



**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. :** 3201

**Distribution No.:** 160-H    **Month/Year:** June/2023

**Instrument ID:** ASX MICROS 60 OT SERIAL NO  
8110T98394

**Name & Contact No. of PT Co-ordinator:** Dr. Manoranjan Mahapatra ( Prof. & Head), Hematology, AIIMS, Delhi,  
Tel: 9013085730 , E-Mail : accuracy2000@gmail.com

**Date of issue & status of the report:** 03-08-2023[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	6.4	6.2	12.6	10.79	0.061	1.91	0.2	0.1	0.010	0.84
RBC x10 <sup>6</sup> /µl	1	4.88	4.87	9.75	8.81	0.015	3.54	0.01	0.04	0.004	-0.67
Hb g/dl	1	14	14	28	25.5	0.044	3.15	0	0.1	0.012	-0.75
HCT%	1	43	42.9	85.9	80.95	0.241	1.19	0.1	0.5	0.043	-0.90
MCV-fl	1	88	88	176	183.6	0.427	-0.97	0	0.25	0.033	-0.67
MCH-Pg	1	28.8	28.7	57.5	57.8	0.092	-0.20	0.1	0.3	0.027	-0.67
MCHC-g/dl	1	32.7	32.7	65.4	62.45	0.178	0.90	0	0.3	0.035	-0.70
Plt. x10 <sup>3</sup> /µl	1	115	111	226	176.5	2.588	0.99	4	5	0.492	-0.17
Retic %	2	12	12	24	18.5	0.496	0.57	0	0.5	0.050	-0.84

### P.S . Assesment

YOUR REPORT		CONSENSUS REPORT
<b>DLC%</b>	3	Nrbcs= , Poly=12 L=76, E=06, Mono/Promono=02 , B1=04 P.M.=, Mye=, Meta=, Other=WBCs - Shift to left upto blast cells. Many smudge cells are also seen. Platelets- Adequate on smear. Blood parasites- Not seen
<b>RBC Morphology</b>	3	Predominantly: Normocytic/Normochromic; Moderate: Anisocytosis, Microcytosis, Hypochromia; Mild: Macrocytosis, Poikilocytosis.

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)
Diagnosis	3	Above hematological findings are suggestive of chronic lymphocytic leukemia. Advice to do cytochemistry and marker studies.				Chronic Lymphoproliferative Disorder/CLL			

### COMBINED DATA VALUES OF TOTAL PARTICIPANTS

Test parameters	S.No.	Total participants covered in the current dist. 160--H	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 <sup>3</sup> /µl	1	146	145	81.38	89.66	6.9	4.83	11.72	5.51
RBC x10 <sup>6</sup> /µl	1	146	146	90.41	92.47	2.05	4.11	7.54	3.42
Hb g/dl	1	146	146	86.99	86.3	6.16	5.48	6.85	8.22
HCT%	1	146	146	89.04	93.84	8.22	2.74	2.74	3.42
MCV-fl	1	146	146	91.78	93.15	6.85	2.74	1.37	4.11
MCH-Pg	1	146	146	81.51	89.04	9.59	6.85	8.9	4.11
MCHC-g/dl	1	146	146	91.1	93.84	6.85	4.11	2.05	2.05
Plt. x10 <sup>3</sup> /µl	1	146	146	91.1	91.78	6.85	4.79	2.05	3.43
ReticCount%	2	146	116	98.28	86.21	1.72	10.34	0	3.45
PS Assessment	3	146	109	Satisfactory :97.96%, Borderline Sat. :1.36%, Unsatisfactory :0.68%					

**\*Comments:**

1). **Among Lab (EQA) : CBC result for RBC & HB unacceptable, please check calibration/human error.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.ishtmaiinseqap.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

-----End Of Report-----



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