



**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. :** 3297

**Distribution No.:** 154-I

**Month/Year:** December/2021

**Instrument ID:** BC30(UD94000739)

**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:** 09-03-2022[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	6.23	6.14	12.37	13.1	0.0630	-0.90	0.09	0.1	0.0190	-0.06
RBC x10 <sup>6</sup> /µl	1	4.29	4.26	8.55	8.46	0.0150	0.45	0.03	0.05	0.0060	-0.39
Hb g/dl	1	13.2	13.2	26.4	26.35	0.0560	0.07	0	0.1	0.0150	-0.67
HCT%	1	43.7	42.7	86.4	81.6	0.3120	0.98	1	0.5	0.0660	0.84
MCV-fl	1	101.7	100.4	202.1	193.65	0.5570	1.02	1.3	0.3	0.0400	2.70
MCH-Pg	1	31	30.8	61.8	62.05	0.1270	-0.13	0.2	0.3	0.0320	-0.34
MCHC-g/dl	1	31	30	61	64.8	0.2700	-0.92	1	0.2	0.0340	2.70
Plt. x10 <sup>3</sup> /µl	1	149	141	290	244	2.15	1.35	8	6	0.59	0.45
Retic %	2			0							

### P.S . Assesment

YOUR REPORT			CONSENSUS REPORT		
<b>DLC%</b>	3	Nrbcs= , Poly= L=, E=, Mono/Promono= , B1= P.M.=, Mye=, Meta=, Other=	Blast: 60-85, Poly: 2-6, Lympho: 6-21, nRBC/mono/Eosino/Myelo/Meta/promyelo: 0-1		
<b>RBC Morphology</b>	3		Predominantly: Normocytic/ Normochromic, Moderate: Anisocytosis, Microcytic		
<b>Diagnosis</b>	3		Acute Leukemia (AL)		

**COMBINED DATA VALUES OF TOTAL PARTICIPANTS**

Test parameters	S.No.	Total participants covered in the current dist. 154--I	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	110	109	79.82	82.57	5.5	4.59	14.68	12.84
<b>RBC x10<sup>6</sup>/µl</b>	1	110	110	83.64	90	10	6.36	6.36	3.64
<b>Hb g/dl</b>	1	110	110	80.91	89.09	5.45	6.36	13.64	4.55
<b>HCT%</b>	1	110	110	93.64	90	4.55	5.45	1.81	4.55
<b>MCV-fl</b>	1	110	110	90	86.36	9.09	4.55	0.91	9.09
<b>MCH-Pg</b>	1	110	110	86.36	90.91	3.64	8.18	10	0.91
<b>MCHC-g/dl</b>	1	110	110	90	82.73	7.27	7.27	2.73	10
<b>Plt. x10<sup>3</sup>/µl</b>	1	110	109	92.66	87.16	5.5	9.17	1.84	3.67
<b>ReticCount%</b>	2	110	110	87.27	90	10	5.45	2.73	4.55
<b>PS Assessment</b>	3	110	104	Satisfactory :95.47%, Borderline Sat. :0.90%, Unsatisfactory :3.63%					

**\*Comments:**

**1). Among Lab (EQA) : Results acceptable. PS Diagnosis not reported**

**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  $\pm 2$ : Acceptable, Z score  $\pm 2$  to  $\pm 3$  :Warning Signal, Z score  $> \pm 3$  : Unacceptable [As per ISO/IEC 13528:2015 standard]

**Note-4:** Z score value between " $0$  to  $\pm 2$ " are texted in green colour. Z score value between " $\pm 2$  to  $\pm 3$ " 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](http://www.ishtmaiimseqap.com).

Report authorized by,



Dr. Seema Tyagi (Prof.)

PT Co-ordinator: ISHTM-AIIMS-EQAP

Department of Hematology, AIIMS, New Delhi

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