



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. : 4043

Distribution No.: 159-k

Month/Year: July/2023

Instrument ID: ERBA H 360

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: 23-08-2023[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	5.4	5.3	10.7	10.88	0.101	-0.10	0.1	0.1	0.012	0.00
RBC x10 ⁶ /μl	1	4.55	4.53	9.08	8.86	0.014	0.93	0.02	0.04	0.004	-0.54
Hb g/dl	1	12.7	12.7	25.4	25.4	0.042	0.00	0	0.1	0.011	-1.35
HCT%	1	42.3	41.6	83.9	80.35	0.223	0.89	0.7	0.4	0.036	0.85
MCV-fl	1	93	91.8	184.8	182.5	0.391	0.33	1.2	0.3	0.037	1.80
MCH-Pg	1	28.1	27.9	56	57.35	0.104	-0.84	0.2	0.2	0.023	0.00
MCHC-g/dl	1	30.6	30	60.6	62.9	0.184	-0.61	0.6	0.3	0.022	1.35
Plt. x10 ³ /μl	1	208	203	411	418.5	2.672	-0.16	5	5	0.519	0.00
Retic %	2	4.6	4.2	8.8	10.4	0.396	-0.20	0.4	0.5	0.050	-0.17

P.S. Assesment

YOUR REPORT			CONSENSUS REPORT		
DLC%	3	Nrbcs=5 , Poly=17 L=22, E=, Mono/Promono=03 , B1=60 P.M.=, Mye=6, Meta=6, Other=	Blast: 26-64, Poly: 6-19, Lympho: 8-15, mono:2-20 , Myelo:0-7 , Meta: 0-7, promyelo: 0-6, Eosino:0-1		
RBC Morphology	3	NC/NC++,ANISO+	Predominantly: Normocytic/ Normochromic, Moderate: Anisocytosis, Microcytic		
Diagnosis	3	ACUTE LEUKEMIA	Acute Leukemia (AL)		

COMBINED DATA VALUES OF TOTAL PARTICIPANTS

Test parameters	S.No.	Total participants covered in the current dist. 156--I	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 x10 ³ /μl	1	138	138	87.68	83.33	5.8	2.9	6.52	13.77
RBC x10 ⁶ /μl	1	138	138	87.68	92.75	7.25	4.35	5.07	2.9
Hb g/dl	1	138	138	92.03	88.41	5.8	5.8	2.17	5.79
HCT%	1	138	138	89.86	90.58	7.25	5.8	2.89	3.62
MCV-fl	1	138	138	93.48	94.2	4.35	1.45	2.17	4.35
MCH-Pg	1	138	138	86.23	90.58	10.87	5.8	2.9	3.62
MCHC-g/dl	1	138	138	93.48	90.58	5.07	2.9	1.45	6.52
Plt. x10 ³ /μl	1	138	138	93.48	92.03	4.35	4.35	2.17	3.62
ReticCount%	2	138	115	96.52	89.57	0.00	6.96	3.48	3.47
PS Assessment	3	138	126	Satisfactory :91.31%, Borderline Sat. :0.72%, Unsatisfactory :7.97%					

***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 > ±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 > ±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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