



# PROFICIENCY TESTING REPORT

# ISHTM-AIIMS EXTERNAL QUALITY ASSURANCE PROGRAMME

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.:** 5822 **Distribution No.:** 160-O **Month/Year:** July/2023

Instrument ID: Mindray BC 3600 (TB89002330)

Name & Contact No. of PT Co-ordinator: Dr. Manoranjan Mahapatra ( Prof. & Head), Hematology, AIIMS, Delhi,

Tel: 9013085730 , E-Mail : accuracy2000@gmail.com **Date of issue & status of the report:** 19-09-2023[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	3.7	3.7	7.4	6.86	0.043	0.62	0	0.08	0.006	-1.20	
RBC x10 <sup>6</sup> /μl	1	5.78	5.59	11.37	9.52	0.016	5.96	0.19	0.04	0.004	2.79	
Hb g/dl	1	11.6	11.6	23.2	25.3	0.036	-2.58	0	0.1	0.011	-0.67	
НСТ%	1	48.1	46.5	94.6	80.7	0.256	2.59	1.6	0.4	0.045	2.02	
MCV-fl	1	83.3	83.1	166.4	169.25	0.451	-0.29	0.2	0.3	0.030	-0.22	
МСН-Рд	1	20.8	20.1	40.9	53	0.104	-6.09	0.7	0.2	0.018	2.25	
MCHC-g/dl	1	25	24.1	49.1	63.1	0.231	-3.02	0.9	0.3	0.027	1.70	
Plt. x10³/μl	1	193	186	379	316.5	2.317	1.22	7	7	0.578	0.00	
Retic %	2	0.02	0.01	0.03	1.58	0.054	-1.07	0.01	0.2	0.014	-0.79	

## P.S. Assesment

		YOUR REPORT	CONSENSUS REPORT				
DLC%	3		Poly: 37 - 52, Myelo: 15 - 27, Meta: 9- 17, Promyelo: 2-8, Lympho: 2- 5, Blast: 1-4, Eosino: 1-3, Mono: 1-2, nRBC/ Baso: 0-5				
RBC Morphology	3	RBC's are normocytic, normochromic WBC series, slow leucocytosis with a number of immature cells. Platelets are reduced	Predominantly: Normocytic/Normochromic; Moderate: Anisocytosis, hypochromic, Mild: Poikilocytosis				
Diagnosis	3	Chronic Myelocytic Leukaemia	Chronic Myeloid Leukemia (Chronic Phase)				

### **COMBINED DATA VALUES OF TOTAL PARTICIPANTS**

Test never eters	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	
Test parameters		current dist. 1600		Among labs	Within lab	Among labs	Within lab	Among labs	Within lab
WBC x10³/μl	1	199	198	86.36	93.43	3.54	1.01	10.1	5.56
RBC x10 <sup>6</sup> /μl	1	199	199	87.44	86.93	5.53	5.53	7.03	7.54
Hb g/dl	1	199	199	86.93	82.91	6.53	5.53	6.54	11.56
HCT%	1	199	1 <mark>98</mark>	92.93	90.4	2.53	3.03	4.54	6.57
MCV-fl	1	199	198	92.42	87.88	6.57	4.04	1.01	8.08
MCH-Pg	1	199	198	83.33	<mark>9</mark> 2.93	9.09	1.01	7.58	6.06
MCHC-g/dl	1	199	198	90.91	85.86	5.56	5.05	3.53	9.09
Plt. x10³/μl	1	199	198	89.39	92.93	6.06	3.54	4.55	3.53
ReticCount%	2	199	151	85.43	92.72	7.28	9.93	7.29	-2.65
PS Assessment	3	199	163	Satisfactory:98.5%, Borderline Sat.:0%, Unsatisfactory:1.50%					

#### \*Comments:

1). Among Lab (EQA): CBC result for *RBC*, *MCH* & *MCHC* unacceptable, please check calibration/human error.Remaining 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  $\pm 2$ : Acceptable, Z score  $\pm 2$  to  $\pm 3$ : Warning Signal, Z score  $> \pm 3$ : Unacceptable [As per ISO/IEC 13528:2015 standard]

**Note-4:** Z score value between "0 to  $\pm 2$ " are texted in green colour. Z score value between " $\pm 2$  to  $\pm 3$ " 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 guarterly 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. Manoranjan Mahapatra (Prof. & Head)

PT Co-ordinator: ISHTM-AIIMS-EQAP

Department of Hematology, AIIMS, New Delhi

-----End Of Report-----