



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.: 1879 **Distribution No.:** 157-E Month/Year: August/2022

Instrument ID: A7125

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: 22-10-2022[Final].

CBC and Retic Assessment

	S.No.			Among Lab (Accuracy Testing)				Within Lab (Precision Testing)				
Test Parameters		Your Result 1		Your Results Sum of 2 Value	Consensus result sum of 2 values (Assigned Value)	Uncertainty		Yours Results Diff. of 2 Values	Consensus Result Diff. of 2 values (Assigned Value)	Uncertainty of Assigned Values	Z Score	
WBC x10³/μl	1	9.9	9.8	19.7	17.38	0.1220	0.74	0.1	0.17	0.0110	-0.47	
RBC x10 ⁶ /μl	1	5.69	5.67	11.36	10.98	0.0110	1.27	0.02	0.04	0.0030	-0.45	
Hb g/dl	1	10.9	10.9	21.8	21.1	0.0220	1.18	0	0.1	0.0070	-1.35	
НСТ%	1	40.3	40.2	80.5	71.3	0.1690	2.01	0.1	0.3	0.0230	-0.54	
MCV-fl	1	70.9	70.8	141.7	129.15	0.2640	1.68	0.1	0.2	0.0120	-0.45	
МСН-Рд	1	19.2	19.2	38.4	38.5	0.0510	-0.08	0	0.1	0.0090	-0.79	
MCHC-g/dl	1	27.1	27.1	54.2	59.2	0.1610	-1.12	0	0.2	0.0130	-0.90	
Plt. x10³/μl	1	288	285	573	465	2.38	1.64	3	7	0.49	-0.49	
Retic %	2	4.3	4	8.3	8.5	0.17	-0.04	0.3	0.4	0.02	-0.34	

P.S. Assesment

		YOUR REPORT	CONSENSUS REPORT				
DLC%	3		Poly: 19 - 36, Myelo: 20 - 40, Meta: 9 - 20, Lympho: 2 - 6, Promyelo: 1 - 6, nRBC/ Baso/ Eos/ Mono /Blast: 0 - 5				
RBC Morphology	3		Predominantly: Normocytic/Normochromic; Moderate: Anisocytosis, hypochromia, Microcytosis; Mild: Macrocytosis, Poikilocytosis				
Diagnosis	3	Plasma Cell Leukemia.	Chronic Myeloid Leukemia (Chronic Phase)				

COMBINED DATA VALUES OF TOTAL PARTICIPANTS

Test never eters	S.No.	Total participants	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		current dist. 157E		Among labs	Within lab	Among labs	Within lab	Among labs	Within lab
WBC x10³/μl	1	313	311	<mark>85</mark> .85	89.07	10.29	4.18	3.86	6.75
RBC x10 ⁶ /μl	1	313	313	87.86	90.42	5.43	3.83	6.71	5.75
Hb g/dl	1	313	313	85.94	90.73	5.11	4.47	8.95	4.8
HCT%	1	313	3 <mark>10</mark>	92.26	89.03	4.52	4.52	3.22	6.45
MCV-fl	1	313	310	91.94	88.39	3.87	9.03	4.19	2.58
MCH-Pg	1	313	309	85.76	91.91	5.5	4.21	8.74	3.88
MCHC-g/dl	1	313	309	91.59	91.91	5.18	4.53	3.23	3.56
Plt. x10³/μl	1	313	311	94.53	87.14	4.5	6.43	0.97	6.43
ReticCount%	2	313	288	97.92	93.4	1.39	1.04	0.69	5.56
PS Assessment	3	313	289	Satisfactory	:74.77%, Bo	rderline Sat	.: 9.58%, Uı	nsatisfactory	:15.65%

*Comments:

- 1). Among Lab (EQA): PS Diagnosis wrongly reported, 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 $> \pm 3$: 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

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