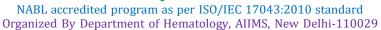




PROFICIENCY TESTING REPORT

ISHTM-AIIMS EXTERNAL QUALITY ASSURANCE PROGRAMME





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

EQAP CODE No.: 5988 **Distribution No.:** 159-F **Month/Year:** June/2023

Instrument ID: CELLENIUM 21 (3020ET-03772)

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

Tel: 9013085730 , E-Mail : accuracy2000@gmail.com **Date of issue & status of the report:** 01-08-2023[Final].

CBC and Retic Assessment

				Among Lab (Accuracy Testing)				Within Lab (Precision Testing)				
Test Parameters	S.No.	Your Result 1	Your Result 2		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	Z Score	
WBC x10³/μl	1	8.2	7.9	16.1	15.33	0.039	0.75	0.3	0.1	0.008	1.80	
RBC x106/μl	1	3.64	3.58	7.22	6.84	0.008	1.90	0.06	0.03	0.002	1.01	
Hb g/dl	1	10.6	10.5	21.1	21.1	0.021	0.00	0.1	0.1	0.007	0.00	
НСТ%	1	37.5	37.3	74.8	67.1	0.167	1.61	0.2	0.4	0.023	-0.67	
MCV-fl	1	90.4	89.4	179.8	195.3	0.416	-1.28	1	0.3	0.022	1.89	
MCH-Pg	1	29.6	28.8	58.4	61.7	0.077	-1.82	8.0	0.3	0.018	1.69	
MCHC-g/dl	1	32.8	32.3	65.1	63.1	0.160	0.43	0.5	0.3	0.021	0.67	
Plt. x10³/μl	1	170	168	338	458	1.916	1.61	0.2	0.4	0.023	-0.67	
Retic %	2	45.1	45	90.1	43	0.840	1.83	0.1	1	0.092	-0.81	

P.S. Assesment

		YOUR REPORT	CONSENSUS REPORT				
DLC%	3	Nrbcs=10 , Poly=74 L=24, E=01, Mono/Promono=01 , B1=0 P.M.=0, Mye=0, Meta=0, Other=0	Poly: 54-67, Lympho: 22-33, Mono: 2-5, Eosino: 1-3, blast/Promyelo/Myelo/Meta: 0-5				
RBC Morphology	3	macrocytosis present anisopolkilocytosis marked, nucleted rbc -: 10/100wbc, platelet-: 82x1000/ul, mp absent, abnormal cell absent,	Predominantly: Normocytic/Normochromic; Moderate: Anisocytosis, Microcytosis, Hypochromia; Mild: Poikilocytosis, Target cells, Sickle shaped cells, tear drop cells				
Diagnosis	3	Macrocytic anemia	Thalassemia/Haemoglobinopathy				

COMBINED DATA VALUES OF TOTAL PARTICIPANTS

To at monomorphous	S.No.	Total participants covered in the current dist. 159F	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 $x10^3/\mu l$	1	317	315	<mark>84</mark> .76	82.22	2.54	10.79	12.7	6.99
RBC x10 ⁶ /μl	1	317	317	83.91	88.01	8.52	5.68	7.57	6.31
Hb g/dl	1	317	317	86.44	92.11	5.68	3.79	7.88	4.1
НСТ%	1	317	3 <mark>15</mark>	93.65	93.02	3.81	2.54	2.54	4.44
MCV-fl	1	317	315	94.29	91.75	5.08	3.49	0.63	4.76
MCH-Pg	1	317	315	85.71	88.89	6.67	4.76	7.62	6.35
MCHC-g/dl	1	317	315	95.24	91.43	3.49	4.76	1.27	3.81
Plt. x10³/μl	1	317	315	89.52	88.89	7.3	5.08	3.18	6.03
ReticCount%	2	317	282	98.23	87.94	1.42	3.55	0.35	8.51
PS Assessment	3	317	282	Satisfactory:89.28%, Borderline Sat.:6.94%, Unsatisfactory:3.78%					

*Comments:

- 1). Among Lab (EQA): PS Diagnosis partially correct, remaining 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 > ±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 $(\bar{x-y})$ 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

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