Deletion of RUNX1 exons 1 and 2 associated with familial platelet disorder with propensity to acute myeloid leukemia

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Highlights

- Deletion of exons 1 and 2 of *RUNX1* associated with FPD/AML is described
- Thrombocytopenia is not mandatory in patients with *RUNX1* exon 1 and 2 deletion
- Thrombocytopenia should not be used as indication for screening in suspected family members with FPD/AML