



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. : 5094

Distribution No.: 157-M

Month/Year: October/2022

Instrument ID: K11051903056

Name & Contact No. of PT Co-ordinator: Dr. Seema Tyagi (Prof.), Hematology, AIIMS, Delhi,
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Date of issue & status of the report: 24-11-2022[Final].

CBC and Retic Assessment

Test Parameters	S.No.	Among Lab (Accuracy Testing)						Within Lab (Precision Testing)			
		Your Result 1	Your Result 2	Your Results Sum of 2 Value	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 x10 ³ /µl	1	5.49	5.43	10.92	11.2	0.0290	-0.35	0.06	0.1	0.0060	-0.39
RBC x10 ⁶ /µl	1	3.85	3.82	7.67	7.55	0.0080	0.56	0.03	0.04	0.0030	-0.22
Hb g/dl	1	11.6	11.6	23.2	23.7	0.0270	-0.76	0	0.1	0.0080	-0.67
HCT%	1	39.3	39.1	78.4	73.3	0.1660	1.06	0.2	0.4	0.0250	-0.39
MCV-fl	1	102.4	102.1	204.5	194.6	0.3960	0.86	0.3	0.3	0.0210	0.00
MCH-Pg	1	30.5	30.2	60.7	62.6	0.0840	-0.80	0.3	0.3	0.0200	0.00
MCHC-g/dl	1	29.7	29.6	59.3	64.5	0.1500	-1.23	0.1	0.3	0.0220	-0.54
Plt. x10 ³ /µl	1	120	120	240	281	1.19	-1.29	0	4	0.28	-0.90
Retic %	2	8	6	14	10.5	0.23	0.53	2	0.5	0.03	2.53

P.S . Assesment

YOUR REPORT		CONSENSUS REPORT
DLC%	3	Nrbcs=2 , Poly=73 L=5, E=5, Mono/Promono=01 , B1=01 P.M.=01, Mye=9, Meta=5, Other=
RBC Morphology	3	Poly: 60 - 77, Myelo: 5 - 12, Meta: 5 - 10, Lympho: 3 - 7, Eos: 1- 3, nRBC/ Baso/ Promyelo, Blast Mono: 0 - 5
Diagnosis	3	microcytic hypochromic predominantly, tear drop cells seen occasionally
		Predominantly: Normocytic/Normochromic; Moderate: Anisocytosis, hypochromia
		Chronic Myeloproliferative Neoplasm favoring Chronic Myeloid Leukemia. however BCR-ABL1 translocation is needed to confirm the diagnosis.
		Chronic Myeloid Leukemia

COMBINED DATA VALUES OF TOTAL PARTICIPANTS

Test parameters	S.No.	Total participants covered in the current dist. 157--M	Total No. responded	% of Labs with Z Score 0-2		% of Labs with Z Score 2-3		% of Labs with Z Score >3	
				Among labs	Within lab	Among labs	Within lab	Among labs	Within lab
WBC x10³/µl	1	334	333	83.18	88.59	6.61	5.11	10.21	6.3
RBC x10⁶/µl	1	334	334	88.62	88.92	5.09	5.69	6.29	5.39
Hb g/dl	1	334	334	86.53	85.93	5.99	6.89	7.48	7.18
HCT%	1	334	332	93.98	91.57	4.22	3.31	1.8	5.12
MCV-fl	1	334	333	95.5	90.99	3	2.4	1.5	6.61
MCH-Pg	1	334	333	90.09	85.59	5.71	7.81	4.2	6.6
MCHC-g/dl	1	334	333	93.69	91.89	3.9	2.1	2.41	6.01
Plt. x10³/µl	1	334	333	91.29	91.89	5.71	4.2	3	3.91
ReticCount%	2	334	297	87.88	88.22	7.41	7.07	4.71	4.71
PS Assessment	3	334	270	Satisfactory :87.66%, Borderline Sat. :11.14%, Unsatisfactory :1.20%					

***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

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