



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. : 2237

Distribution No.: 151-E

Month/Year: September 2020

Instrument ID: Mindray 5 part BC5100,RE-4A100300

Name & Contact No. of PT Co-ordinator: Dr. Seema Tyagi (Prof.), Hematology, AIIMS, Delhi,
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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	6.15	6.01	12.16	11.4	0.0740	0.40	0.14	0.1	0.0090	0.34
RBC x10 ⁶ /µl	1	4.84	4.74	9.58	9.61	0.0090	-0.11	0.1	0.04	0.0020	1.16
Hb g/dl	1	11.4	11.2	22.6	23.5	0.0220	-1.52	0.2	0.1	0.0080	0.67
HCT%	1	39.7	39.3	79	75.6	0.1710	0.76	0.4	0.3	0.0260	0.22
MCV-fl	1	82.8	82	164.8	157.25	0.2890	1.02	0.8	0.2	0.0190	2.02
MCH-Pg	1	23.7	23.6	47.3	48.9	0.0520	-1.35	0.1	0.2	0.0130	-0.45
MCHC-g/dl	1	28.7	28.6	57.3	62.1	0.1420	-1.30	0.1	0.2	0.0220	-0.34
Plt. x10 ³ /µl	1	193	186	379	340	1.66	0.92	7	6	0.42	0.15
Retic %	2	25	22	47	29.9	0.63	1.04	3	1	0.06	1.69

P.S . Assesment

YOUR REPORT			CONSENSUS REPORT		
DLC%	3	Nrbcs=NIL , Poly=05% L=05%, E=, Mono/Promono= , B1=50% P.M.=40%, Mye=, Meta=, Other=	Blasts: 65-85, Lymph: 2-6, nRBC/Poly/Eo/Mono: 0-5, Pro: 0-10, Myelo/Meta: 0-5.		
RBC Morphology	3	Are Normocytic, Normochromic with mild anisocytosis. Few macrocytes and microcytes cells are seen..	Predominantly: Normocytic, Normochromic. Moderate: Microcytic. Mild Anisocytosis.		
Diagnosis	3	Acute Myeloid Leukemia	Acute Leukemia (Myeloid Lineage)		

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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	450	272	86.4	75.74	6.25	7.72	4.04	9.19
RBC x10 ⁶ /µl	1	450	272	84.19	84.56	4.41	4.04	8.09	5.15
Hb g/dl	1	450	272	86.4	81.99	5.15	5.15	4.78	9.19
HCT%	1	450	272	89.71	83.46	5.51	4.41	1.47	6.25
MCV-fl	1	450	272	90.07	92.28	3.68	0.37	2.57	4.04
MCH-Pg	1	450	272	80.51	82.72	8.46	3.68	7.35	6.25
MCHC-g/dl	1	450	272	89.34	78.68	4.41	6.62	2.57	6.99
Plt. x10 ³ /µl	1	450	272	88.97	85.29	4.04	4.78	3.68	6.62
ReticCount%	2	450	254	96.06	93.31	2.36	2.76	2.76	5.12
PS Assessment	3	450	261	Acceptable:93.5%,Warning Signal:4.6%,Unacceptable :1.9%					

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.

Report authorized by,



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