# DISEASE:
# Acute myeloid leukemia with t(8;16)(p11;p13) translocation

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| **NAME:** | Acute myeloid leukemia with t(8;16)(p11;p13) translocation |
| **DESCRIPTION:** | A distinct form of Acute myeloid leukemia (AML) in which this chromosomal anomaly is found de novo or in therapy-related AML cases, and is characterized by frequent extramedullary involvement (mainly hepatomegaly, splenomegaly, lymphadenopathies, cutaneous infiltration, but also gum, bone, central nervous system, testicles involvement), severe coagulation disorder (disseminated intravascular coagulopathy or primary fibrinolysis) and poor prognosis. Morphologically, a blast population with a myelomonocytic stage of differentiation is observed. |
| **ORPHACODE:** | 370026 |
| **SYNONYMS:** | AML with t(8;16)(p11;p13) translocation |
| **XREF(S):** | Orphanet |
| **ANALYTE(S):** | CREBBP<br>KAT6A |
| **CREATED:** | 13 May 2019 - 01:02 |
| **CHANGED:** | 22 Jun 2023 - 16:14 |

## RELATED CONTENT

### Related Analytes

- CREB binding protein
- lysine acetyltransferase 6A