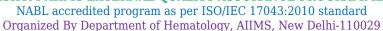




PROFICIENCY TESTING REPORT

ISHTM-AIIMS EXTERNAL QUALITY ASSURANCE PROGRAMME





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

EQAP CODE No.: 5081 **Distribution No.:** 158-M **Month/Year:** January/2023

Instrument ID: Eurocount 5L 5part 952305022IEJCP

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:** 28-02-2023[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	9.11	8.97	18.08	13.31	0.0350	5.46	0.14	0.1	0.0070	0.42	
RBC x10 ⁶ /μl	1	5	4.99	9.99	10.25	0.0120	-0.72	0.01	0.05	0.0030	-0. 77	
Hb g/dl	1	13.5	13.2	26.7	26.5	0.0270	0.27	0.3	0.1	0.0080	1.35	
НСТ%	1	46.6	46.5	93.1	85.7	0.2190	0.99	0.1	0.4	0.0250	-0.77	
MCV-fl	1	93.4	93.1	186.5	169.35	0.3530	1.49	0.3	0.3	0.0190	0.00	
MCH-Pg	1	27.1	26.5	53.6	51.5	0.0590	1.29	0.6	0.2	0.0140	1.80	
MCHC-g/dl	1	29	28.5	57.5	61	0.1400	-0.64	0.5	0.3	0.0210	0.67	
Plt. x10³/μl	1	414	383	797	781	2.99	0.18	31	9	0.52	2.70	
Retic %	2	8.1	7.4	15.5	15.35	0.22	0.03	0.7	0.5	0.03	0.34	

P.S. Assesment

		YOUR REPORT	CONSENSUS REPORT				
DLC%	3		Blast: 65-88, Lympho: 5-14, Poly: 2-5, nRBC/Eos/Baso/Mono /Myelo/Meta/ Promyelo: 0-5				
RBC Morphology	3		Predominantly: Normocytic/ Normochromic, Moderate: Anisocytosis, Microcytic				
Diagnosis	3	Acute myeloid leukemia with thrombocytopenia .	Acute Leukemia (AL)				

COMBINED DATA VALUES OF TOTAL PARTICIPANTS

Test never eters	S.No.	Total participants covered in the current dist. 158M	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	338	335	83.88	84.18	6.27	6.27	9.85	9.55	
RBC x10 ⁶ /μl	1	338	338	89.64	89.35	5.92	5.03	4.44	5.62	
Hb g/dl	1	338	338	88.76	85.21	5.03	4.44	6.21	10.35	
HCT%	1	338	3 <mark>36</mark>	97.92	89.88	0.89	5.36	1.19	4.76	
MCV-fl	1	338	336	97.62	87.2	1.79	6.55	0.59	6.25	
MCH-Pg	1	338	336	88.39	<mark>8</mark> 9.58	7.44	4.76	4.17	5.66	
MCHC-g/dl	1	338	336	98.21	86.9	0.89	7.74	0.9	5.36	
Plt. x10³/μl	1	338	336	94.64	92.26	3.27	2.98	2.09	4.76	
ReticCount%	2	338	296	91.22	85.14	6.08	9.8	2.7	5.06	
PS Assessment	3	338	283	Satisfactory: 97.93%, Borderline Sat.: 1.18%, Unsatisfactory: 0.890%						

*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 $> \pm 3$: 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

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