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Leukemia translocations

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Jonathan B Weitzman

Email: jonathanweitzman@hotmail.com

Chromosomal **translocations** involving the mixed-lineage leukemia gene (*MLL/ALL1*) define a subset of human acute leukemias with poor prognosis. In the Advanced Online Publication of **Nature Genetics**, Armstrong *et al.* describe a gene-expression profile analysis of leukemic cells with *MLL* translocations (DOI:10.1038/ng765). They compared the expression profiles of over 12,000 genes in B-precursor acute lymphoblastic leukemia (ALL) cells from **patients** with or without a rearranged *MLL* gene. The *MLL* samples had a distinct profile, with about 1,000 genes underexpressed and 200 highly expressed. The gene expression pattern was characteristic of maturational arrest at an early lymphoid progenitor stage; several *HOX* genes were highly expressed in *MLL* compared to other ALL samples. Armstrong *et al.* used principal component analysis (PCA) to cluster the *MLL*, ALL (acute lymphoblastic leukemia) and AML (acute myelogenous leukemia) expression profiles. They found that *MLL* profiles were distinct from ALL and AML samples, and defined a set of 100 genes that could distinguish *MLL* from the other leukemic classes. Such studies may lead to the development of translocation-specific therapies for *MLL* patients.

References

1. The critical role of chromosome translocations in human leukemias.
2. *Nature Genetics*, [<http://genetics.nature.com>]
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