



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.: 853

Distribution No.: 164-A

Month/Year: May/2024

Instrument ID: SYSMEX

Model Name.: XN-330

Serial No.: 15454

Name & Contact No. of PT Co-ordinator: Dr. Manoranjan Mahapatra (Prof. & Head), Hematology, AIIMS, Delhi,

Tel: 9013085730, E-Mail: info@ishtmaiimseqap.com

Date of issue & status of the report: 05-07-2024[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 | | Results | Consensus Result Diff. of 2 values (Assigned Value) | Uncertainty of Assigned Values | | |
| WBC x10³/μl | 1 | 5.28 | 5.18 | 10.46 | 9.84 | 0.044 | 0.49 | 0.1 | 0.1 | 0.006 | 0.00 | |
| RBC x10 ⁶ /μl | 1 | 4.42 | 4.39 | 8.81 | 9.06 | 0.008 | -1.12 | 0.03 | 0.03 | 0.002 | 0.00 | |
| Hb g/dl | 1 | 14.3 | 14.3 | 28.6 | 28.6 | 0.026 | 0.00 | 0 | 0.1 | 0.007 | -1.35 | |
| НСТ% | 1 | 43.5 | 43.1 | 86.6 | 89.7 | 0.215 | -0.49 | 0.4 | 0.4 | 0.023 | 0.00 | |
| MCV-fl | 1 | 98.4 | 98.2 | 196.6 | 197.8 | 0.419 | -0.09 | 0.2 | 0.2 | 0.019 | 0.00 | |
| МСН-Рд | 1 | 32.6 | 32.4 | 65 | 63 | 0.056 | 1.28 | 0.2 | 0.2 | 0.011 | 0.00 | |
| MCHC-g/dl | 1 | 33.2 | 32.9 | 66.1 | 63.3 | 0.150 | 0.60 | 0.3 | 0.3 | 0.016 | 0.00 | |
| Plt. x10³/μl | 1 | 117 | 113 | 230 | 231 | 1.429 | -0.02 | 4 | 3 | 0.222 | 0.22 | |
| Retic % | 2 | 10.8 | 10.3 | 21.1 | 31 | 0.457 | -0.70 | 0.5 | 0.8 | 0.044 | -0.58 | |

P.S. Assesment

| YOUR REPORT | | | CONSENSUS REPORT | | | | |
|-------------------|---|------------------------------------------------------------------------------------------|----------------------------------------------------------------------------------------------------------------------------|--|--|--|--|
| DLC% | 3 | Nrbcs=8 , Poly=22 L=3, E=4, Mono/Promono=0 , B1=3 P.M.=25, Mye=22, Meta=18, Other= | Poly: 20 - 35, Lympho: 4 - 10, Myelo: 19 - 39, Meta: 7-15, Promyelo: 2-9, Eosino: 2-7, Mono: 1-2, Blast: 1 - 5, Baso: 0-5 | | | | |
| RBC Morphology | 3 | | Predominantly: Normocytic/Normochromic; Moderate: Anisocytosis, Mild: Polychromatophils (+), Poikilocytosis, Macrocytes | | | | |
| Diagnosis | 3 | CHRONIC MYELOID LEUKEMIA | MPN likely CML-CP | | | | |

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COMBINED DATA VALUES OF TOTAL PARTICIPANTS

| Test parameters | C N- | Total participants covered in the current dist. 164A | Total No. responded | % of Labs with Z Score 0-2 | | % of Labs with Z Score 2-3 | | % of Labs with Z Score >3 | | |
|---------------------------|-------|------------------------------------------------------|------------------------|---------------------------------------------------------------------|---------------|-------------------------------|---------------|------------------------------|---------------|--|
| rest parameters | 5.No. | | | Among labs | Within lab | Among labs | Within lab | Among labs | Within lab | |
| WBC x10 ³ /μl | 1 | 365 | 365 | 81.64 | 92.6 | 7.67 | 2.19 | 10.69 | 5.21 | |
| RBC x10 ⁶ /μl | 1 | 365 | 365 | 89.32 | 91.51 | 6.03 | 3.84 | 4.65 | 4.65 | |
| Hb g/dl | 1 | 365 | 365 | 86.58 | 89.04 | 7.4 | 3.56 | 6.02 | 7.4 | |
| НСТ% | 1 | 365 | 365 | 93.7 | 90.41 | 4.66 | 3.84 | 1.64 | 5.75 | |
| MCV-fl | 1 | 365 | 365 | 94.79 | 93.15 | 3.01 | 3.84 | 2.2 | 3.01 | |
| MCH-Pg | 1 | 365 | 365 | 85.75 | 93.7 | 7.67 | 2.74 | 6.58 | 3.56 | |
| MCHC-g/dl | 1 | 365 | 365 | 93.7 | 90.96 | 3.56 | 3.01 | 2.74 | 6.03 | |
| Plt. x10 ³ /μl | 1 | 365 | 365 | 89.59 | 89.86 | 6.58 | 5.21 | 3.83 | 4.93 | |
| ReticCount% | 2 | 365 | 350 | 96.86 | 92.29 | 2.57 | 0.86 | 0.57 | 6.85 | |
| PS Assessment | 3 | 365 | 345 | Satisfactory :89.05%, Borderline Sat. :2.46%, Unsatisfactory :8.49% | | | | | | |

'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 $> \pm 3$ 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 $(\overline{x}-\overline{y})$ 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. Manoranjan Mahapatra (Prof. & Head)

PT Co-ordinator: ISHTM-AIIMS-EOAP

Department of Hematology, AIIMS, New Delhi

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

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NABL accredited program as per ISO/IEC 17043:2010 standard
Organized By Department of Hematology, AIIMS, New Delhi-110029



PARTICIPATION CERTIFICATE

[Certificate No. EQAP/853/2023/30]

Date 30.12.2023

This is to certify that" AGILUS PATH LABS-DR.PHADKE LABS - DR.TECKCHANDANI LAB., THANE, MAHARASHTRA, 400706 "has participated in the "ISHTM-AIIMS External Quality Assurance Program" for the period "January 2023 to December 2023".

Report authorized by,

Dr. Manoranjan Mahapatra (Prof. & Head)

PT Co-ordinator: ISHTM-AIIMS-EQAP

Department of Hematology, AIIMS, New Delhi



Date: 10/07/2024

TO WHOMSOEVER IT MAY CONCERN

I Dr Suresh Teckchandani, Technical Head of Agilus Pathlabs Pvt. Ltd. Dr. Teckchandani Lab, Vashi, New Mumbai, confirm that the lab located at Office-12, 1st Floor, Above Golden Punjab Hotel, Sector-17, Vashi, Navi Mumbai is a unit of Agilus Pathlabs Pvt. Ltd.- Dr. Avinash Phadke labs.

For Agilus Pathlabs Pvt. Ltd, Dr. Teckchandani, Vashi Lab.

Dr. Suresh Teckchandani

Lab Head