



**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. : 2986

Distribution No.: 159-G

Month/Year: March/2023

Instrument ID: BENESPERA H33s TH-84004555

Name & Contact No. of PT Co-ordinator: Dr. Seema Tyagi (Prof.), Hematology, AIIMS, Delhi,  
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Date of issue &amp; status of the report: 01-05-2023[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 <sup>3</sup> /µl	1	4.5	4.5	9	8.21	0.046	0.71	0	0.1	0.007	-0.90
RBC x10 <sup>6</sup> /µl	1	4.13	4.13	8.26	8.8	0.010	-2.75	0	0.04	0.003	-1.08
Hb g/dl	1	13.1	13	26.1	25.9	0.024	0.32	0.1	0.1	0.008	0.00
HCT%	1	38.7	38.5	77.2	80.5	0.180	-0.74	0.2	0.3	0.027	-0.27
MCV-fl	1	93.8	93.2	187	183.65	0.290	0.43	0.6	0.3	0.025	0.81
MCH-Pg	1	31.7	31.6	63.3	59	0.073	2.52	0.1	0.2	0.017	-0.45
MCHC-g/dl	1	34.1	33.7	67.8	64.05	0.136	1.09	0.4	0.3	0.020	0.34
Plt. x10 <sup>3</sup> /µl	1	140	138	278	293	1.455	-0.40	2	4.5	0.344	-0.48
Retic %	2	14	12	26	8.15	0.186	3.81	2	0.4	0.026	6.17

### P.S . Assesment

YOUR REPORT		CONSENSUS REPORT
DLC%	3	Nrbcs=53 , Poly=56 L=41, E=2, Mono/Promono=1 , B1=0 P.M.=, Mye=, Meta=, Other=
RBC Morphology	3	Predominantly: Normocytic/Normochromic; Moderate: Anisocytosis, Microcytosis, Hypochromia; Mild: Poikilocytosis , Target cells , Sickle shaped cells ,tear drop cells
Diagnosis	3	Thalassemia/Haemoglobinopathy

**COMBINED DATA VALUES OF TOTAL PARTICIPANTS**

Test parameters	S.No.	Total participants covered in the current dist. 159--G	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
<b>WBC x10<sup>3</sup>/µl</b>	1	248	248	87.5	88.31	4.44	3.63	8.06	8.06
<b>RBC x10<sup>6</sup>/µl</b>	1	248	248	79.84	87.9	10.48	6.05	9.68	6.05
<b>Hb g/dl</b>	1	248	248	88.71	92.34	4.84	3.63	6.45	4.03
<b>HCT%</b>	1	248	248	94.35	89.92	4.44	4.84	1.21	5.24
<b>MCV-fl</b>	1	248	248	96.77	91.53	2.42	3.23	0.81	5.24
<b>MCH-Pg</b>	1	248	247	91.9	94.74	6.07	0.4	2.03	4.86
<b>MCHC-g/dl</b>	1	248	248	94.76	90.73	4.84	2.82	0.4	6.45
<b>Plt. x10<sup>3</sup>/µl</b>	1	248	248	92.74	93.95	5.24	4.03	2.02	2.02
<b>ReticCount%</b>	2	248	216	91.2	84.26	5.56	9.26	3.24	6.48
<b>PS Assessment</b>	3	248	218	Satisfactory :90.74%, Borderline Sat. :8.06%, Unsatisfactory :1.20%					

**\*Comments:**

1). **Among Lab (EQA) : RETIC result is unacceptable, may be due to random/human error.**

2). **Within Lab (IQA) : RETIC result is 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.ishtmaimseqap.com](http://www.ishtmaimseqap.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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