



## 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

**Instrument ID:** ERBA H-360, 3 PART (K10012116054)

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

Tel: 9013085730 , E-Mail : accuracy2000@gmail.com **Date of issue & status of the report:** 03-03-2022[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	12.41	11.94	24.35	10.4	0.0520	8.38	0.47	0.1	0.0110	2.94	
RBC x10 <sup>6</sup> /μl	1	4.13	3.81	7.94	8.44	0.0130	-1.73	0.32	0.04	0.0030	6.30	
Hb g/dl	1	12.6	12.1	24.7	25.4	0.0300	-0.94	0.5	0.1	0.0090	5.40	
НСТ%	1	39	36.1	75.1	79.85	0.1810	-1.05	2.9	0.4	0.0280	5.62	
MCV-fl	1	94.6	94.5	189.1	189.3	0.3200	-0.03	0.1	0.4	0.0300	-0.51	
MCH-Pg	1	33.1	29.4	62.5	60	0.0910	1.04	3.7	0.3	0.0200	11.47	
MCHC-g/dl	1	34.9	31.1	66	63.45	0.1430	0.67	3.8	0.3	0.0220	11.80	
Plt. x10³/μl	1	205	171	376	374	1.39	0.05	34	6	0.38	5.40	
Retic %	2	10	8	18	10.3	0.21	1.30	2	0.39	0.03	2.63	

#### P.S. Assesment

		YOUR REPORT	CONSENSUS REPORT				
DLC%	3		Poly: 30-50 ,Myelo: 14-29, blast: 1-8, Lympho: 3-7, Eosino: 2-5 , Promyelo: 2-8, nRBC/mono/Meta: 0-1				
RBC Morphology	3	MILD ANISOPOIKILOCYTOSIS, MICROCYTIC+, HYPOCHROMIC+, FEW POLYCHROMATIC RBCS	Predominantly: Normocytic Normochromic. Moderate: Anisocytosis. Mild: Microcytic				
Diagnosis	3	S/O CHRONIC MYELOID LEUKEMIA	Chronic Myeloid Leukemia (Chronic Phase) : CML-CP				

#### **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. 154F		Among labs	Within lab	Among labs	Within lab	Among labs	Within lab
WBC x10³/μl	1	272	268	<mark>76.</mark> 12	84.33	7.09	6.72	16.79	8.95
RBC x10 <sup>6</sup> /μl	1	272	272	87.5	88.24	6.62	2.94	5.88	8.82
Hb g/dl	1	272	272	86.76	86.4	6.99	4.04	6.25	9.56
HCT%	1	272	2 <mark>68</mark>	93.28	89.93	2.99	3.73	3.73	6.34
MCV-fl	1	272	268	89.55	95.15	7.09	1.87	3.36	2.98
MCH-Pg	1	272	268	92.91	<mark>8</mark> 7.31	4.48	5.22	2.61	7.47
MCHC-g/dl	1	272	268	94.4	87.31	2.99	5.97	2.61	6.72
Plt. x10³/μl	1	272	268	93.28	89.93	4.85	5.97	1.87	4.1
ReticCount%	2	272	269	97.03	91.45	1.12	1.49	1.85	7.06
PS Assessment	3	272	248	Satisfactory:84.32%, Borderline Sat.:15.32%, Unsatisfactory:0.36%					

#### \*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  $\pm 2$ : 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

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