



PROFICIENCY TESTING REPORT

ISHTM-AHMS 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.: 4035

Distribution No.: 152-K

Month/Year: March/2021

Instrument ID: ERBA H560- SR k11042109048

Tel: 9013085730, E-Mail: accuracy2000@gmail.com Date of issue & status of the report: 24-05-2021[Final].

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

CBC and Retic Assessment

					Tab (Acc	curacy Testin	ia)	With	nin Lab (Pre	cision Testin	9)
Test arameters	S.No.	Your Result 1	Your Result 2	Your Results Sum of 2 Value	Consensus result sum of 2 values (Assigned	Uncertainty of Assigned Values	7	Yours Results Diff. of 2 Values	values	Uncertainty of Assigned Values	Z Score
		10.13	9,93	20.05	Value) 15.4	0.3590	0.64	0.19	0.13	0.0180	0.37
WBC x10³/μl	1	10.12		-	8.28	0.0130	1.86	0.04	0.05	0.0040	-0.27
RBC x10 ⁶ /μl	1	4.36	4.32	8.68		0.0420	0.27	0.1	0.1	0.0110	0.00
Hb g/dl	1	12.1	12	24.1	23.9		0.53	+	0.4	0.0370	0.00
нст%	1	38.9	38.5	77.4	74.3	0.3300	+		0.2	0.0310	-0.27
MCV-fl	1	89.3	89.	1 178.3	3 178.7	0.5920	-0.0			0.0260	-1.3
мсн-Рд	1	27.	7 27.	8 55.5	57.8	0.1200	-1.1	-0.	0.2		-
MCHC-g/	+	3	31	.2 62.	2 63.7	0.2690	-0.	31 -0.	2 0.2	0.0280	-1.3
	+	1 22	22 23	31 45	360	3.51	1.	60 -	9	0.85	-2.
Plt. x103	ш	+	.2	2 4	.2 5	0.15	-0	.27	0.3	0.10	-0.

P.S . Assesment

		PEROPT	CONSENSUS REPORT				
DLC%		YOUR REPORT Nrbcs=02, Poly=30 L=10, E=, Mono/Promono=, B1=04 P.M.=10, Mye=28, Meta=14, Other=baso 02	Poly: 30 - 65, Myelo: 10 - 35, Meta: 5 - 20, Promyelo/Blast/Lympho: 1 - 10, nRBC/Baso/Eos/Mono: 0 - 5				
RBC Morphology	3	Red blood cells predominately normocytic	Predominantly: Normocytic/Normochromic; Moderate: Anisocytosis, Hypochromia, Microcytosis; Mild: Poikilocytosis, Macrocytosis				
Diagnosis		Chronic myeloid leukemia	Chronic Myeloid Leukemia (CML)				

COMBINED DATA VALUES OF TOTAL PARTICIPANTS

Test parameters	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		
	0.110.	the current dist.		Among labs	Within lab	Among labs	Within lab	Among labs	Within lab	
WBC x10³/μl	1	124	137	85.4	85.4	8.03	8.03	5.84	5.84	
RBC x10 ⁶ /μl	1	124	137	87.59	91.24	5.11	3.65	6.57	4.38	
Hb g/dl	1	124	137	89.78	89.78	5.84	1.46	3.65	8.03	
HCT%	1	124	137	92.7	90.51	5.11	2.92	1.46	5.84	
MCV-fl	1	124	137	93.43	86.86	4.38	3.65	1.46	8.76	
MCH-Pg	1	124	137	91.24	90.51	3.65	2.92	4.38	5.84	
MCHC-g/dl	1	124	137	93.43	86.13	4.38	5.11	1.46	8.03	
Plt. x10³/μl	1	124	137	89.05	86.13	5.84	5.11	3.65	7.3	
ReticCount%	2	124	122	93.44	85.25	3.28	0.82	3.28	13.93	
PS Assessment	3	124	128	Acceptable:92, Warning Signal:4, Unacceptable:4						

'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)

 $IQR = Quartile 3 - Quartile 1 of participant data, Normalised <math>IQR = 0.7413 \times IQR$

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.

Report authorized by,

Dr. Seema Tyagi (Prof.)

PT Co-ordinator: ISHTM-AIIMS-EQAP

Department of Hematology, AIIMS, New Delhi

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