# Easy to understand reports facilitate treatment options.



# **TOPA Diagnostics**

351 East Rolling Oaks Drive, Suite 100 Thousand Oaks, CA 91361 805-373-8582 CLIA Number 05D1022855

# Directors

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Our reports use standardized formatting and terminology with color images that make them easy to comprehend and foster patient-physician communication.

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# PATH #: TB00-00000

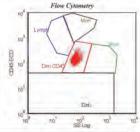
oblastic leukemia (AML-M5a) CD13 (partial), CD15, CD33, CD34, CD64; dim to nagative HLA-DR, CD117

tional information)

# 5:11)(q27:q23)[20] Abnormal Karyotype\*

ly seen in AMML and AMoL and is usually associated with a poor prognosis.





# hine) biopsy

two separate containers, each labeled Your, Patient. Specimen container #1 is cm in diameter tan white hard irregular fragment of bony tissue with minim

ed "BM clot" and consists of a 0.3 cm in diameter aggregate of a dark red sette 1B. Also received with the specimen are nine unstained slides. beled Your, Patient - peripheral blood. The specimen consists of

Note: A purple-topped tube containing blood is sent for flow cytometry processing to Clarient and a green-topped tube is sent to Clarient for Cytogenetics.

# SURGICAL PATHOLOGY REPORT

PATIENT: YOUR, PATIENT 10/10/1930 AGE: PHYSICIAN: Code Doctor, M.D. COPY TO:

TB00-00000 DATE COLLECTED: 01/12/2000 DATE RECEIVED: 01/12/2000 DATE REPORTED:

# LOCATION: DIAGNOSIS:

- BONE MARROW ASPIRATION AND CORE, RIGHT ILIAC CREST, (TREPHINE) BIOPSY: -Acute monoblastic leukemia, AML-M5a (see Microscopic Description and Comments)
  - -Decresed iron stores
  - -Abnormal karyotype (see Description)

# COMMENTS:

COMMENTS:
The bone marrow is replaced by sheets of blasts displaying high N/C ratio, immature chromatin, and prominent nucleoli. The peripheral blood smears show numerous blasts with morphologic features resembling monoblasts. The flow cytometry analysis demonstrates acute myeloid leukemia with immunophenotypic features most consistent with acute monoblastic leukemia (AML-M5a). Abnormal karyotype is noted, which is associated with a poor prognosis. Clinical correlation is reco

PERIPHERAL BLOOD:

-Acute monoblastic leukemia, AML-M5a -Normocytic anemia and thrombocytopenia

# MICROSCOPIC DESCRIPTION:

A recent CBC (1/1/2006) shows WBC 345,000; HGB 11.1; MCV 94; MCHC 34; RDW 16.5; PLAT 119,000.

## PERIPHERAL BLOOD SMEAR:

The peripheral blood smears show numerous blasts with high N/C ratio, immuture chramatin, and 1-3 prominent nucleoli. Some blasts show nuclear folds. Occasional blasts show cytoplasmic granules. No Auer rods are identified. The red blood cells are normocytic and normochromic with mild anisopoikilocytosis. Occasional nucleated red blood cells are noted. Moturing leukocytes are markedly decreased consisting of rare segmented neutrophils, monocytes and lymphocytes. No eosinophils or basophils are noted. The platelets are decreased in number. The manual platelet count is about 60,000 - 80,000. No platelet clumping is noted. Some cytoplasmic fragments are noted.

# BONE MARROW ASPIRATE:

The bone marrow aspirate smears show numerous blasts with the same morphologic features as described in the peripheral blood smear. Normal marrow elements are essentially absent.

# BONE MARROW BIOPSY AND CLOT SECTIONS:

DONG MARKOF BUFST AND CLOU SECTIONS:
The decalefied bore marrow bipsys section shows hypercellular marrow (85%). The marrow space is entirely replaced by sheets of blasts displaying high N/C ratio, immature chromatin and prominent nucleoli. Normal marrow elements are essentially absent. No basophils or cosinophils are identified. The bone trabectulae are normal. No fibrosis or granulomas are identified. The clot sections show a few hypercellular particles with findings similar to those in the biopsy section.

Iron stain shows decreased storage iron (trace amount of stainable iron). No ringed sideroblasts are identified. PAS stain shows no megakaryocytes. Reticulin stain shows no reticulin fibrosis.



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