

PROFICIENCY TESTING REPORT





NABL accredited program as per ISO/IEC 17043:2010 standard Organized By Department of Hematology, AIIMS, New Delhi-110029

Duration of stability testing - minimum upto 8 days at ambient temp. after dispatch of specimens

EQAP CODE No.: 4285 **Distribution No.:** 156-K **Month/Year:** July/2022

Instrument ID: Mindray BC -6200, TW-9A000724

Name & Contact No. of PT Co-ordinator: Dr. Seema Tyagi (Prof.), Hematology, AIIMS, Delhi,

Tel: 9013085730 , E-Mail : accuracy2000@gmail.com **Date of issue & status of the report:** 25-09-2022[Final].

CBC and Retic Assessment

				Among Lab (Accuracy Testing)				Within Lab (Precision Testing)				
Test Parameters	S.No.	Your Result 1		Your Results Sum of 2 Value	Consensus result sum of 2 values (Assigned Value)	Uncertainty		Results		Uncertainty of Assigned Values	Z Score	
WBC x10³/μl	1	4.51	4.31	8.82	9.2	0.0610	-0.24	0.2	0.1	0.0110	0.57	
RBC x10 ⁶ /μl	1	4.53	4.47	9	8.88	0.0110	0.45	0.06	0.05	0.0030	0.19	
Hb g/dl	1	14	13.7	27.7	27	0.0370	0.73	0.3	0.1	0.0100	1.35	
НСТ%	1	44.7	44.2	88.9	83.2	0.2210	1.06	0.5	0.5	0.0370	0.00	
MCV-fl	1	98.8	98.5	197.3	186.65	0.3960	1.18	0.3	0.3	0.0270	0.00	
MCH-Pg	1	30.8	30.6	61.4	60.9	0.0870	0.26	0.2	0.3	0.0190	-0.34	
MCHC-g/dl	1	31.3	30.9	62.2	65.15	0.1890	-0.67	0.4	0.3	0.0210	0.32	
Plt. x10³/μl	1	111	106	217	211.5	1.74	0.12	5	4	0.32	0.22	
Retic %	2	1.4	1.2	2.6	3.8	0.10	-0.48	0.2	0.2	0.01	0.00	

P.S. Assesment

		YOUR REPORT	CONSENSUS REPORT				
DLC%	3		Poly: 35 - 48, Myelo: 14 - 26, Meta: 8 - 15, Promyelo: 3-7, nRBC/ Lympho /Blast/Eos/Baso/Mono: 0 - 5				
RBC Morphology	3		Predominantly: Normocytic/Normochromic; Moderate: Anisocytosis, hypochromia, Microcytosis; Mild: Macrocytosis, Poikilocytosis				
Diagnosis		Chronic Myeloid Leukemia(CML) - Chronic phase	Chronic Myeloid Leukemia (Chronic Phase)				

COMBINED DATA VALUES OF TOTAL PARTICIPANTS

Test parameters	S.No.	Total participants	Total No. responded	% of Labs with Z Score 0-2		% of Labs with Z Score 2-3		% of Labs with Z Score >3		
rest parameters		current dist. 156K		Among labs	Within lab	Among labs	Within lab	Among labs	Within lab	
WBC x10³/μl	1	252	248	84.27	87.1	6.85	2.42	8.88	10.48	
RBC x10 ⁶ /μl	1	252	252	86.11	92.46	6.75	1.59	7.14	5.95	
Hb g/dl	1	252	252	90.87	82.54	1.98	7.94	7.15	9.52	
HCT%	1	252	248	94.35	91.94	2.42	2.82	3.23	5.24	
MCV-fl	1	252	248	89.11	94.35	6.85	2.42	4.04	3.23	
MCH-Pg	1	252	248	87.1	<mark>9</mark> 1.53	6.05	2.82	6.85	5.65	
MCHC-g/dl	1	252	248	91.13	91.94	4.44	2.42	4.43	5.64	
Plt. x10³/μl	1	252	248	92.34	89.11	3.63	5.24	4.03	5.65	
ReticCount%	2	252	214	93.46	89.72	4.67	7.48	1.87	2.80	
PS Assessment	3	252	213	Satisfactory:93.65%, Borderline Sat.:4.76%, Unsatisfactory:1.58%						

*Comments:

1). Among Lab (EQA): Results acceptable.

2). Within Lab (IQA): Precision acceptable.

Note-1: EQA (External Quality Assurance): Your Performance among various of participating labs in PT, to determine the accuracy of your results.

IQA (Internal Quality Assurance): Your Performance of comparison of two consecutive measurement values within your lab to test the precision of your autoanalyzer.

Note-2: Z score among & within lab were calculated, as per to ISO/IEC 13528:2015 standard. Z score among lab (EQA)= (Your Result Sum of two values - Consensus Result sum of two values)/(Normalised IQR)

Z score within lab (IQA)= (Your Result Difference of two values - Consensus Result difference of two values)/(Normalised IQR)

IQR = Quartile 3 - Quartile 1 of participant data, Normalised IQR = 0.7413 x IQR

Note-3: Z score 0 to ± 2 : Acceptable, Z score ± 2 to ± 3 : Warning Signal, Z score $> \pm 3$: Unacceptable [As per ISO/IEC 13528:2015 standard]

Note-4: Z score value between "0 to ± 2 " are texted in green colour. Z score value between " ± 2 to ± 3 " are texted in orange colour. Z score value $> \pm 3$ are texted in red colour.

Note-5: Homogeneity and stability testing of PT sample were done as per ISO 13528:2015 standard. To pass homogeneity test, between sample SD (Ss) should be smaller than the check value (0.3*SDPA). To pass the stability test, average difference in measurement values of first and last day sample $(\bar{x}-\bar{y})$ should be smaller than the check value (0.3*SDPA).

Note-6: ISHTM-AIIMS-EQAP does not subcontract any task of its scheme

Note-7: Participants are free to use methods/analyzer of their own choice.

Note-8: Proficiency testing (PT) samples are sent quarterly to each participant.

Note-9: All the necessary details regarding design and implementation of PT, are provided in the instruction sheet as well as on programme's website www.ishtmaiimseqap.com.

Note 10: Reports are kept confidential.

Report authorized by,

Dr. Seema Tyagi (Prof.)

PT Co-ordinator: ISHTM-AIIMS-EQAP

Department of Hematology, AIIMS, New Delhi

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