



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. : 4496 **Distribution No.:** 161-L Month/Year: October/2023

Instrument ID: Medonic

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

 $Tel: 9013085730 \; , \; E\text{-Mail}: accuracy 2000@gmail.com$ Date of issue & status of the report: 31-01-2024[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	2.9	2.7	5.6	4.95	0.040	0.58	0.2	0.07	0.005	1.95	
RBC x10 ⁶ /μl	1	4.62	4.45	9.07	9.22	0.011	-0.52	0.17	0.04	0.003	2.92	
Hb g/dl	1	14.8	14	28.8	30.1	0.028	-1.93	0.8	0.1	0.008	4.72	
НСТ%	1	46.9	45.8	92.7	96.2	0.206	-0.58	1.1	0.4	0.027	1.89	
MCV-fl	1	102.9	101.6	204.5	209.5	0.364	-0.49	1.3	0.3	0.024	2.70	
MCH-Pg	1	32	31.5	63.5	65.3	0.069	-1.01	0.5	0.2	0.017	1.35	
MCHC-g/dl	1	31.5	30.6	62.1	62.2	0.126	-0.03	0.9	0.2	0.017	2.36	
Plt. x10³/μl	1	133	128	261	306	1.239	-1.29	5	5	0.314	0.00	
Retic %	2	4.6	4.4	9	10.25	0.216	-0.19	0.2	0.5	0.044	-0.51	

P.S. Assesment

		YOUR REPORT	CONSENSUS REPORT				
DLC%	3	Nrbcs=2 , Poly=80 L=16, E=2, Mono/Promono=2 , B1= P.M.=, Mye=, Meta=, Other=	Poly: 53-65, Lympho: 25-35, Eosino: 2-6, Mono: 2-5, blast/Promyelo/Myelo/Meta: 0-5				
RBC Morphology	3	Rbcs are normocytic normochromic with moderate degree of anisopoikilocytosi composed of tear drop cells, elliptocytes ,schiztocytes	Predominantly: Microcytic, Hypochromic, Moderate: Anisopoikilocytosis Mild:Target cells , Tear drop cells				
Diagnosis	3	Hemolytic anaemia	Thalassemia Haemoglobinopathy				

COMBINED DATA VALUES OF TOTAL PARTICIPANTS

Test parameters	S.No.	Total participants covered in the current dist. 161L	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	301	301	<mark>87</mark> .38	91.03	7.64	2.99	4.98	5.98	
RBC x10 ⁶ /μl	1	301	301	87.71	88.37	6.64	5.98	5.65	5.65	
Hb g/dl	1	301	301	86.71	83.39	7.31	6.64	5.98	9.97	
HCT%	1	301	3 <mark>01</mark>	94.35	87.71	2.66	4.65	2.99	7.64	
MCV-fl	1	301	301	94.68	94.02	3.99	2.66	1.33	3.32	
MCH-Pg	1	301	301	86.38	90.7	7.31	3.65	6.31	5.65	
MCHC-g/dl	1	301	301	93.36	91.36	3.99	4.65	2.65	3.99	
Plt. x10³/μl	1	301	301	94.02	90.37	3.32	3.65	2.66	5.98	
ReticCount%	2	301	186	95.7	87.1	3.23	8.06	1.07	4.84	
PS Assessment	3	301	177	Satisfactory:94.03%, Borderline Sat.:3.32%, Unsatisfactory:2.65%						

*Comments:

1). Among Lab (EQA): Results acceptable.

2). Within Lab (IQA): Difference in the CBC measurement values for *HB* unacceptable, may be due to random/human error.

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 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,

Dr. Manoranjan Mahapatra (Prof. & Head)

PT Co-ordinator: ISHTM-AIIMS-EQAP

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