

**Patient Name:** Sample Patient  
**D.O.B. & Gender:** 10/10/20XX (xx yrs.) - Male  
**Collection Date:** 06/26/2014  
**Received Date:** 06/27/2014  
**Report Date:** 06/27/2014  
**Specimen Type:** Bone Marrow

**Ordering Physician:** Smith, Jane  
**Ordering Facility:** John Doe Medical Group  
**Account Number:** 0000  
**VantagePoint ID #:** 200000  
**VantagePoint Case #:** FLG13-000000  
**Medical Record #:**

## Flow Cytometry Report

**Diagnosis: 1) POSITIVE FOR ACUTE MYELOID LEUKEMIA WITH 35% BLASTS DETECTED.  
 2) LEUKEMIA LACKS CD34 AND HLA-DR; FEATURES WORRISOME FOR ACUTE PROMYELOCYTIC LEUKEMIA  
 SUBTYPE OF AML (PLEASE SEE INTERPRETATION).**

Interpretation: The blasts percentage by flow cytometry is consistent with an acute myeloid leukemia. The lack of HLA-DR & CD34 is a phenotype which is seen in AML-M3 (a.k.a. acute promyelocytic leukemia), but can occasionally be seen post-therapy. Limited morphologic review of the smear is inconclusive in determining AML-M3 versus AML, not otherwise specified. Fluorescence in-situ hybridization for translocation (15;17) - (PML/RARA gene re-arrangement), can be performed if needed (or if no previous diagnosis present). This translocation would be diagnostic for acute promyelocytic leukemia if present. Please contact Vantage Point Laboratories, if testing needed. Please also correlate with core biopsy, if available.

The aberrant CD7 and CD56 expression have utility in future detection of residual disease.

**PHENOTYPE OF BLASTS:**  
**POSITIVE FOR:** CD4, CD11B, CD13, CD33, CD38, CD56, CD117, ABERRANT CD7  
**NEGATIVE FOR:** CD34, HLA-DR

CD45 Dim: 35% blasts of with myeloid phenotype mentioned above.

Lymphocytes are unremarkable.

Monocytes: mature monocytic phenotype. Only 4%.

Granulocytes: The mature granulocytes are decreased with variable phenotypic abnormalities.

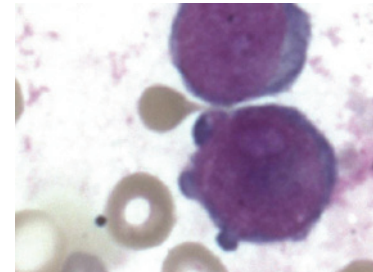
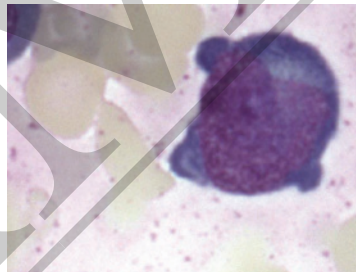
Clinical Data: xx year old male

**Specimen Yield:** Adequate **Viability:** Adequate

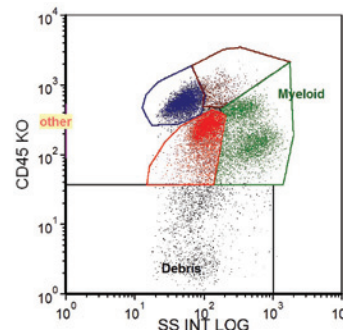
Submitted Dx: 205.00

Additional Clinical Data Submitted: RBC 2.8 mm<sup>3</sup>; WBC 5.1 mm<sup>3</sup>; HGB 10.1 g/dL; HCT 30.6%; PLT 22 mm<sup>3</sup>

Flow Cytometry Differential (% of Total Cells)	
<b>Lymphocytes</b>	30%
<b>B-cells</b>	10%
Kappa	19%
Lambda	12%
<b>Kappa:Lambda Ratio</b>	1.6
<b>T-cells</b>	19%
CD4	30%
CD8	31%
<b>CD4:CD8 Ratio</b>	1.0
<b>Monocytes</b>	4%
<b>Granulocytes</b>	24%
<b>CD45 Dim</b>	35.0%
<b>Viability</b>	95%
Coexpressions	
CD19+CD5+	3%
CD19+CD10+	2%
CD4+CD8+	0%



**Markers:** Antibodies against the following antigens were used in comprehensive ten-color multiparameter flow cytometric Screen panel analysis: CD2, CD3, CD4, CD5, CD7, CD8, CD10, CD11b, CD11c, CD13, CD14, CD16, CD19, CD20, CD23, CD25, CD33, CD34, CD38, CD45, CD56, CD64, CD103, CD117, HLA-DR, Kappa, Lambda, CD22, FMC-7. (29 Markers)



Electronically Signed by:

Anand Kunda M.D.  
 Hematopathologist

This test was performed at VantagePoint Laboratory Partners, Inc., 4980 Carroll Canyon Road, San Diego, CA 92121. This test has not been cleared or approved by the US Food and Drug Administration. The FDA has determined that such clearance is not necessary. The test is used for clinical purposes and should not be regarded as investigational or for research. This laboratory is certified under the Clinical Laboratory Improvement Amendments of 1988 (CLIA-88) as qualified to perform high complexity clinical testing. This document contains private and confidential health information protected by state and federal law. If you received this document in error, please call 888.VANTAGE.