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

NABL Certificate No. PC - 1002

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

EQAP CODE No. : 1196 **Distribution No.:** 152-D **Month/Year:** February/2021

Instrument ID: SYSMEX XN 350 (11359)

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: 02-03-2021[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 x10³/µl	1	2.43	2.37	4.8	5.5	0.0200	-1.23	0.06	0.1	0.0050	-0.62
RBC x10⁶/µl	1	4.66	4.66	9.32	9.32	0.0080	0.00	0	0.03	0.0020	-1.01
Hb g/dl	1	13.4	13.4	26.8	27	0.0200	-0.34	0	0.1	0.0070	-1.35
HCT%	1	40.6	40.6	81.2	83.9	0.1780	-0.46	0	0.3	0.0200	-0.81
MCV-fl	1	87.1	87.1	174.2	179.9	0.3170	-0.55	0	0.2	0.0170	-0.90
MCH-Pg	1	28.8	28.8	57.6	57.7	0.0540	-0.07	0	0.2	0.0140	-0.90
MCHC-g/dl	1	33	33	66	64.15	0.1300	0.41	0	0.3	0.0120	-1.01
Plt. x10³/µl	1	96	94	190	196	0.75	-0.28	2	4	0.26	-0.45
Retic %	2	2.7	2.5	5.2	5	0.08	0.09	0.2	0.2	0.01	0.00

P.S . Assesment

		YOUR REPORT	CONSENSUS REPORT
DLC%	3	Nrbcs=NORMOBLASTS 60% , Poly=68 L=18, E=1, Mono/Promono=5 , B1= P.M.=01, Mye=04, Meta=03, Other=	nRBC: 30 - 65, Poly: 60 - 75, Lympho: 15-30, Eos/Mono: 1-5, Blast/Myelo/Meta: 0-1
RBC Morphology	3	ANISOPOIKILOCYTOSIS,SPHEROCYTES,MICROCYTOSIS,MACROCYTOSIS,TARGET & TEAR DROP CELLS,S HOWELL JOLLY BODIESPHEROCYTES	Predominantly: Macrocytosis, Microcytosis, Spherocytosis, Polychromasia, Anisocytosis: Moderate: Normocytic/Normochromic, Hypo.
Diagnosis	3	HEREDITARY SPHEROCYTOSIS	Hemolytic Anemia

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 $x10^3/\mu l$	1	312	347	89.05	91.93	2.31	1.44	8.36	6.34
RBC $x10^6/\mu l$	1	312	347	89.63	90.78	6.92	3.46	3.17	5.19
Hb g/dl	1	312	347	91.07	93.08	6.92	2.59	2.02	4.32
HCT%	1	312	347	97.41	91.93	1.73	3.46	0.58	4.32
MCV-fl	1	312	347	97.12	85.59	1.44	8.93	0.86	4.9
MCH-Pg	1	312	347	91.35	90.78	6.05	3.46	2.31	5.48
MCHC-g/dl	1	312	347	98.27	91.93	0.29	3.75	1.15	3.75
Plt. $x10^3/\mu l$	1	312	347	93.08	91.64	3.46	5.48	3.17	2.59
ReticCount%	2	312	318	93.71	86.48	4.09	2.2	2.2	11.64
PS Assessment	3	312	335	Acceptable:91.4,Warning Signal:7.7,Unacceptable :0.9					

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 > ±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----------

ISHTM-AIIMS EXTERNAL QUALITY ASSURANCE PROGRAMME
An ISO/IEC 17043:2010 certified programme
Organized by
Department of Hematology, AIIMS, New Delhi-110029

Email: accuracy2000@gmail.com

NABL Certificate No. PC-1002

PARTICIPATION CERTIFICATE

[Certificate No. EQAP/1196/2021/04]

Dated: 16.4.2021

This is to certify that, Vinamra Swaraj Hospital, Navi Mumbai-400703has participated in the "ISHTM-AIIMS External Quality Assurance Program" for the period "Jan 2020 to Dec 2020".

Dr. Seema Tyagi (Prof, Hematology)
Chief Coordinator
ISHTM-AIIMS-EQAP