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

NABL Certificate No. MC - 1002

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

EQAP CODE No. : 2921 **Distribution No.:** 152-G **Month/Year:** March/2021

Instrument ID: 2900pet02683

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: 12-05-2021[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	17.8	16.4	34.2	29.2	0.0870	2.76	1.4	0.2	0.0150	7.47
RBC x10^6/µl	1	4.64	4.5	9.14	9.12	0.0090	0.10	0.14	0.04	0.0020	2.70
Hb g/dl	1	12.2	11.9	24.1	23.4	0.0260	1.35	0.3	0.1	0.0080	2.70
HCT%	1	41	39.7	80.7	77.8	0.1760	0.73	1.3	0.4	0.0280	2.43
MCV-fl	1	88.4	88.3	176.7	171.3	0.3280	0.67	0.1	0.3	0.0250	-0.60
MCH-Pg	1	26.4	26.2	52.6	51.5	0.0620	0.76	0.2	0.2	0.0140	0.00
MCHC-g/dl	1	29.9	29.7	59.6	59.8	0.1410	-0.06	0.2	0.3	0.0200	-0.34
Plt. x10^3/µl	1	953	948	1901	1724	5.77	1.36	5	12	0.95	-0.56
Retic %	2	10.2	10.2	20.4	23.5	0.46	-0.27	0	0.5	0.05	-0.63

P.S . Assesment

		YOUR REPORT	CONSENSUS REPORT
DLC%	3	Nrbcs=00 , Poly=24 L=14, E=00, Mono/Promono=00 , B1=62 P.M.=00, Mye=00, Meta=00, Other=00	Blast: 20-80, Mono: 1-30, Poly: 10-25, Lympho: 5-15, Myelo/Promyelo/Meta: 1-5, nRBC/Eos: 0-1
RBC Morphology	3	mild anisocytosis & poikilocytosis. microcytosis.	Predominantly: Normocytic/Normochromic; Moderate: Microcytosis, Hypochromia; Mild: Anisocytosis, Macrocytosis
Diagnosis	3	ACUTE LEUKEMIA.	Acute Myeloid Leukemia (AML)

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COMBINED DATA VALUES OF TOTAL PARTICIPANTS

Test parameters	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	
				Among labs	Within lab	Among labs	Within lab	Among labs	Within lab
WBC $x10^3$/µl	1	214	233	87.12	90.56	3	3	9.44	5.58
RBC $x10^6$/µl	1	214	233	88.41	91.42	6.87	2.58	4.29	4.72
Hb g/dl	1	214	233	84.55	92.27	6.87	3	7.3	3.86
HCT%	1	214	233	93.13	88.41	5.58	5.58	0.86	5.15
MCV-fl	1	214	233	95.71	84.12	3.43	9.44	0	6.01
MCH-Pg	1	214	233	86.27	76.39	8.15	18.45	5.15	3.86
MCHC-g/dl	1	214	233	93.13	87.55	4.72	6.01	1.72	4.72
Plt. $x10^3$/µl	1	214	233	85.41	88.84	9.87	3.43	4.29	7.3
ReticCount%	2	214	214	96.73	92.99	2.34	5.14	0.93	2.34
PS Assessment	3	214	221	Acceptable:80.8,Warning Signal:10.3,Unacceptable :8.9					

*Comments:

1). Among Lab (EQA) : Results acceptable.

2). Within Lab (IQA) : Difference in the CBC measurement values for *WBC* unacceptable, may be due to random/human error.

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.

Report authorized by,

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