



PROFICIENCY TESTING REPORT
 ISHBT-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. : 2661
 Instrument ID: SYSMEX

Distribution No.: 164-E
 Model Name.: XP-100

Month/Year: June/2024
 Serial No.: A9675

Name & Contact No. of PT Co-ordinator: Dr. Manoranjan Mahapatra (Prof. & Head), Hematology, AIIMS, Delhi,
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Date of issue & status of the report: 05-08-2024[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 | 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 /µl | 1 | 5.3 | 5.1 | 10.4 | 28.25 | 0.639 | -0.83 | 0.2 | 0.3 | 0.021 | -0.25 |
| RBC x10 ⁶ /µl | 1 | 4.98 | 4.95 | 9.93 | 9.7 | 0.011 | 0.78 | 0.03 | 0.04 | 0.003 | -0.19 |
| Hb g/dl | 1 | 12.8 | 12.7 | 25.5 | 27.6 | 0.029 | -2.60 | 0.1 | 0.1 | 0.008 | 0.00 |
| HCT% | 1 | 40.1 | 39.7 | 79.8 | 86.85 | 0.235 | -1.03 | 0.4 | 0.4 | 0.025 | 0.00 |
| MCV-fl | 1 | 80.5 | 80.2 | 160.7 | 178.8 | 0.448 | -1.54 | 0.3 | 0.3 | 0.023 | 0.00 |
| MCH-Pg | 1 | 25.9 | 25.5 | 51.4 | 56.9 | 0.082 | -2.56 | 0.4 | 0.3 | 0.013 | 0.34 |
| MCHC-g/dl | 1 | 32.2 | 31.7 | 63.9 | 63.5 | 0.181 | 0.08 | 0.5 | 0.3 | 0.022 | 0.54 |
| Plt. x10 ³ /µl | 1 | 129 | 122 | 251 | 175 | 2.475 | 1.04 | 7 | 5 | 0.342 | 0.34 |
| Retic % | 2 | 0.91 | 0.68 | 1.59 | 21.6 | 0.244 | -2.90 | 0.23 | 0.7 | 0.047 | -0.49 |

P.S . Assesment

| YOUR REPORT | | | CONSENSUS REPORT | | |
|----------------|---|--|---|--|--|
| DLC% | 3 | Nrbcs=16 , Poly=39 L=14, E=4, Mono/Promono=8 , B1=14 P.M.=1, Mye=1, Meta=3, Other= | Poly: 53-64, Lympho: 28-38, Eosino: 1-3, Mono: 2-5, blast/Promyelo/Myelo/Meta: 0-5 | | |
| RBC Morphology | 3 | MODERATE POIKILOCYTOSIS.CELLS ARE MICROCYTIC AND MACROCYTIC WITH TEAR DROP CELLS,TARGET CELLS. | Predominantly: Microcytic, Hypochromic, Moderate: Anisopoikilocytosis Mild:Target cells , Tear drop cells | | |
| Diagnosis | 3 | ACUTE MYELOCYTIC LEUMIA. | Thalassemia Hemoglobinopathy | | |

COMBINED DATA VALUES OF TOTAL PARTICIPANTS

| Test parameters | S.No. | Total participants covered in the current dist. 164--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 /μl | 1 | 329 | 325 | 84 | 85.85 | 3.69 | 6.46 | 12.31 | 7.69 |
| RBC x10 ⁶ /μl | 1 | 329 | 329 | 87.54 | 92.1 | 7.9 | 3.95 | 4.56 | 3.95 |
| Hb g/dl | 1 | 329 | 329 | 88.15 | 82.37 | 5.47 | 8.81 | 6.38 | 8.82 |
| HCT% | 1 | 329 | 326 | 95.09 | 90.18 | 2.45 | 5.21 | 2.46 | 4.61 |
| MCV-fI | 1 | 329 | 325 | 91.38 | 94.46 | 5.54 | 2.15 | 3.08 | 3.39 |
| MCH-Pg | 1 | 329 | 325 | 88.31 | 93.23 | 5.54 | 4 | 6.15 | 2.77 |
| MCHC-g/dl | 1 | 329 | 325 | 92.92 | 92.62 | 4.31 | 2.77 | 2.77 | 4.61 |
| Plt. x10 ³ /μl | 1 | 329 | 327 | 92.97 | 88.38 | 3.06 | 5.2 | 3.97 | 6.42 |
| ReticCount% | 2 | 329 | 289 | 87.89 | 95.5 | 10.03 | 2.42 | 2.08 | 2.08 |
| PS Assessment | 3 | 329 | 290 | Satisfactory :95.75%, Borderline Sat. :1.82%, Unsatisfactory :2.43% | | | | | |

Comments:

1). Among Lab (EQA) : **PS Diagnosis wrongly reported, remaining 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.

Note 10: Reports are kept confidential.

Report authorized by,



Dr. Manoranjan Mahapatra (Prof. & Head)

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