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. : 1937 **Distribution No.:** 155-E **Month/Year:** March/2022

Instrument ID: H360-K10012102151

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: 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 $x10^3/\mu l$	1	3.1	2.9	6	6.4	0.0320	**-0.45**	0.2	0.09	0.0050	**1.65**
RBC $x10^6/\mu l$	1	2.96	2.94	5.9	6.14	0.0070	**-1.30**	0.02	0.03	0.0020	**-0.34**
Hb g/dl	1	10.9	10.8	21.7	22.4	0.0210	**-1.28**	0.1	0.1	0.0070	**0.00**
HCT%	1	33.6	33.3	66.9	69.4	0.1290	**-0.71**	0.3	0.3	0.0230	**0.00**
MCV-fl	1	113.5	113.3	226.8	225.1	0.3410	**0.18**	0.2	0.4	0.0310	**-0.34**
MCH-Pg	1	37.1	36.5	73.6	73.1	0.0970	**0.19**	0.6	0.3	0.0220	**1.01**
MCHC-g/dl	1	32.7	32.2	64.9	64.65	0.1280	**0.07**	0.5	0.3	0.0150	**0.67**
Plt. $x10^3/\mu l$	1	185	170	355	351	1.21	**0.12**	15	5	0.31	**1.93**
Retic %	2										

P.S . Assesment

YOUR REPORT			CONSENSUS REPORT
DLC%	3	Nrbcs=2 , Poly=69 L=4, E=1, Mono/Promono=05 , B1=00 P.M.=02, Mye=08, Meta=09, Other=-	Poly: 37 - 50, Myelo: 16 - 32, Meta: 8 - 16, Promyelo: 1-10, nRBC/Lympho/Blast/Eos/Baso/Mono: 0 - 5
RBC Morphology	3	HYPOCHRONIC AND NORMOCYTIC TO MACROCYTIC.FEW MICROCYTES.ANISOCYTOSIS	Predominantly: Normocytic/Normochromic; Moderate: Anisocytosis, hypochromia, Microcytosis; Mild: Macrocytosis, Poikilocytosis
Diagnosis	3	Suggestive of chronic myeloproliferative disorder.Immunophenotyping advised for confirmation	Chronic Myeloid Leukemia (Chronic Phase)

Result satisfactory

Anju Kackar

COMBINED DATA VALUES OF TOTAL PARTICIPANTS

Test parameters	S.No.	Total participants covered in the current dist. 155--E	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	320	318	89.94	90.25	5.66	3.77	4.4	5.98
RBC $x10^6/\mu l$	1	320	320	88.13	87.81	6.56	3.13	5.31	9.06
Hb g/dl	1	320	320	85	91.25	5.63	3.13	9.37	5.62
HCT%	1	320	318	93.4	88.36	5.03	5.97	1.57	5.67
MCV-fl	1	320	317	94.01	95.58	4.42	1.58	1.57	2.84
MCH-Pg	1	320	317	90.22	88.01	4.1	5.36	5.68	6.63
MCHC-g/dl	1	320	318	93.08	90.25	4.09	4.72	2.83	5.03
Plt. $x10^3/\mu l$	1	320	318	89.94	88.99	6.92	5.66	3.14	5.35
ReticCount%	2	320	320	84.69	82.19	5.63	5.94	9.68	11.87
PS Assessment	3	320	300	Satisfactory :87.16%, Borderline Sat. :6.89%, Unsatisfactory :5.95%					

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----------

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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. : 1937 **Distribution No.:** 154-E **Month/Year:** November/2021

Instrument ID: H360

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: 23-02-2022[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 $x10^3/\mu l$	1	3.51	3.4	6.91	10.99	0.1670	**-0.68**	0.11	0.18	0.0140	-0.31
RBC $x10^6/\mu l$	1	4.24	4.2	8.44	8.53	0.0100	**-0.36**	0.04	0.06	0.0040	-0.34
Hb g/dl	1	13	12.9	25.9	25.9	0.0300	**0.00**	0.1	0.1	0.0100	0.00
HCT%	1	41.2	41.2	82.4	79.3	0.1750	**0.66**	0	0.4	0.0290	-0.77
MCV-fl	1	98	97.3	195.3	185.2	0.3170	**1.12**	0.7	0.3	0.0250	0.90
MCH-Pg	1	30.8	30.8	61.6	60.5	0.0690	**0.69**	0	0.3	0.0250	-0.70
MCHC-g/dl	1	31.6	31.4	63	64.8	0.1380	**-0.48**	0.2	0.4	0.0270	-0.39
Plt. $x10^3/\mu l$	1	122	96	218	278	1.30	**-1.76**	26	5	0.34	3.54
Retic %	2	0.8	0.7	1.5	8.8	0.20	**-1.17**	0.1	0.36	0.02	-0.64

P.S . Assesment

YOUR REPORT			CONSENSUS REPORT
DLC%	3	Nrbcs=1.00 , Poly=4.00 L=5.00, E=0.00, Mono/Promono=1.00 , B1=80.00 P.M.=9.00, Mye=, Meta=, Other=	Blast: 60-88, Poly: 2-6, Lympho: 4-12, Mono: 0-4, Myelo/Promyelo/Meta: 1-5, nRBC/Eos: 0-1
RBC Morphology	3	HYPOCHROMIC AND MICROCYTIC .ANISONUCLEOSIS .FEW MACROCYTES.TEAR DROP CELLS	Predominantly: Normocytic/Normochromic; Moderate: Microcytosis, Hypochromia; Mild: Anisocytosis, Macrocytosis
Diagnosis	3	AML.IMMUNOPHENOTYPING ADVISED FOR CONFIRMATION	Acute Myeloid Leukemia (AML)

Arjit Kesh (?) Randomemr (?) Result satisfactory Platelet ILC done satisfactory

COMBINED DATA VALUES OF TOTAL PARTICIPANTS

rameters	S.No.	Total participants covered in the current dist. 154--E	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
BC x10^3/µl	1	298	290	96.55	88.62	3.79	4.14	-0.34	7.24
RBC x10^6/µl	1	298	298	83.89	86.91	6.38	5.37	9.73	7.72
Hb g/dl	1	298	298	81.88	83.22	10.4	4.36	7.72	12.42
HCT%	1	298	291	93.81	86.6	3.44	7.9	2.75	5.5
MCV-fl	1	298	291	92.78	94.85	5.15	2.75	2.07	2.4
MCH-Pg	1	298	291	83.51	91.75	8.59	6.19	7.9	2.06
MCHC-g/dl	1	298	291	91.75	94.16	6.19	3.44	2.06	2.4
Plt. x10^3/µl	1	298	291	93.13	94.85	3.78	3.78	3.09	1.37
ReticCount%	2	298	298	95.3	88.59	3.02	1.68	1.68	9.73
PS Assessment	3	298	275	Satisfactory :78.18%, Borderline Sat. :21.14%, Unsatisfactory :0.67%					

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.
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Report authorized by,

Dr. Seema Tyagi (Prof.)
PT Co-ordinator: ISHTM-AIIMS-EQAP
Department of Hematology, AIIMS, New Delhi

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

Healthplus Wellness DIAGNOSTICS

ISO 9001 2015

Contact No.: 011-42441274, 8130018328
Email : healthplusxx@gmail.com

Date	13/07/2022		Srl No.	1	
Name	Ms. MEGHA		Age	23 Yrs.	Sex F
Ref. By	Dr. SELF				

COMPLETE HAEMOGRAM

HAEMOGLOBIN (Hb)	13.7	gm/dl	11.0 - 16.0
TOTAL LEUCOCYTE COUNT (TLC)	5,800	/cumm	4000 - 11000
DIFFERENTIAL LEUCOCYTE COUNT (DLC)			
NEUTROPHIL	51	%	40 - 75
LYMPHOCYTE	40	%	20 - 45
EOSINOPHIL	04	%	01 - 06
MONOCYTE	05	%	02 - 10
BASOPHIL	00	%	0 - 0
ESR (WESTEGREN`s METHOD) Westegren`s	16	mm/Ist hr.	0 - 20
R B C COUNT	4.13	Millions/cmm	3.8 - 4.8
P.C.V / HAEMATOCRIT	42.0	%	35 - 45
M C V	**101.7**	fl.	80 - 100
M C H	**33.1**	Picogram	27.0 - 31.0
M C H C	**32.6**	gm/dl	33 - 37
PLATELET COUNT	**174**	Lakh/cmm	1.50 - 4.50
RDW	**54.4**	FL	39.0 - 46.0

Anju Kackar

DR. ANJU KACKAR
MBBS,MD
SENIOR PATHOLOGIST

In case of any discrepancies in the report, please contact the laboratory immediately.
(This is professional opinion and not the final diagnosis. It should be clinically correlated)

Timings 8 am to 8 pm
Sundays 8 am to 1.30 pm

• Free Home Collection • Not For Medico Legal Cases

Pocket C2/34, Sector 11, Rohini, Delhi-110085

Patient MRN:	296160	Patient ID:	2712207130001
Name:	**Ms. MEGHA**	Sample Drawn Date:	13/Jul/2022 04:22PM
Age/Gender:	23 Y/Female	Lab Accession Date:	13/Jul/2022 04:22PM
Order ID:	296160130722130921	Report Date & Time:	13/Jul/2022 05:18PM
Booked By:	Healthplus Wellness Diagnostics	Ref By:	Self
Sample Type:		BarcodeID/Slide No:	2665685/

MC-2606

Test Name	Results	Units	Bio. Ref. Interval	Test Method
Complete Blood Count (CBC)				
Hemoglobin^	13.5	g/dL	12.0-15.0	Non Cyanide - SLS
Total Leucocyte Count (TLC / WBC)^	5.75	10^3/uL	4.0-10.0	Flocytometry
Packed Cell Volume (PCV / HCT)^	42.1	%	36.0-46.0	Cumulative Pulse Height Detection
Mean Corpuscular Volume (MCV)^	98.1	fl	83.0-101.0	Calculated
Mean Corpuscular Hemoglobin (MCH)^	31.5	pg	27.0-32.0	Calculated
Mean Copuscular Hb Conc (MCHC)^	32.1	g/dL	31.5-34.5	Calculated
Platelet count^	161	10^3/uL	150-410	DC Detection
RDW-SD^	**47.3**	fL	36.4-46.3	Calculated
RDW-CV^	13.6	%	11.7-14.4	Calculated
Neutrophils^	68.00	%	40-80	Semiconductor Laser Flocytometry/ Light Microscopy
Lymphocytes^	25.00	%	20-40	Semiconductor Laser Flocytometry/ Light Microscopy
Monocytes^	5.00	%	2-10	Semiconductor Laser Flocytometry/ Light Microscopy
Eosinophils^	2.00	%	1-6	Semiconductor Laser Flocytometry/ Light Microscopy
Basophils^	0.00	%	0-2	Semiconductor Laser Flocytometry/ Light Microscopy
Absolute Neutrophils^	3.91	10^3/uL	2.00-8.00	Calculated
Absolute Lymphocytes^	1.44	10^3/uL	1.00-3.00	Calculated
Absolute Monocytes^	0.29	10^3/uL	0.20-1.00	Calculated
Absolute Eosinophils^	0.12	10^3/uL	0.02-0.50	Calculated
Absolute Basophils^	**0.00**	10^3/uL	0.02-0.10	Calculated

Above Results are of the Tests performed at NirAmaya Pathlabs with Commitment to provide Accurate Pathology Services

Approved By Dr. Ashok Malhotra

Dr. Indu Sardana
MD Pathology
Lab director & Senior Pathologist

Dr. Ashok Malhotra
MBBS, MD.
Sr. Consultant Biochemist

Dr. Surbhi
MBBS. MD. Microbiologist

Test result marked 'BOLD/RED' indicates abnormal results i.e higher or lower than normal.
All Lab results are subject to clinical interpretation by a qualified medical professional & This report is not subject to use for any medico-legal purpose