



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. : 1910

Distribution No.: 159-F

Month/Year: March/2023

Instrument ID: BENESPHERA H31 (RQ-75122029)

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: 27-04-2023[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	8.2	7.9	16.1	15.33	0.039	0.75	0.3	0.1	0.008	1.80
RBC x10 ⁶ /µl	1	3.64	3.58	7.22	6.84	0.008	1.90	0.06	0.03	0.002	1.01
Hb g/dl	1	10.6	10.5	21.1	21.1	0.021	0.00	0.1	0.1	0.007	0.00
HCT%	1	37.5	37.3	74.8	67.1	0.167	1.61	0.2	0.4	0.023	-0.67
MCV-fl	1	90.4	89.4	179.8	195.3	0.416	-1.28	1	0.3	0.022	1.89
MCH-Pg	1	29.6	28.8	58.4	61.7	0.077	-1.82	0.8	0.3	0.018	1.69
MCHC-g/dl	1	32.8	32.3	65.1	63.1	0.160	0.43	0.5	0.3	0.021	0.67
Plt. x10 ³ /µl	1	170	168	338	458	1.916	-2.40	2	6	0.351	-0.67
Retic %	2	45.1	45	90.1	43	0.840	1.83	0.1	1	0.092	-0.81

P.S . Assesment

YOUR REPORT		CONSENSUS REPORT
DLC%	3	Nrbcs=10 , Poly=74 L=24, E=01, Mono/Promono=01 , B1=0 P.M.=0, Mye=0, Meta=0, Other=0
RBC Morphology	3	macrocytosis present anisopolkilocytosis marked, nucleated rbc -: 10/100wbc, platelet-: 82x1000/ul, mp absent, abnormal cell absent,
Diagnosis	3	Macrocytic anemia
		Thalassemia/Haemoglobinopathy

COMBINED DATA VALUES OF TOTAL PARTICIPANTS

Test parameters	S.No.	Total participants covered in the current dist. 159--F	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	317	315	84.76	82.22	2.54	10.79	12.7	6.99
RBC x10⁶/µl	1	317	317	83.91	88.01	8.52	5.68	7.57	6.31
Hb g/dl	1	317	317	86.44	92.11	5.68	3.79	7.88	4.1
HCT%	1	317	315	93.65	93.02	3.81	2.54	2.54	4.44
MCV-fl	1	317	315	94.29	91.75	5.08	3.49	0.63	4.76
MCH-Pg	1	317	315	85.71	88.89	6.67	4.76	7.62	6.35
MCHC-g/dl	1	317	315	95.24	91.43	3.49	4.76	1.27	3.81
Plt. x10³/µl	1	317	315	89.52	88.89	7.3	5.08	3.18	6.03
ReticCount%	2	317	282	98.23	87.94	1.42	3.55	0.35	8.51
PS Assessment	3	317	282	Satisfactory :89.28%, Borderline Sat. :6.94%, Unsatisfactory :3.78%					

***Comments:**

1). Among Lab (EQA) : PS Diagnosis partially correct, 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 > ±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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