

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.: 1207 **Distribution No.**: 156-C **Month/Year**: May/2022

Instrument ID: YUMIZEN H550 (006YAXH03049)

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:** 12-07-2022[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		Uncertainty of Assigned Values	Z Score	
WBC x10³/μl	1	4.13	3.81	7.94	41.35	0.2760	-1.56	0.32	0.21	0.0140	0.37	
RBC x10 ⁶ /μl	1	4.35	4.2	8.55	8.63	0.0080	-0.35	0.15	0.03	0.0020	3.24	
Hb g/dl	1	13.4	13	26.4	26.4	0.0330	0.00	0.4	0.1	0.0080	2.02	
НСТ%	1	40.4	39.7	80.1	83.6	0.2070	-0.56	0.7	0.4	0.0240	0.81	
MCV-fl	1	94.5	92.8	187.3	194.1	0.4040	-0.57	1.7	0.3	0.0250	2.36	
MCH-Pg	1	30.9	30.7	61.6	61.1	0.0840	0.22	0.2	0.2	0.0160	0.00	
MCHC-g/dl	1	33.1	32.7	65.8	62.6	0.1590	0.70	0.4	0.3	0.0170	0.34	
Plt. x10³/μl	1	211	198	409	346	3.20	0.62	13	5	0.31	1.44	
Retic %	2	7.7	7.2	14.9	7.85	0.12	2.08	0.5	0.3	0.02	0.90	

P.S. Assesment

		YOUR REPORT	CONSENSUS REPORT				
DLC%	3		Blast: 49-70, Lympho: 12-27 ,Poly: 9-17,/mono:1-5 nRBC/Eosino/Myelo/Meta/promyelo: 0-2				
RBC Morphology	.5		Predominantly: Normocytic/ Normochromic, Moderate: Anisocytosis, Microcytic, macrocytes, Tear drop cells				
Diagnosis	3	Acute Leukemia S/o AML	Acute Leukemia (AL)				

COMBINED DATA VALUES OF TOTAL PARTICIPANTS

Test never eters	S.No.	Total participants covered in the current dist. 156C	Total No. responded	% of Labs with Z Score 0-2		% of Labs with Z Score 2-3		% of Labs with Z Score >3		
rest parameters				Among labs	Within lab	Among labs	Within lab	Among labs	Within lab	
WBC x10³/μl	1	348	346	99.13	89.88	0.29	3.47	0.58	6.65	
RBC x10 ⁶ /μl	1	348	348	91.09	90.8	4.6	3.74	4.31	5.46	
Hb g/dl	1	348	348	85.92	87.93	6.61	4.6	7.47	7.47	
HCT%	1	348	3 <mark>47</mark>	88.47	89.05	4.9	3.75	6.63	7.2	
MCV-fl	1	348	347	88.47	91.93	3.75	4.61	7.78	3.46	
MCH-Pg	1	348	347	87.32	<mark>8</mark> 7.61	5.76	6.92	6.92	5.47	
MCHC-g/dl	1	348	347	91.64	86.46	4.9	4.9	3.46	8.64	
Plt. x10³/μl	1	348	347	95.39	91.64	3.46	3.46	1.15	4.9	
ReticCount%	2	348	330	90.91	82.12	7.27	12.73	1.82	5.15	
PS Assessment	3	348	327	Satisfactory:94.27%, Borderline Sat.:4.59%, Unsatisfactory:1.14%						

*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 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 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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