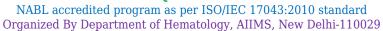




PROFICIENCY TESTING REPORT

ISHTM-AIIMS EXTERNAL QUALITY ASSURANCE PROGRAMME





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

Instrument ID: SYSMEX **Model Name.:** XN-350 **Serial No.:** 17249

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

Tel: 9013085730 , E-Mail : info@ishtmaiimseqap.com **Date of issue & status of the report:** 01-04-2024[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		Yours Results Diff. of 2 Values	Consensus Result Diff. of 2 values (Assigned Value)	Uncertainty of Assigned Values	Z Score	
WBC x10³/μl	1	6.06	5.7	11.76	11.59	0.034	0.20	0.36	0.1	0.007	2.19	
RBC x10 ⁶ /μl	1	4.22	4.18	8.4	8.58	0.011	-0.66	0.04	0.04	0.003	0.00	
Hb g/dl	1	10.7	10.6	21.3	21.9	0.021	-1.01	0.1	0.1	0.007	0.00	
НСТ%	1	36.9	36. <mark>5</mark>	73.4	73.1	0.207	0.05	0.4	0.3	0.022	0.27	
MCV-fl	1	87.4	87.3	174.7	171.85	0.394	0.24	0.1	0.2	0.021	-0.34	
МСН-Рд	1	25.4	25.4	50.8	51	0.057	-0.13	0	0.2	0.013	-0.90	
MCHC-g/dl	1	29	29	58	59.45	0.159	-0.30	0	0.3	0.017	-1.25	
Plt. x10³/μl	1	196	193	389	409	1.547	-0.48	3	7	0.389	-0.65	
Retic %	2	15.4	12.13	27.53	24.9	0.441	0.18	3.27	1	0.061	1.91	

P.S. Assesment

		YOUR REPORT	CONSENSUS REPORT				
DLC%	3		Blast: 22-49.75, Poly: 14-30, Lympho: 4-11, Myelo: 3-15, Meta: 2-10, Mono: 1-5, Eos: 1-4, Promyelo: 0-9.25, nRBC/Baso: 0-5				
RBC Morphology	3		Predominantly: Normocytic/ Normochromic, Moderate: Anisocytosis, Microcytic, poikilocytosis				
Diagnosis	3	CML-CMPD	Acute Leukemia (AL)				

COMBINED DATA VALUES OF TOTAL PARTICIPANTS

Test never eters	S.No.	Total participants covered in the current dist. 162L	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	316	316	80.38	89.56	4.75	4.75	14.87	5.69	
RBC x10 ⁶ /μl	1	316	316	87.97	92.41	7.91	5.38	4.12	2.21	
Hb g/dl	1	316	316	88.92	92.41	6.65	2.53	4.43	5.06	
HCT%	1	316	3 <mark>16</mark>	96.84	91.77	2.85	4.43	0.31	3.8	
MCV-fl	1	316	316	95.89	89.24	3.48	5.06	0.63	5.7	
MCH-Pg	1	316	316	91.14	<mark>74.</mark> 37	4.75	19.94	4.11	5.69	
MCHC-g/dl	1	316	316	96.2	90.82	3.48	4.43	0.32	4.75	
Plt. x10³/μl	1	316	316	91.77	92.09	6.65	4.75	1.58	3.16	
ReticCount%	2	316	212	93.87	93.4	4.72	2.83	1.41	3.77	
PS Assessment	3	316	198	Satisfactory:89.57%, Borderline Sat.:3.79%, Unsatisfactory:6.64%						

*Comments:

1). Among Lab (EQA): PS Diagnosis wrongly reported, 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 ± 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

Department of Hematology, AIIMS, New Delhi

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