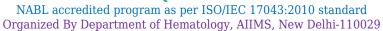




PROFICIENCY TESTING REPORT

ISHTM-AIIMS EXTERNAL QUALITY ASSURANCE PROGRAMME





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

EQAP CODE No.: 1198 Distribution No.: 153-D Month/Year: August/2021 Instrument ID: MEK 7300K, Celltac ES, 5 Part Automated Hematology, Analyser, Nihon Kohden, Japan. Serial

No.994

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: 13-09-2021[Final].

CBC and Retic Assessment

				Among Lab (Accuracy Testing)				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 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 x10³/μl	1	2.9	2.9	5.8	5.5	0.0230	0.51	0	0.1	0.0050	-1.69	
RBC x10 ⁶ /μl	1	3.54	3.53	7.07	7.54	0.0070	-2.54	0.01	0.04	0.0020	-0.81	
Hb g/dl	1	11.7	11.7	23.4	23.8	0.0200	-0. 77	0	0.1	0.0070	-1.35	
НСТ%	1	36.8	36. <mark>6</mark>	73.4	74.3	0.1290	-0.25	0.2	0.4	0.0230	-0.54	
MCV-fl	1	104	103	207	196.55	0.2900	1.19	1	0.4	0.0250	1.62	
MCH-Pg	1	33.1	33.1	66.2	63.1	0.0300	2.02	0	0.3	0.0160	-1.01	
MCHC-g/dl	1	32	31.8	63.8	64.15	0.1170	-0.11	0.2	0.3	0.0170	-0.45	
Plt. x10³/μl	1	117	117	234	222	1.09	0.38	0	5	0.28	-1.12	
Retic %	2	0.7	0.5	1.2	1.2	0.03	0.00	0.2	0.2	0.01	0.00	

P.S. Assesment

		YOUR REPORT	CONSENSUS REPORT				
DLC%	3		Poly: 40 - 60, Myelo: 10 - 25, Meta: 5 - 20, Promyelo: 1-10, nRBC/Lympho/Blast/Eos/Baso/Mono: 0 - 5				
RBC Morphology	3	Anisocytosis +, normocytic normochromic +, macrocytosis +, hypochromia +	Predominantly: Normocytic/Normochromic; Moderate: Anisocytosis, hypochromia, Microcytosis; Mild: Macrocytosis, Poikilocytosis				
Diagnosis		Chronic myeloproferative neoplasm, CML is the likely possibility	Chronic Myeloid Leukemia (Chronic Phase)				

EQAP Code Distribution No.: No.: 1198 153-D

Month/Year: August/2021

Instrument ID: MEK 7300K, Celltac ES, 5 Part Automated Hematology, Analyser, Nihon Kohden, Iapan.Serial No.994

COMBINED DATA VALUES OF TOTAL PARTICIPANTS

Test parameters	S.No.	Total participants covered in the current dist.	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 x10³/μl	1	317	314	84.71	87.9	3.5	1.59	11.78	9.87
RBC x10 ⁶ /μl	1	317	314	92.68	91.08	4.14	3.82	3.18	4.78
Hb g/dl	1	317	314	88.22	91.72	6.69	0.32	5.1	4.46
HCT%	1	317	314	95.22	90.13	2.23	3.5	2.55	6.05
MCV-fl	1	317	314	96.5	92.68	2.23	3.18	1.27	4.14
MCH-Pg	1	317	314	88.85	91.72	7.96	3.5	3.18	4.78
MCHC-g/dl	1	317	314	94.59	88.22	3.5	3.18	1.91	8.6
Plt. x10³/μl	1	317	314	93.31	90.45	4.46	4.14	2.23	5.41
ReticCount%	2	317	276	92.39	89.49	5.07	6.88	3.99	5.8
PS Assessment	3	317	295	Acceptable:98.65%, Warning Signal:1.35%, Unacceptable:0%					

*Comments:

1). Among Lab (EQA): 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 $> \pm 3$: 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 > ± 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-----