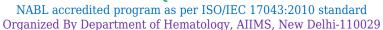




PROFICIENCY TESTING REPORT

ISHTM-AIIMS EXTERNAL QUALITY ASSURANCE PROGRAMME





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

EQAP CODE No.: 3104 **Distribution No.:** 149-H Month/Year: November/2019

Instrument ID: DXH 500 (BB020119)

Name & Contact No. of PT Co-ordinator: Dr. Renu Saxena (Prof & Head), Hematology, AIIMS, Delhi,

Tel: 9013085730, E-Mail: accuracy2000@gmail.com Date of issue & status of the report: 11-12-2019[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	15.93	15.93	31.86	11.25	0.1350	13.30	0	0.11	0.0210	-0.74	
RBC x10 ⁶ /μl	1	3.3	3.28	6.58	10.05	0.0210	-12.32	0.02	0.04	0.0060	-0.28	
Hb g/dl	1	12.11	12.1	24.21	25.4	0.0570	-1.78	0.01	0.1	0.0160	-0.61	
НСТ%	1	36.3	36	72.3	78.85	0.3710	-1.30	0.3	0.3	0.0560	0.00	
MCV-fl	1	110.1	109.7	219.8	156.85	0.6470	7.57	0.4	0.2	0.0410	0.67	
МСН-Рд	1	36.9	36.7	73.6	50.6	0.1190	16.33	0.2	0.2	0.0280	0.00	
MCHC-g/dl	1	33.6	33.3	66.9	64	0.3040	0.81	0.3	0.3	0.0440	0.00	
Plt. x10³/μl	1	451	418	869	735	5.55	1.93	33	9	1.15	1.96	
Retic %	2	0.7	0.6	1.3	6.5	0.39	-0.56	0.1	0.3	48.00	-0.30	

P.S. Assesment

		YOUR REPORT	CONSENSUS REPORT				
DLC%	3	Nrbcs=4 , Poly=11 L=04, E=02, Mono/Promono=01 , B1=24 P.M.=14, Mye=30, Meta=09, Other=01	Blast: 80-90, Poly: 5-7, Lymph: 2-7, Eo/Mono/Pro/My/Meta: 0-4				
RBC Morphology	3	$[\Lambda]$	Predominantly: Normocytic Normochromic. Moderate: Anisocytosis. Mild: Microcytic				
Diagnosis	3	Acute Myeloid Leukemia	Acute Leukemia (myeloid lineage)				

COMBINED DATA VALUES OF TOTAL PARTICIPANTS

Test	S.No.	Total participants covered in the current dist.	Total No. responded	% of Labs with Z Score 0-2		% of Labs with Z Score 2-3		% of Labs with Z Score >3	
parameters				Among labs	Within lab	Among labs	Within lab	Among labs	Within lab
WBC x10³/μl	1	100	82	80.49	87.8	10.98	4.88	12.2	10.98
RBC x10 ⁶ /μl	1	100	82	84.15	91.46	6.1	0	13.41	8.54
Hb g/dl	1	100	82	87.8	86.59	4.88	7.32	7.32	9.76
HCT%	1	100	82	90.24	89.02	3.66	0	6.1	13.41
MCV-fl	1	100	82	87.8	91.46	2.44	1.22	9.76	10.98
MCH-Pg	1	100	82	84.15	87.8	7.32	3.66	10.98	10.98
MCHC-g/dl	1	100	82	86.59	90.24	6.1	1.22	7.32	9.76
Plt. x10³/μl	1	100	82	87.8	89.02	4.88	6.1	10.98	8.54
ReticCount%	2	100	62	100	93.55	4.84	8.06	3.23	6.45
PS Assessment	3	100	68	Acceptable:93.5%, Warning Signal:3.9%, Unacceptable:2.6%					

*Comments:

- 1). Among Lab (EQA): CBC result for WBC, RBC, MCV & MCH unacceptable, please check calibration/human error.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.

Report authorized by,

Dr. R. Saxena

Prof & Head, Hematology, AIIMS, Delhi.

PT Co-ordinator: ISHTM-AIIMS-EQAP

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