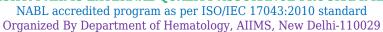




# PROFICIENCY TESTING REPORT

# ISHTM-AIIMS EXTERNAL QUALITY ASSURANCE PROGRAMME





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

**EQAP CODE No.:** 1433 **Distribution No.:** 159-C Month/Year: February/2023

Instrument ID: A380 CELLT SN 665000215

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: 05-04-2023[Final].

## **CBC** and Retic Assessment

				Among Lab (Accuracy Testing)				Within Lab (Precision Testing)				
Test Parameters	S.No.	Your Result 1		Results	Consensus result sum of 2 values (Assigned Value)	Uncertainty		Yours Results Diff. of 2 Values	Consensus Result Diff. of 2 values (Assigned Value)	Uncertainty of Assigned Values	Z Score	
WBC x10³/μl	1	2.23	2.1	4.33	4.23	0.017	0.24	0.13	0.08	0.004	0.75	
RBC x10 <sup>6</sup> /μl	1	3.09	3.04	6.13	6.06	0.006	0.43	0.05	0.03	0.002	0.67	
Hb g/dl	1	10.1	10	20.1	20.8	0.018	-1.57	0.1	0.1	0.007	0.00	
НСТ%	1	29.81	29.6 <mark>4</mark>	59.45	64.7	0.116	-1.34	0.17	0.3	0.020	-0.44	
MCV-fl	1	98	96	194	213.4	0.339	-1.64	2	0.3	0.024	3.82	
МСН-Рд	1	32.7	32.5	65.2	68.7	0.071	-1.69	0.2	0.3	0.018	-0.34	
MCHC-g/dl	1	33.7	33.6	67.3	63.9	0.115	0.81	0.1	0.3	0.018	-0.67	
Plt. x10³/μl	1	78	71	149	174	0.707	-1.20	7	3	0.194	1.35	
Retic %	2	0.45	0.4	0.85	8.9	0.145	-1.86	0.05	0.4	0.021	-1.57	

## P.S. Assesment

		YOUR REPORT	CONSENSUS REPORT				
DLC%	3		Poly: 43-54 , Lympho: 40-50 , Mono: 2-6, Eosino: 1-3 , nRBC:0-2, blast/Promyelo/Myelo/Meta: 0				
RBC Morphology	3	HYPOCHROMIC,FEW SICKLE SHAPED	Predominantly: Normocytic/Normochromic; Moderate: Anisocytosis, Microcytosis, Hypochromia; Mild: Poikilocytosis, Target cells, Sickle shaped cells, tear drop cells				
Diagnosis	3	SICKLE CELL DISEASE	Hemoglobinopathy Likely sickle cell-Beta Thalassemia				

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

Test never eters	S.No.	Total participants covered in the current dist. 159C	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				Among labs	Within lab	Among labs	Within lab	Among labs	Within lab
WBC x10³/μl	1	349	347	84.73	92.8	6.05	1.15	9.22	6.05
RBC x10 <sup>6</sup> /μl	1	349	349	91.4	87.39	5.73	5.73	2.87	6.88
Hb g/dl	1	349	349	90.54	92.84	5.44	3.44	4.02	3.72
HCT%	1	349	3 <mark>47</mark>	98.56	88.18	0.86	6.34	0.58	5.48
MCV-fl	1	349	347	99.42	92.22	0.58	3.75	0	4.03
MCH-Pg	1	349	347	93.95	<mark>8</mark> 7.03	4.03	5.48	2.02	7.49
MCHC-g/dl	1	349	347	97.69	87.03	2.02	6.92	0.29	6.05
Plt. x10³/μl	1	349	347	90.2	91.35	6.63	4.03	3.17	4.62
ReticCount%	2	349	331	94.56	82.48	4.23	13.29	1.21	4.23
PS Assessment	3	349	314	Satisfactory :88.83%, Borderline Sat. :8.02%, Unsatisfactory :3.15%					

#### \*Comments:

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

2). Within Lab (IQA): Difference in the CBC measurement values for MCV unacceptable, may be due to random/human error.

**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 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

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