



**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. : 4042

Distribution No.: 152-K

Month/Year: March/2021

Instrument ID: Dymind DH 36, 3 Part Differential Automated Hematology Analyser, Equip S.N. - DM10011813072

Name &amp; Contact No. of PT Co-ordinator: Dr. Seema Tyagi (Prof.), Hematology, AIIMS, Delhi,

Tel: 9013085730, E-Mail : accuracy2000@gmail.com

Date of issue &amp; status of the report: 24-05-2021[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 x10 <sup>3</sup> /µl	1	5.18	5.05	10.23	15.4	0.3590	-0.71	0.13	0.13	0.0180	0.00	
RBC x10 <sup>9</sup> /µl	1	4.38	4.33	8.71	8.28	0.0130	2.00	0.05	0.05	0.0040	0.00	
Hb g/dl	1	11.6	11.5	23.1	23.9	0.0420	-1.08	0.1	0.1	0.0110	0.00	
HCT%	1	38	37.6	75.6	74.3	0.3300	0.22	0.4	0.4	0.0370	0.00	
MCV-fl	1	86.8	86.7	173.5	178.7	0.5920	-0.51	0.1	0.2	0.0310	-0.27	
MCH-Pg	1	26.6	26.4	53	57.8	0.1200	-2.31	0.2	0.2	0.0260	0.00	
MCHC-g/dl	1	30.6	30.5	61.1	63.7	0.2690	-0.54	0.1	0.2	0.0280	-0.34	
Plt. x10 <sup>3</sup> /µl	1	198	195	393	360	3.51	0.57	3	9	0.85	-0.67	
Retic %	2	5	4.5	9.5	5	0.15	1.53	0.5	0.3	0.10	0.90	

### P.S . Assesment

YOUR REPORT			CONSENSUS REPORT	
DLC%	3	Nrbcs=00, Poly=13 L=02, E=00, Mono/Promono=00, B1=62 P.M.=08, Mye=10, Meta=05, Other=00	Poly: 30 - 65, Myelo: 10 - 35, Meta: 5 - 20, Promyelo/Blast/Lympho: 1 - 10, nRBC/Baso/Eos/Mono: 0 - 5	
RBC Morphology	3	Normocytic, Hypochromic	Predominantly: Normocytic/Normochromic; Moderate: Anisocytosis, Hypochromia, Microcytosis; Mild: Poikilocytosis, Macrocytosis	
Diagnosis	3	Chronic MyeloblastidLeukemia ( CML )	Chronic Myeloid Leukemia (CML)	

EQAP Code Distribution No.: Month/Year:  
No.: 4042 152-K March/2021

Instrument ID: Dymind DH 36, 3 Part Differential  
Automated Hematology Analyser, Equip S.N. -  
DM10011813072

### 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 <sup>3</sup> /µl	1	124	136	86.03	86.03	8.09	8.09	5.88	5.88
RBC x10 <sup>6</sup> /µl	1	124	136	88.24	91.91	5.15	3.68	6.62	4.41
Hb g/dl	1	124	136	90.44	253.68	5.88	8.09	3.68	1.47
HCT%	1	124	136	93.38	91.18	5.15	2.94	1.47	5.88
MCV-fl	1	124	136	94.12	87.5	4.41	3.68	1.47	8.82
MCH-Pg	1	124	136	91.91	91.18	3.68	2.94	4.41	5.88
MCHC-g/dl	1	124	136	94.12	86.76	4.41	5.15	1.47	8.09
Plt. x10 <sup>3</sup> /µl	1	124	135	90.37	87.41	5.93	5.19	3.7	7.41
ReticCount%	2	124	122	93.44	85.25	3.28	0.82	3.28	13.93
PS Assessment	3	124	124	Acceptable:92,Warning Signal:4,Unacceptable :4					

#### 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.ishtmaimseqap.com.

Report authorized by,



Dr. Seema Tyagi (Prof.)

PT Co-ordinator: ISHTM-AIIMS-EQAP

Department of Hematology, AIIMS, New Delhi

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Reviewed &  
found satisfactory  
30/5/2021



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

**EQAP CODE No. :** 4042

**Distribution No.:** 153-K

**Month/Year:** October/2021

**Instrument ID:** DYMIND DH 36, S.N. DM10011813072

**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:** 10-12-2021[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 x10 <sup>3</sup> /μl	1	3.97	3.85	7.82	8.7	0.0440	-0.91	0.12	0.13	0.0100	-0.08
RBC x10 <sup>6</sup> /μl	1	5.61	5.46	11.07	10.4	0.0130	2.08	0.15	0.05	0.0040	1.69
Hb g/dl	1	15.3	15.3	30.6	32.5	0.0460	-1.83	0	0.2	0.0100	-2.70
HCT%	1	53.9	52.5	106.4	99.6	0.2350	1.05	1.4	0.5	0.0390	1.62
MCV-fI	1	96.1	96	192.1	190.7	0.3630	0.15	0.1	0.4	0.0290	-0.51
MCH-Pg	1	28	27.3	55.3	62.4	0.0880	-1.42	0.7	0.3	0.0190	1.80
MCHC-g/dl	1	29.2	28.4	57.6	65.35	0.1640	-1.84	0.8	0.3	0.0210	1.69
Plt. x10 <sup>3</sup> /μl	1	142	138	280	284	1.84	-0.08	4	6	0.36	-0.39
Retic %	2	6	5.5	11.5	7	0.14	1.24	0.5	0.3	0.02	0.90

**P.S. Assesment**

YOUR REPORT			CONSENSUS REPORT
<b>DLC%</b>	3	Nrbcs=0, Poly=4 L=96, E=0, Mono/Promono=0, B1=0 P.M.=0, Mye=0, Meta=0, Other=0	Lymph: 85-94, Poly: 4-12, blast: 1-8, nRBC/mono/Eosino/Myelo/Meta: 0-1
<b>RBC Morphology</b>	3	Predominantly Microcytic, Hypochromic, Few normocytic, normochromic and moderate Anisocytosis	Predominantly: Normocytic/Normochromic; Moderate: Anisocytosis, Microcytosis, Hypochromia; Mild: Macrocytosis, Poikilocytosis
<b>Diagnosis</b>	3	Chronic Lymphoproliferative Disorder - probably Chronic Lymphocytic Leukemia	Chronic Lymphocytic Leukemia (CLL)

## COMBINED DATA VALUES OF TOTAL PARTICIPANTS

Test parameters	S.No.	Total participants covered in the current dist. 153--K	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 <sup>3</sup> /μl	1	267	267	77.9	88.01	6.74	2.25	15.36	9.74
RBC x10 <sup>6</sup> /μl	1	267	267	91.01	86.52	6.37	7.49	2.62	5.99
Hb g/dl	1	267	267	88.39	61.42	7.87	32.21	3.74	6.37
HCT%	1	267	266	95.86	87.97	2.26	5.26	1.88	6.77
MCV-fl	1	267	266	94.36	95.49	4.89	2.26	0.75	2.25
MCH-Pg	1	267	266	88.35	89.47	7.14	3.38	4.51	7.15
MCHC-g/dl	1	267	266	95.49	89.85	3.38	4.14	1.13	6.01
Plt. x10 <sup>3</sup> /μl	1	267	266	95.11	90.98	3.38	4.89	1.51	4.13
ReticCount%	2	267	267	95.13	84.27	2.62	1.5	2.25	14.23
PS Assessment	3	267	232	Satisfactory :93.14%, Borderline Sat. :0.86%, Unsatisfactory :6%					

**\*Comments:**

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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-----

Reviewed &  
found Satisfactory  
Amul  
15/12/2021