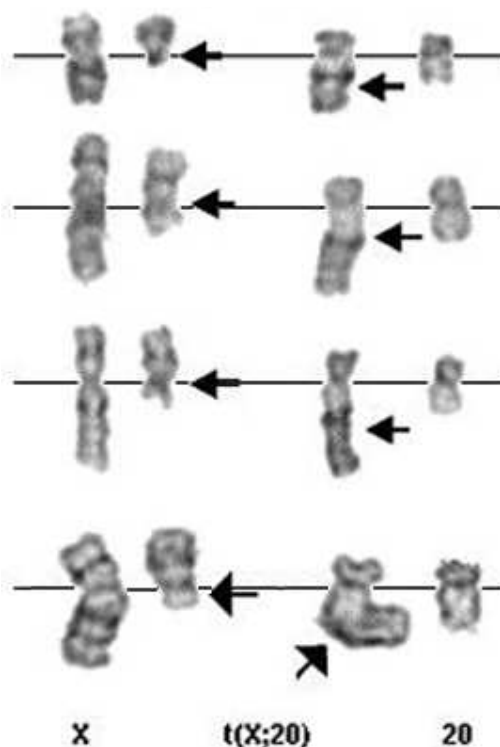


Fig. 1. Partial karyotypes of the translocation $t(X;20)(q13;q13.3)$ for cases 1-4 (top to bottom). Arrows indicate the derivatives 20 and X.

References

Gray BA, Cornfield D, Bent-Williams A, Zori RT. Translocation (X;20)(q13.1;q13.3) as a primary chromosomal finding in two patients with myelocytic disorders. *Cancer Genet Cytogenet.* 2003 Mar;141(2):169-74

Reddy KS, Richkind K, Ross M, Seirra R. Translocation (X;20)(q13;q13.3): a nonrandom abnormality in four patients with myeloid disorders. *Cancer Genet Cytogenet.* 2005 Feb;157(1):70-3

Michaux L, Wlodarska I, Mecucci C, Hernandez JM, Van Orshoven A, Michaux JL, Van den Berghe H. Characterization by chromosome painting of balanced and unbalanced X chromosome translocations in myelodysplastic syndromes. *Cancer Genet Cytogenet.* 1995 Jul 1;82(1):17-22

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