



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.: 5598 **Distribution No.:** 164-N Month/Year: August/2024

Model Name.: XP1003P **Instrument ID: SYSMEX** Serial No.: A9049

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

 $Tel: 9013085730 \; , \; E\text{-Mail}: info@ishtmaiimseqap.com$ Date of issue & status of the report: 24-09-2024 [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	Consensus Result Diff. of 2 values (Assigned Value)	Uncertainty of Assigned Values	Z Score	
WBC x10³/μl	1											
RBC x10 ⁶ /μl	1	4.05	4.01	8.06	7.59	0.016	1.01	0.04	0.03	0.002	0.27	
Hb g/dl	1	11.2	11.1	22.3	21.1	0.038	0.90	0.1	0.1	0.008	0.00	
НСТ%	1	37.7	37. <mark>3</mark>	75	68.8	0.219	1.05	0.4	0.3	0.020	0.34	
MCV-fl	1	93.1	93	186.1	181.4	0.458	0.39	0.1	0.3	0.024	-0.54	
МСН-Рд	1	27.7	27.7	55.4	55.6	0.081	-0.10	0	0.2	0.012	-0.90	
MCHC-g/dl	1	29.8	29.7	59.5	61.85	0.187	-0.44	0.1	0.3	0.021	-0.83	
Plt. x10³/μl	1	168	164	332	335	2.347	-0.05	4	7	0.420	-0.51	
Retic %	2	1.5	1	2.5	3.05	0.087	-0.22	0.5	0.2	0.015	1.01	

P.S. Assesment

		YOUR REPORT	CONSENSUS REPORT				
DLC%		Nrbcs=9 , Poly=29 L=32, E=2, Mono/Promono=2 , B1=27 P.M.=3, Mye=0, Meta=0, Other=SMUDGE CELLS	Blast: 20-45, Lympho: 26.25-53, Poly: 10-18, Mono: 1-5 Myelo/Meta/Promyelo/Eos/Baso : 0-5				
RBC Morphology	3	NORMOCYTIC HYDOCHROMIC	Predominantly: Normocytic/ Normochromic, Moderate: Microcytic, Anisopoikilocytosis, Mild: Tear drop cells, Spherocytes				
Diagnosis	3	ACUTE LEUKAEMIA	Acute Leukemia (AL)				

COMBINED DATA VALUES OF TOTAL PARTICIPANTS

Test never eters	S.No.	Total participants covered in the current dist. 164N	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	285	283	<mark>85</mark> .16	91.17	14.84	3.53	0	5.3
RBC x10 ⁶ /μl	1	285	285	95.44	94.39	3.51	2.46	1.05	3.15
Hb g/dl	1	285	285	98.25	91.23	1.4	3.16	0.35	5.61
HCT%	1	285	284	95.07	88.03	3.52	6.69	1.41	5.28
MCV-fl	1	285	284	93.31	91.9	5.63	2.82	1.06	5.28
MCH-Pg	1	285	284	86.97	<mark>75</mark> .35	9.15	16.55	3.88	8.1
MCHC-g/dl	1	285	284	96.48	86.27	2.11	4.93	1.41	8.8
Plt. x10³/μl	1	285	285	96.84	88.42	2.46	7.37	0.7	4.21
ReticCount%	2	285	210	90	85.71	6.67	1.43	3.33	12.86
PS Assessment	3	285	212	Satisfactory	:82.76%, Bo	orderline Sat	.: :13.02%, U	Insatisfactor	y :4.22%

*Comments:

Among Lab (EQA): Results acceptable.
 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. Manoranjan Mahapatra (Prof. & Head)

PT Co-ordinator: ISHTM-AIIMS-EQAP

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