

### **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. : 289

Distribution No.: 158-A

Month/Year: October/2022

Instrument ID: BC 6200 (TW-04001042)

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Date of issue & status of the report: 21-12-2022[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 of Assigned Values	Z Score	Deculto		Uncertainty of Assigned Values	Z Score	
WBC x10³/µl	1	7.84	7.59	15.43	14.5	0.0300	1.07	0.25	0.1	0.0070	1.35	
RBC x10 <sup>6</sup> /µl	1	4.39	4.31	8.7	8.5	0.0070	1.07	0.08	0.03	0.0020	1.69	
Hb g/dl	1	13.2	13.1	26.3	25.1	0.0200	2.02	0.1	0.1	0.0070	0.00	
HCT%	1	45.1	44. <mark>3</mark>	89.4	79.6	0.1720	1.69	0.8	0.3	0.0060	1.69	
MCV-fl	1	102.6	102. <mark>6</mark>	205.2	188	0.3400	1.38	0	0.2	0.0180	-0.67	
MCH-Pg	1	30.3	30.1	60.4	59.2	0.6740	0.89	0.2	0.2	0.0130	0.00	
MCHC-g/dl	1	29.5	29.3	58.8	62.7	0.1290	-0.87	0.2	0.2	0.0150	0.00	
Plt. x10³/µl	1	240	239	479	503	1.50	-0.56	1	6	0.31	-0.96	
Retic %	2	3.8	3.6	7.4	15.8	0.26	-1.12	0.2	0.4	0.02	-0.34	

### P.S . Assesment

		YOUR REPORT	CONSENSUS REPORT				
DLC%	3	Nrbcs=5 , Poly=67 L=24, E=4, Mono/Promono=5 , B1=0 P.M.=0, Mye=0, Meta=0, Other=0	Lympho: 37-47, Poly: 44-54, Mono: 2-5, Eosino: 1-5, blast/Promyelo/Myelo/Meta: 0				
RBC Morphology	3		Predominantly: Normocytic/Normochromic; Moderate: Anisocytosis, Microcytosis, Hypochromia; Mild: Poikilocytosis , Target cells , Sickle shaped cells ,tear drop cells				
Diagnosis	3	correlation with clinical features,hemato morphological features suggestive of hemolytic anemia possibility of thalassemia correlate with HB Electrophoresis	Diagnosis- Haemoglobinopathy/Thalassemia				

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

Test never store	S No	Total participants covered in the	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	5.NU.	current dist. 158A		Among labs	Within lab	Among labs	Within lab	Among labs	Within lab
WBC x10 <sup>3</sup> /µl	1	362	356	<mark>84</mark> .55	85.39	4.21	7.3	11.24	7.31
RBC x10 <sup>6</sup> /µl	1	362	362	84.25	90.61	8.56	3.59	7.19	5.8
Hb g/dl	1	362	362	87.85	91.71	4.97	2.49	7.18	5.8
HCT%	1	362	3 <mark>57</mark>	98.04	86.27	0.84	6.72	1.12	7.01
MCV-fl	1	362	357	99.44	87.11	0.28	9.52	0.28	3.37
MCH-Pg	1	362	357	87.96	<mark>9</mark> 2.44	5.6	2.52	6.44	5.04
MCHC-g/dl	1	362	357	97.2	93	1.12	3.08	1.68	3.92
Plt. x10³/µl	1	362	357	92.16	90.2	5.32	5.88	2.52	3.92
ReticCount%	2	362	340	93.53	93.24	4.12	5	2.35	1.76
PS Assessment	3	362	339	Satisfactory :96.68%, Borderline Sat. :2.76%, Unsatisfactory :0.552%					

#### \*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  $IQR = 0.7413 \times IQR$ 

**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,

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Dr. Seema Tyagi (Prof.) PT Co-ordinator: ISHTM-AIIMS-EQAP Department of Hematology, AIIMS, New Delhi

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