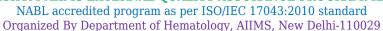




PROFICIENCY TESTING REPORT

ISHTM-AIIMS EXTERNAL QUALITY ASSURANCE PROGRAMME





Duration of stability testing - minimum upto 8 days at ambient temp. after dispatch of specimens

EQAP CODE No.: 5215 **Distribution No.**: 158-M **Month/Year**: January/2023

Instrument ID: BC-3000plus (RJ8C127453)

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:** 28-02-2023[Final].

CBC and Retic Assessment

				Amo	ng Lab (Aco	curacy Testin	Within Lab (Precision Testing)				
Test Parameters	S.No.	Your Result 1		Your Results Sum of 2 Value	Consensus result sum of 2 values (Assigned Value)	Uncertainty		Results	Consensus Result Diff. of 2 values (Assigned Value)	Uncertainty of Assigned Values	Z Score
WBC x10³/μl	1	7	6.5	13.5	13.31	0.035	0.22	0.5	0.1	0.007	4.15
RBC x10 ⁶ /μl	1	5.17	4.97	10.14	10.25	0.012	-0.30	0.2	0.05	0.003	2.89
Hb g/dl	1	12.6	12.2	24.8	26.5	0.027	-2.29	0.4	0.1	0.008	2.02
НСТ%	1	37.8	36. <mark>5</mark>	74.3	85.7	0.219	-1.53	1.3	0.4	0.025	2.31
MCV-fl	1	73.5	73.2	146.7	169.35	0.353	-1.97	0.3	0.3	0.019	0.00
MCH-Pg	1	24.5	24.3	48.8	51.5	0.059	-1.66	0.2	0.2	0.014	0.00
MCHC-g/dl	1	33.4	33.3	66.7	61	0.140	1.05	0.1	0.3	0.021	-0.67
Plt. x10³/μl	1	355	326	681	781	2.986	-1.11	29	9	0.518	2.45
Retic %	2	2.5	2	4.5	15.35	0.222	-1.82	0.5	0.5	0.034	0.00

P.S. Assesment

		YOUR REPORT	CONSENSUS REPORT
DLC%	3		Blast: 65-88, Lympho: 5-14, Poly: 2-5, nRBC/Eos/Baso/Mono /Myelo/Meta/ Promyelo: 0-5

				Among Lab (Accuracy Testin			ng)	With	in Lab (Pre	ecision Testi	ng)
Test Parameters		Your Result 1		Ioui	Consensus result sum of 2 values (Assigned Value)	Uncertainty of Assigned Values	Z Score	Results	Consensus Result Diff. of 2 values (Assigned Value)	Uncertainty of Assigned Values	Z Score
RBC Morphology	3	Normocytic Normochromic RBC's predominantly with occational microcytic hypochromic RBC's .Nucleated RBC's noted				Predominantly: Normocytic/ Normochromic, Moderate: Anisocytosis, Microcytic					
Diagnosis	3	Normocytic normochromic anemia.Acute leukemia (accure myeloid leukemia) thrombocytophenia				Acute Leukemia	a (AL)				

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COMBINED DATA VALUES OF TOTAL PARTICIPANTS

Test parameters	S No	Total participants covered in the	Total No.	% of Labs with Z Score 0-2		% of Labs with Z Score 2-3		% of Labs with Z Score >3	
rest parameters	3.NU.	current dist. 158M	responded	Among labs	Within lab	Among labs	Within lab	Among labs	Within lab
WBC x10³/μl	1	338	335	83.88	84.18	6.27	6.27	9.85	9.55
RBC x10 ⁶ /μl	1	338	338	89.64	89.35	5.92	5.03	4.44	5.62
Hb g/dl	1	338	338	88.76	85.21	5.03	4.44	6.21	10.35
HCT%	1	338	336	97.92	89.88	0.89	5.36	1.19	4.76
MCV-fl	1	338	336	97.62	87.2	1.79	6.55	0.59	6.25
MCH-Pg	1	338	336	88.39	89.58	7.44	4.76	4.17	5.66
MCHC-g/dl	1	338	336	98.21	86.9	0.89	7.74	0.9	5.36
Plt. x10³/μl	1	338	336	94.64	92.26	3.27	2.98	2.09	4.76
ReticCount%	2	338	296	91.22	85.14	6.08	9.8	2.7	5.06
PS Assessment	3	338	283	Satisfactory	:97.93%, Bo	rderline Sat	:1.18%, Uı	nsatisfactory	:0.890%

*Comments:

- 1). Among Lab (EQA): Results acceptable.
- 2). Within Lab (IQA): Difference in the CBC measurement values for WBC 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 $> \pm 3$: 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 $> \pm 3$ 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.

Report authorized by,

Dr. Seema Tyagi (Prof.)

PT Co-ordinator: ISHTM-AIIMS-EQAP

Department of Hematology, AIIMS, New Delhi

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

ISHBT





PROFICIENCY TESTING REPORT

ISHTM-AIIMS EXTERNAL QUALITY ASSURANCE PROGRAMME
NABL accredited program as per ISO/IEC 17043:2010 standard

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.: 5215 **Distribution No.:** 157-M **Month/Year:** October/2022

Instrument ID: BC-3000plus (RJ8C127453)

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:** 24-11-2022[Final].

CBC and Retic Assessment

				Amo	ng Lab (Aco	curacy Testin	Within Lab (Precision Testing)				
Test Parameters	S.No.	Your Result 1		Your Results Sum of 2 Value	Consensus result sum of 2 values (Assigned Value)	Uncertainty		Results		Uncertainty of Assigned Values	Z Score
WBC x10³/μl	1	5.4	5.2	10.6	11.2	0.029	-0.74	0.2	0.1	0.006	0.96
RBC x10 ⁶ /μl	1	3.73	3.73	7.46	7.55	0.008	-0.42	0	0.04	0.003	-0.90
Hb g/dl	1	10.9	10.8	21.7	23.7	0.027	-3.03	0.1	0.1	0.008	0.00
НСТ%	1	31	30.9	61.9	73.3	0.166	-2.37	0.1	0.4	0.025	-0.58
MCV-fl	1	83.3	82.9	166.2	194.6	0.396	-2.46	0.4	0.3	0.021	0.34
МСН-Рд	1	29.2	28.9	58.1	62.6	0.084	-1.90	0.3	0.3	0.020	0.00
MCHC-g/dl	1	35.1	34.9	70	64.5	0.150	1.30	0.2	0.3	0.022	-0.27
Plt. x10³/μl	1	143	129	272	281	1.191	-0.28	14	4	0.275	2.25
Retic %	2	1.7	1	2.7	10.5	0.232	-1.18	0.7	0.5	0.033	0.34

P.S. Assesment

		YOUR REPORT	CONSENSUS REPORT
DLC%	2		Poly: 60 - 77, Myelo: 5 - 12, Meta: 5 - 10, Lympho: 3 - 7, Eos: 1- 3, nRBC/ Baso/ Promyelo, Blast Mono: 0 - 5
RBC Morphology		NORMOCYTIC IN NORMOCHROMIC ,RBCs WITH OCCASIONAL TEAR DROP CELLS AND FEW PENCIL CELLS .NRBCs SEEN	Predominantly: Normocytic/Normochromic; Moderate: Anisocytosis, hypochromia
Diagnosis	3	CHRONIC MYELOID LEUKEMIA	Chronic Myeloid Leukemia

COMBINED DATA VALUES OF TOTAL PARTICIPANTS

Test parameters	S.No.	Total participants	Total No.	% of Labs with Z Score 0-2		% of Labs with Z Score 2-3		% of Labs with Z Score >3	
Test parameters		current dist. 157M	responded	Among labs	Within lab	Among labs	Within lab	Among labs	Within lab
WBC x10³/μl	1	334	333	83.18	88.59	6.61	5.11	10.21	6.3
RBC x10 ⁶ /μl	1	334	334	88.62	88.92	5.09	5.69	6.29	5.39
Hb g/dl	1	334	334	86.53	85.93	5.99	6.89	7.48	7.18
HCT%	1	334	3 <mark>32</mark>	93.98	91.57	4.22	3.31	1.8	5.12
MCV-fl	1	334	333	95.5	90.99	3	2.4	1.5	6.61
MCH-Pg	1	334	333	90.09	<mark>85</mark> .59	5.71	7.81	4.2	6.6
MCHC-g/dl	1	334	333	93.69	91.89	3.9	2.1	2.41	6.01
Plt. x10³/μl	1	334	333	91.29	91.89	5.71	4.2	3	3.91
ReticCount%	2	334	297	87.88	88.22	7.41	7.07	4.71	4.71
PS Assessment	3	334	270	Satisfactory	:87.66%, Bo	rderline Sat	.:11.14%, U	Insatisfactor	y:1.20%

*Comments:

- 1). Among Lab (EQA): CBC result for HB unacceptable, may be due to random/human error
- 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)

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IQR = Quartile 3 - Quartile 1 of participant data, Normalised IQR = 0.7413 x IQR

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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 $> \pm 3$ 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.

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-----