

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

Month/Year: February/2021

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

Distribution No.: 152-D

Instrument ID: 710YOXH01128

EQAP CODE No.: 1302

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

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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 of Assigned Values	Z Score	Doculte	Consensus Result Diff. of 2 values (Assigned Value)	Uncertainty of Assigned Values	Z Score	
WBC x10³/μl	1	1.06	0.92	1.98	5.5	0.0200	-6.17	0.14	0.1	0.0050	0.62	
RBC x10 ⁶ /μl	1	4.57	4.47	9.04	9.32	0.0080	-1.35	0.1	0.03	0.0020	2.36	
Hb g/dl	1	13.7	13.7	27.4	27	0.0200	0.67	0	0.1	0.0070	-1.35	
НСТ%	1	39.1	38.3	77.4	83.9	0.1780	-1.11	0.8	0.3	0.0200	1.35	
MCV-fl	1	85.8	85.7	171.5	179.9	0.3170	-0.80	0.1	0.2	0.0170	-0.45	
МСН-Рд	1	30.6	29.9	60.5	57.7	0.0540	1.84	0.7	0.2	0.0140	2.25	
MCHC-g/dl	1	35.7	34.9	70.6	64.15	0.1300	1.44	0.8	0.3	0.0120	1.69	
Plt. x10³/μl	1	127	121	248	196	0.75	2.44	6	4	0.26	0.45	
Retic %	2	2	1.7	3.7	5	0.08	-0.57	0.3	0.2	0.01	0.34	

P.S. Assesment

		YOUR REPORT	CONSENSUS REPORT				
DLC%	3	Mono/Promono=3 , B1=- P.M.=-, Mye=-, Meta=-, Other=	nRBC: 30 - 65, Poly: 60 - 75, Lympho: 15-30, Eos/Mono: 1-5, Blast/Myelo/Meta: 0-1				
RBC Morphology	3	macrocytes, microcytic hypochromic cells, target cells, tear drop cells, polychromatophils and nucleated erythrocytes	Predominantly: Macrocytosis, Microcytosis, Spherocytosis, Polychromasia, Anisocytosis; Moderate: Normocytic/Normochromic, Hypo.				
Diagnosis	3	Haemolytic Anaemia	Hemolytic Anemia				

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COMBINED DATA VALUES OF TOTAL PARTICIPANTS

Test	S.No.	Total participants covered in	Total No. responded	% of Labs with Z Score 0-2		% of Labs with Z Score 2-3		% of Labs with Z Score >3		
parameters		the current dist.		Among labs	Within lab	Among labs	Within lab	Among labs	Within lab	
WBC x10³/μl	1	312	346	89.31	92.2	2.31	1.45	8.38	6.36	
RBC x10 ⁶ /μl	1	312	346	89.88	91.04	6.94	3.47	3.18	5.2	
Hb g/dl	1	312	347	91.07	99.42	6.92	3.17	2.02	0.58	
НСТ%	1	312	346	97.69	92.2	1.73	3.47	0.58	4.34	
MCV-fl	1	312	345	97.68	86.09	1.45	8.99	0.87	4.93	
MCH-Pg	1	312	346	91.62	91.04	6.07	3.47	2.31	5.49	
MCHC-g/dl	1	312	346	98.55	92.2	0.29	3.76	1.16	3.76	
Plt. x10³/μl	1	312	346	93.35	91.91	3.47	5.49	3.18	2.6	
ReticCount%	2	312	318	93.71	86.48	4.09	2.2	2.2	11.64	
PS Assessment	3	312	312	Acceptable:91.4,Warning Signal:7.7,Unacceptable:0.9						

*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 $> \pm 3$ 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.ishtmaiimsegap.com.

Report authorized by,

Dr. Seema Tyagi (Prof.)

PT Co-ordinator: ISHTM-AIIMS-EQAP

Department of Hematology, AIIMS, New Delhi

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