



**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. : 1834

Distribution No.: 155-F

Month/Year: March/2022

Instrument ID: ERBA ELITE-580 ( S.No- K11052120035)

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Date of issue &amp; status of the report: 29-04-2022[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	8.08	7.74	15.82	16.51	0.0810	-0.34	0.34	0.14	0.0110	1.35
RBC x10 <sup>6</sup> /µl	1	4.35	4.05	8.4	7.41	0.0340	0.97	0.3	0.06	0.0040	4.05
Hb g/dl	1	12.1	12	24.1	24	0.0220	0.17	0.1	0.1	0.0090	0.00
HCT%	1	39.1	37.4	76.5	70.05	0.1940	1.19	1.7	0.5	0.0260	2.31
MCV-fl	1	92.4	90	182.4	185.7	0.6860	-0.16	2.4	0.6	0.0430	2.43
MCH-Pg	1	29.6	27.8	57.4	64.4	0.2860	-0.80	1.8	0.4	0.0300	3.15
MCHC-g/dl	1	32	30.9	62.9	68.65	0.1770	-1.21	1.1	0.4	0.0250	1.89
Plt. x10 <sup>3</sup> /µl	1	539	518	1057	852	3.93	1.89	21	10	0.75	0.82
Retic %	2	3	2.5	5.5	14.55	0.24	-1.34	0.5	0.4	0.03	0.17

### P.S . Assesment

YOUR REPORT		CONSENSUS REPORT
DLC%	3	Nrbcs=2 , Poly=10 L=10, E=5, Mono/Promono=20 , B1=35 P.M.=10, Mye=5, Meta=5, Other=-
RBC Morphology	3	Blast: 45-80, Poly: 7-13, Lympho: 5-14, Promyelo: 0-6.25, Myelo/Mono/Meta: 1-5, nRBC/Eos: 0-1
Diagnosis	3	moderate anisopoikilocytosis predominantly microcytes with moderate to severe hypochromia. Acute Myeloid Leukemia (AML)

**COMBINED DATA VALUES OF TOTAL PARTICIPANTS**

Test parameters	S.No.	Total participants covered in the current dist. 155--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 x10 <sup>3</sup> /µl	1	314	307	86.32	85.67	6.19	6.19	7.49	8.14
RBC x10 <sup>6</sup> /µl	1	314	314	94.59	86.31	3.18	3.82	2.23	9.87
Hb g/dl	1	314	314	84.39	81.21	7.01	8.6	8.6	10.19
HCT%	1	314	309	89.32	88.67	5.83	4.85	4.85	6.48
MCV-fl	1	314	309	98.38	90.61	1.29	3.24	0.33	6.15
MCH-Pg	1	314	308	96.1	93.18	1.95	2.92	1.95	3.9
MCHC-g/dl	1	314	309	88.67	88.67	4.21	4.21	7.12	7.12
Plt. x10 <sup>3</sup> /µl	1	314	309	94.17	91.91	3.24	2.59	2.59	5.5
ReticCount%	2	314	314	86.94	82.17	3.18	6.05	9.88	11.78
PS Assessment	3	314	279	Satisfactory :86.27%, Borderline Sat. :2.55%, Unsatisfactory :11.18%					

**\*Comments:**

1). **Among Lab (EQA) : PS Diagnosis partially correct, remaining results acceptable**

2). **Within Lab (IQA) : Difference in the CBC measurement values for RBC & MCH unacceptable, please check precision/human error. Remaining 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](http://www.ishtmaiimseqap.com).

Report authorized by,



Dr. Seema Tyagi (Prof.)

PT Co-ordinator: ISHTM-AIIMS-EQAP

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