



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. : 3529

Distribution No.: 162-J

Month/Year: January/2024

Instrument ID: HORIBA

Model Name.: YUMIZEN H550

Serial No.: 909YAXH02625

Name & Contact No. of PT Co-ordinator: Dr. Manoranjan Mahapatra (Prof. & Head), Hematology, AIIMS, Delhi,
 Tel: 9013085730 , E-Mail : accuracy2000@gmail.com

Date of issue & status of the report: 21-03-2024[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 | 1.69 | 1.53 | 3.22 | 10.41 | 0.030 | -10.21 | 0.16 | 0.1 | 0.006 | 0.62 |
| RBC x10 ⁶ /μl | 1 | 4.88 | 4.79 | 9.67 | 9.51 | 0.011 | 0.55 | 0.09 | 0.05 | 0.003 | 0.90 |
| Hb g/dl | 1 | 12.7 | 12.7 | 25.4 | 24.79 | 0.022 | 1.19 | 0 | 0.1 | 0.008 | -0.71 |
| HCT% | 1 | 39.2 | 38.2 | 77.4 | 78.9 | 0.207 | -0.24 | 1 | 0.4 | 0.028 | 1.35 |
| MCV-fl | 1 | 80.2 | 79.9 | 160.1 | 168.4 | 0.394 | -0.67 | 0.3 | 0.3 | 0.021 | 0.00 |
| MCH-Pg | 1 | 26.4 | 26 | 52.4 | 51.9 | 0.065 | 0.30 | 0.4 | 0.3 | 0.015 | 0.45 |
| MCHC-g/dl | 1 | 33.1 | 32.4 | 65.5 | 62 | 0.166 | 0.69 | 0.7 | 0.3 | 0.020 | 1.66 |
| Plt. x10 ³ /μl | 1 | 205 | 198 | 403 | 421 | 2.119 | -0.36 | 7 | 7 | 0.417 | 0.00 |
| Retic % | 2 | 18 | 14 | 32 | 22.55 | 0.336 | 1.06 | 4 | 0.7 | 0.057 | 2.78 |

P.S . Assesment

| YOUR REPORT | | CONSENSUS REPORT |
|-----------------------|---|--|
| DLC% | 3 | Nrbcs=1 , Poly=76 L=05, E=2, Mono/Promono=01 , B1=02 P.M.=05, Mye=02, Meta=05, Other=Nil |
| RBC Morphology | 3 | NORMOCYTIC NORMOCHROMIC RBCs ADMIXED WITH FEW MICROCYTIC MILD TO MODERATE HYPOCHROMIC RBCs |
| Diagnosis | 3 | Differential Diagnosis-1) CML- CHRONIC PHASE or 2) Myeloid Leukemoid Reaction |

COMBINED DATA VALUES OF TOTAL PARTICIPANTS

| Test parameters | S.No. | Total participants covered in the current dist. 162--J | 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 | 303 | 300 | 81 | 93.33 | 4.67 | 1.67 | 14.33 | 5 |
| RBC x10⁶/µl | 1 | 303 | 303 | 89.11 | 89.11 | 4.95 | 4.95 | 5.94 | 5.94 |
| Hb g/dl | 1 | 303 | 303 | 84.49 | 88.45 | 6.93 | 4.62 | 8.58 | 6.93 |
| HCT% | 1 | 303 | 300 | 97 | 91 | 1.67 | 3.67 | 1.33 | 5.33 |
| MCV-fl | 1 | 303 | 300 | 97.67 | 90.33 | 1 | 3.33 | 1.33 | 6.34 |
| MCH-Pg | 1 | 303 | 300 | 89.33 | 78 | 6.67 | 15.33 | 4 | 6.67 |
| MCHC-g/dl | 1 | 303 | 300 | 96.67 | 91.67 | 2.33 | 4.33 | 1 | 4 |
| Plt. x10³/µl | 1 | 303 | 300 | 88.67 | 86.67 | 9.33 | 5.67 | 2 | 7.66 |
| ReticCount% | 2 | 303 | 260 | 95 | 93.08 | 3.85 | 1.92 | 1.15 | 5.00 |
| PS Assessment | 3 | 303 | 252 | Satisfactory :80.01%, Borderline Sat. :10.66%, Unsatisfactory :9.33% | | | | | |

***Comments:**

- 1). Among Lab (EQA) : CBC result for WBC unacceptable, may be due to random/human error**
- 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)

PT Co-ordinator: ISHTM-AIIMS-EQAP

Department of Hematology, AIIMS, New Delhi

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