



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.: 3953

Distribution No.: 157-H

Month/Year: September/2022

Instrument ID: B6923

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

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Date of issue & status of the report: 07-11-2022[Final].

CBC and Retic Assessment

Test Parameters	S.No.			Among Lab (Accuracy Testing)				Within Lab (Precision Testing)				
		Your Result 1		Your Results Sum of 2 Value	Consensus result sum of 2 values (Assigned Value)	Uncertainty of Assigned Values	Z Score	Yours Results Diff. of 2 Values	Consensus Result Diff. of 2 values (Assigned Value)	Uncertainty of Assigned Values		
WBC x10³/μl	1	5.6	5.4	11	9.03	0.0420	3.14	0.2	0.1	0.0090	1.04	
RBC x10 ⁶ /μl	1	4.84	4.74	9.58	9.38	0.0150	0.76	0.1	0.05	0.0050	0.84	
Hb g/dl	1	12.2	12 -	24.2	24.9	0.0410	-1.05	0.2	0.1	0.0110	1.35	
НСТ%	1	43	42.4	85.4	80.25	0.2890	1.04	0.6	0.4	0.0430	0.45	
MCV-fl	1	89.5	88.8	178.3	170.45	0.4740	0.84	0.7	0.3	0.0410	0.77	
МСН-Рд	1	25.3	25.2	50.5	52.9	0.1010	-1.52	0.1	0.2	0.0230	-0.45	
MCHC-g/dl	1	28.4	28.3	56.7	61.75	0.2320	-1.29	0.1	0.3	0.0300	-0.54	
Plt. x10³/μl	1	204	194	398	367.5	2.38	0.71	10	7	0.61	0.45	
Retic %	2	5.5	4.8	10.3	9.5	0.28	0.14	0.7	. 0.4	0.04	0.77	

P.S. Assesment

YOUR REPORT			CONSENSUS REPORT				
DLC%	3	Nrbcs=33, Poly=37 L=23, E=05, Mono/Promono=02, B1=00 P.M.=00, Mye=00, Meta=00, Other=	Poly: 42-56, Lympho: 28-40, Eosino: 5-12, Mono: 2-5, blast/Promyelo/Myelo/Meta: 0				
RBC Morphology	3	Microcytic hypochromic, target cells, macrocytes, sickled RBC's.	Predominantly: Normocytic/Normochromic; Moderate: Anisocytosis, Microcytosis, Hypochromia; Mild: Poikilocytosis, Target cells, Sickle shaped cells, tear drop cells				
Diagnosis	3	Hemoglobinopathy (Sickle cell Anemia or sickle cell-Thalassemia compound heterozygote) with secondary megaloblastic changes.	Hemoglobinopathy possible sickle cell anemia				

COMBINED DATA VALUES OF TOTAL PARTICIPANTS

T	S.No.	Total participants covered in the current dist. 157H	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	136	136	86.76	90.44	4.41	2.94	8.83	6.62
RBC x10 ⁶ /µl	1	136	136	90.44	91.18	3.68	2.94	5.88	5.88
Hb g/dl	1	136	136	91.91	91.18	3.68	5.15	4.41	3.67
HCT%	1	136	136	89.71	88.97	5.88	5.88	4.41	5.15
MCV-fl	1	136	136	91.18	93.38	5.88	1.47	2.94	5.15
MCH-Pg	1	136	136	83.82	90.44	8.82	5.15	7.36	4.41
MCHC-g/dl	1	136	136	88,97	88.24	5.88	6.62	5.15	5.14
Plt. x10³/µl	1	136	136	93.38	95.59	3.68	2.21	2.94	2.2
ReticCount%	2	136	107	97.2	96.26	1.87	1.87	0.93	1.87
PS Assessment	3	136	105	Satisfactory	:89.71%, B	orderline Sat	.: 7.35%, U1	nsatisfactory	:2.94%

*Comments:

- 1). Among Lab (EQA): CBC result for WBC unacceptable, may be due to random/human error
- 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 > ± 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 > ± 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. Seema Tyagi (Prof.)

PT Co-ordinator: ISHTM-AIIMS-EQAP

Department of Hematology, AIIMS, New Delhi

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