



PROFICIENCY TESTING REPORT

ISHTM-AHMS EXTERNAL QUALITY ASSURANCE PROGRAMME NABL accredited program as per ISO/IEC 17043:2010 standard Organized By Department of Hematology, AIIMS, New Delhi-110029



 $Duration\ of\ stability\ testing\ -\ minimum\ up to\ 8\ days\ at\ ambient\ temp.\ after\ dispatch\ of\ specimens$

EQAP CODE No.: 433

Distribution No.: 163-A

Month/Year: January/2024

Instrument ID: Sysmex J K Biomed

Model Name.: Sysmex XN 1000

Serial No.: 51194

Name & Contact No. of PT Co-ordinator: Dr. Manoranjan Mahapatra (Prof. & Head), Hematology, AIIMS, Delhi, Tel: 9013085730 , E-Mail : accuracy2000@gmail.com

Date of issue & status of the report: 05-04-2024[Final].

CBC and Retic Assessment

Test Parameters	S.No.			Amo	ng Lab (Ac	curacy Testi	Within Lab (Precision Testing)				
		Your Result	Your Result 2	Your Results Sum of 2 Value		Uncertainty of Assigned Values		Yours Results Diff. of 2 Values	Consensus Result Diff. of 2 values (Assigned Value)	Uncertainty of Assigned Values	
WBC x10³/µl	1	8.28	8.27	16.55	16.6	0.050	-0.04	0.01	0.11	0.007	-0.96
RBC x10 ⁶ /μl	1	5.14	5.13	10.27	10.18	0.009	0.35	0.01	0.04	0.003	-0.67
Hb g/dl	1	13.5	13.4	26.9	26.53	0.019	0.71	0.1	0.1	0.007	0.00
НСТ%	1	42.1	41.9	84	83.9	0.145	0.02	0.2	0.3	0.022	-0.27
MCV-fl	1	81.9	81.7	163.6	165.15	0.234	-0.21	0.2	0.2	0.018	0.00
МСН-Рд	1	26.3	26.1	52.4	52.2	0.045	0.15	0.2	0.2	0.012	0.00
MCHC-g/dl	1	32.1	32	64.1	63.1	0.111	0.31	0.1	0.2	0.015	-0.45
Plt. x10³/μl	1	95	85	180	116.5	1.599	1.26	10	5	0.302	0.96
Retic %	2	10.3	10.2	20.5	27.4	0.394	-0.62	0.1	0.6	0.047	w().71

P.S. Assesment

YOUR REPORT			CONSENSUS REPORT				
DLC%	3	Nrbcs=07, Poly=59 L=32, E=03, Mono/Promono=06, B1=00 P.M.=00, Mye=00, Meta=00, Other=	Poly: 56 - 65, Lympho: 30- 38, Mono: 2-4, Eosino: 1-3, Myelo/Meta/Promyelo/Blast/Baso: 0-5				
RBC Morphology	3	Normocytic nomrochromic with anisopoikilocytosis showing good number of spherocytes and nRBCs.	Predominantly: Normocytic/Normochromic with Mild Anisocytosis				
Diagnosis	3	Normocytic normochromic anemia with features of hemolysis. D/D: Hereditary spehrocytosis/ Autoimmune hemolytic anemia	SPherocytic Hemolytic Anemia				

COMBINED DATA VALUES OF TOTAL PARTICIPANTS

	S.No.	163A	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
				86.52	91.01	6.74	1.69	6.74	7.3
WBC x10³/μl	1	357		90.2	89.36	5.04	3.92	4.76	6.72
RBC x10 ⁶ /µl	1	357	357		89.92	4.76	4.76	5.04	5.32
Hb g/dl	1	357	357	90.2	91.01	4.49	1.97	2.53	7.02
HCT%	1	357	356	92.98		3.65	6.18	0.84	5.62
MCV-fl	1	357	356	95.51	88.2	5.62	3.37	4.21	5.34
MCH-Pg	1	357	356	90.17	91.29		3.37	1.68	7.02
	1	357	356	92.98	89.61	5.34		2.8	6.74
MCHC-g/dl	1	357	356	95.51	89.89	1.69	3.37		4.18
Plt. x10³/µl	1		335	93.13	91.94	5.07	3.88	1.8	
ReticCount%	2	357	317	Satisfactory:89.64%, Borderline Sat.:5.32%, Unsatisfactory:5.04%					
PS Assessment	3	357	317	Juliotavoor					

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

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 ± 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 $(\overline{x}-\overline{y})$ 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

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