



PT EQAS RESULT REVIEW FORM

Issued by QA:

Department: Haematology	Month / Year: MAY-2023
PT Provider: METROPOLIS MHL EQAS	Distribution / Lot No: 230104/03
Analyte(Test): TLL, RBC Count, Haemoglobin, MCV, MCH, MCHC, RBC Count, HCT, RDW C%.	
Machine used: ADVIA 560 (SIEMENS)	Kit used: ADVIA 560 (SIEMENS)
Test done by: RHIKA	Done on: 15/5/2023
Report uploaded by: AMET KUMAR	Report uploaded on: 19/5/2023
Software used: METROPOLIS MHL EQAS	NA
EQAS Report received on 19/5/2023	NA
Reviewed by: AMET KUMAR / KOLDDP	Approved by: DR. SHWETA ANUJA
Observations: ALL EQAS RESULT IS Satisfactory with 88.89% score EXCEPT TLL RESULT, due to SI unit issue After SI unit correction TLL Result value is within acceptable Range.	

\*If results are ungraded, please attach self-assessment. If satisfactory no corrective and preventive action required. But if unsatisfactory, corrective and preventive action is required with optimum evidence.

Self-Assessment Result:  Satisfactory  Unsatisfactory

Attachment:  Yes  No

Please investigate failure for unsatisfactory performance and submit a report to QA and Medical Director in 3 Days.

Approved by DR. SHWETA ANUJA

Signature  
(Lab Director/Designee)



Analyte	Instrument	Result Value	Standard Unit	Z-Score
* Total Leucocytes (WBC) count	ADVIA 560	1.50	/c.mm	*
Legend @ : z score $\leq$ 2.0 - Acceptable ! : 2.0 < z score < 3.0 - Warning X : z score $\geq$ 3 - Unacceptable # : Not Evaluated ⊙ : Delayed Result Entry * : Not considered for evaluation.				

*unit 21/05/2023*

**Problem Classification:** *TLC SI unit issue. TLC value wrongly filled as per SI during EQAS RESULT submitted. i.e 10<sup>3</sup>/cu is filled. in EQAS SI unit is 1/cu.mm.*

**Corrective Action:** *follow the same in next cycle.*

**Reviewed by:** *[Signature]*  
 21/5/2023

**Dated:** 21/5/2023.

*[Signature]*



Instrument : ADVIA 560

Analyte	Standard Unit	Result Value	Mean	Z-Score	RMZ
@ Erythrocyte (RBC) Count	mill/cu.mm	1.03	1.04	-0.04	-0.61
* Total Leucocytes (WBC) count	/c.mm	1.50	1280.00	-13.99 *	0.00
@ Haemoglobin	g/dL	3.10	2.84	1.73	1.16
@ MCV (Mean Corpuscular Volume)	fL	116.70	112.06	0.54	0.30
@ MCH (Mean Corpuscular Hb)	pg	29.70	27.37	1.42	1.29
@ MCHC (Mean Corpuscular Hb Concentration)	g/dL	25.40	24.39	1.16	1.12
@ Platelet Count	1000/ $\mu$ L	27.00	30.00	-0.70	-0.81
@ Hematocrit	%	12.00	11.70	0.58	-0.06
@ RDW CV%	%	25.50	24.53	1.43	0.66

Legend @ : Acceptable

! :  $2.0 < z \text{ score} < 3.0$  - Warning

X :  $z \text{ score} \geq 3$  - Unacceptable

# : Not Evaluated

⊙ : Delayed Result Entry

\* : Not considered for evaluation.

Total Parameters	9
Not Evaluated Parameters	0
Evaluated Parameters	9
Outlier Parameters (X)	1
<b>EQAS Score Haematology - CBC</b>	<b>88.89 %</b>



*Dr Puneet Kumar Nigam*

Dr Puneet Kumar Nigam  
PT coordinator & Technical Manager, MHL EQAS  
Unit No. 409-416.  
Commercial Building - 1A  
Kohinoor Mall, Kiroi Road, Kurla (W),  
Mumbai - 400070



## PT EQAS RESULT REVIEW FORM

Issued by QA:

Department: <u>Hematology</u>	Month / Year: <u>MAY-2023</u>
PT Provider: <u>METROPOLIS MHL EQAS</u>	Distribution / Lot No: <u>230104/04</u>
Analyte(Test): <u>TLC, Rbc count, Haemoglobin, MW, MCV, MCHC, Rf Count, HCT, Rbc CV.</u>	
Machine used: <u>ADVIA 560 (SIEMENS)</u>	Kit used: <u>ADVIA 560 (SIEMENS)</u>
Test done by: <u>19/5/2023 RIHKA</u>	Done on: <u>19/5/2023</u>
Report uploaded by: <u>19/5/2023 AMET KUMAR</u>	Report uploaded on: <u>19/5/2023</u>
Software used: <u>METROPOLIS MHL EQAS</u>	
EQAS Report received on <u>19/5/2023</u>	
Reviewed by: <u>AMET KUMAR/KULDEEP</u>	Approved by: <u>DR. SHWETA AHUJA</u>
Observations: <u>ALL EQAS RESULT IS Satisfactory with 88.89% Score EXCEPT TLC RESULT, due to SI UNIT issue After SI UNIT correction TLC RESULT value is within Acceptable Range.</u>	

\*If results are ungraded, please attach self-assessment. If satisfactory no corrective and preventive action required. But if unsatisfactory, corrective and preventive action is required with optimum evidence.

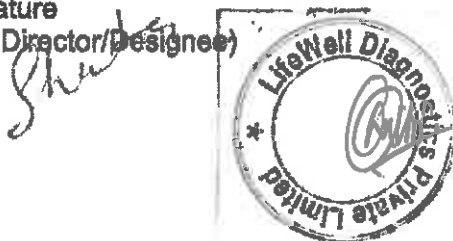
Self-Assessment Result:  Satisfactory  Unsatisfactory

Attachment:  Yes  No

Please investigate failure for unsatisfactory performance and submit a report to QA and Medical Director in 3 Days.

Approved by DR. SHWETA AHUJA

Signature  
(Lab Director/Designee)



Analyte	Instrument	Result Value	Standard Unit	Z-Score
* Total Leucocytes (WBC) count	ADVIA 560	8.48	/c.mm	.
Legend @ : z score $\leq$ 2.0 - Acceptable ! : 2.0 < z score < 3.0 - Warning X : z score $\geq$ 3 - Unacceptable # : Not Evaluated ⊕ : Delayed Result Entry * : Not considered for evaluation.				

Problem Classification: TLC SI unit issue. TLC value <sup>units of 10<sup>3</sup>/cmm</sup> wrong filled  
as per SI during EQAS result submitted  
i.e 10<sup>3</sup>/cmm is filled in EQAS SI unit is  
1/cmm

Corrective Action: Follow the same in next cycle

Reviewed by: Amit  
21/5/2023

Dated: 21/5/2023





**Metropolis Healthcare Ltd., EQAS**  
**LIFEWELL DIAGNOSTICS PVT LTD (CHANDIGARH) (880)**  
**Outlier And Analyte Summary Report**  
**Result Details For Cycle No 230104 and Sample No 04**

Report Date : 19/05/2023

Instrument : ADVIA 560

Analyte	Standard Unit	Result Value	Mean	Z-Score	RMZ
@ Erythrocyte (RBC) Count	mill/cu.mm	2.15	2.25	-1.17	-0.61
• Total Leucocytes (WBC) count	/c.mm	8.48	9720.00	-10.24 *	0.00
@ Haemoglobin	g/dL	7.70	7.78	0.59	1.16
@ MCV (Mean Corpuscular Volume)	fL	131.50	132.10	0.06	0.30
@ MCH (Mean Corpuscular Hb)	pg	35.90	34.61	1.15	1.29
@ MCHC (Mean Corpuscular Hb Concentration)	g/dL	27.30	26.20	1.07	1.12
@ Platelet Count	1000/ µL	211.00	224.00	-0.92	-0.81
@ Hematocrit	%	28.30	29.88	-0.70	-0.06
@ RDW CV%	%	16.90	19.10	-0.11	0.66

Legend @ : Acceptable

: 2.0 < z score < 3.0 - Warning

X : z score ≥ 3 - Unacceptable

# : Not Evaluated

⊙ : Delayed Result Entry

\* : Not considered for evaluation.

Total Parameters	9
Not Evaluated Parameters	0
Evaluated Parameters	9
Outlier Parameters (X)	1
EQAS Score Haematology - CBC	88.89 %



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 Commercial Building - 1A  
 Kohinoor Mall, Kirod Road, Kurla (W),  
 Mumbai - 400070



PT EQAS RESULT REVIEW FORM

Issued by QA:

Department: <i>Haematology</i>	Month / Year: <i>MAY- 2023</i>
PT Provider: <i>METROPOLIS MHL EQAS -</i>	Distribution / Lot No: <i>23010401</i>
Analyte(Test): <i>Neutrophil, Lymphocyte, Eosinophil.</i>	
Machine used: <i>MICROSCOPIC</i>	Kit used: <i>MICROSCOPIC</i>
Test done by: <i>MICROSCOPY by DR. SHWETA</i>	Done on: <i>12/5/2023</i>
Report uploaded by: <i>AMIT KUMAR</i>	Report uploaded on: <i>14/5/2023</i>
Software used: <i>METROPOLIS MHL EQAS</i>	
EQAS Report received on <i>19/5/2023</i>	
Reviewed by: <i>DR. SHWETA AHUJA</i>	Approved by: <i>DR. SHWETA AHUJA</i>
Observations: <i>ALL BEAS RESULT WITHIN ACCEPTABLE RANGE. WITH 100.0% SCORE. BEAS SATISFACTORY</i>	

\*If results are ungraded, please attach self-assessment. If satisfactory no corrective and preventive action required. But if unsatisfactory, corrective and preventive action is required with optimum evidence.

Self-Assessment Result:  Satisfactory  Unsatisfactory

Attachment  Yes  No

Please investigate failure for unsatisfactory performance and submit a report to QA and Medical Director in 3 Days.

Approved by *DR. SHWETA AHUJA*

Signature  
(Lab Director/Designee)





**Motropolis Healthcare Ltd., EQAS**  
LIFEWELL DIAGNOSTICS PVT LTD (CHANDIGARH) (880)  
Outlier And Analyte Summary Report

Outlier Details For Cycle No 230104 and Sample No 01

Report Date : 19/05/2023

Analyte	Instrument	Result Value	Standard Unit
No Outlier			
Legend @ : Acceptable X : Unacceptable # : Not Evaluated ⊙ : Delayed Result Entry			

Problem Classification: N.A All results found Satisfactory

Corrective Action: N.A

Reviewed by: [Signature]  
21/5/2023

Dated: 21/5/2023



[Signature]  
21/5/2023





**Metropolis Healthcare Ltd., EQAS**  
**LIFEWELL DIAGNOSTICS PVT LTD (CHANDIGARH) (880)**  
**Outlier And Analyte Summary Report**  
**Result Details For Cycle No 230104 and Sample No 01**

Report Date : 19/05/2023

Instrument : -

Analyte	Standard Unit	Result Value	Accepted Value
@ Neutrophil.	-	40-70%	40-70%
@ Lymphocyte.	-	20-40%	20-40%
@ Eosinophil.	-	<6%	<6%

Legend @ : Acceptable

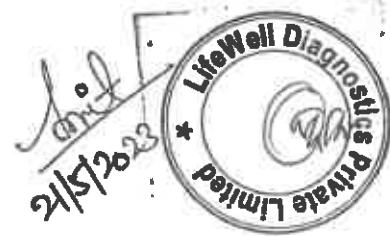
Ⓢ : Acceptable

X : Unacceptable

# : Not Evaluated

Ⓢ : Delayed Result Entry

Total Parameters	3
Not Evaluated Parameters	0
Evaluated Parameters	3
Outlier Parameters (X)	0
EQAS Score Haematology-PS Slide Count Qualitative	100.00 %



*Dr Puneet Kumar Nigam*

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**PT coordinator & Technical Manager, MHL EQAS**  
**Unit No. 409-416.**  
**Commercial Building - 1A**  
**Kohinoor Mall, Kiroi Road, Kuria (W),**  
**Mumbai - 400070**



**INVESTIGATION FORM FOR UNACCEPTABLE PT/EQAS RESULTS**

Issued by QUA:

Department	Haematology		
PT provider	MHL EQAS		
Survey Name:	MHL EQAS 230104		
Date Survey received	07-5-2023	Date Analysis Performed:	12-5-2023
Date Survey Results Submitted	12-5-2023	Date Results received:	22-5-2023

Investigation performed by	AMIT KUMAR		
Lab director	DR. SHWETA	Date	22-5-2023

Unacceptable Result 1			
Specimen No.	03	Analyte	TLC
Reported result	1.5	Intended result/range	1500
Acceptable Lower Limit	775.62	Acceptable Higher Limit	1672.9
SD	224.32	CV%	20.50

Unacceptable Result 2			
Specimen No.	04	Analyte	TLC
Reported result	848	Intended result/range	8480
Acceptable Lower Limit	7824.96	Acceptable Higher Limit	11615.04
SD	947.52	CV%	9.75

Unacceptable Result 3			
Specimen No.		Analyte	
Reported result		Intended result/range	
Acceptable Lower Limit		Acceptable Higher Limit	
SD		CV%	

**EVALUATION OF POSSIBLE SOURCES OF ERROR**

Clerical	Yes	No	N/A
Incorrect transcription of the result from the instrument read-out or report? (Check the raw data.)			✓
Incorrect instrument/method/reagent reported on the result form (Check instrument log book)			✓
Mis-match of the units of measure between the result form and the instrument results			✓
Incorrect decimal placement			✓
Errors in calculations.			✓
Mis-match between result reported on the 'result form' and on the 'proficiency testing evaluation report'			✓
Mis-match between result reported on the 'result form' and on the 'proficiency testing evaluation report'			✓

Any other clerical problem?			✓
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A response of "Yes" to any of these questions may indicate a clerical error.

Although reporting of PT results is unlike those for patient results, clerical errors may indicate a need for additional staff training, review of instructions provided with PT or investigation of the reporting format provided by the testing device. If results reported on the result form do not match the results found on the evaluation report, please contact PT provider.

Procedural	Yes	No	N/A
Written procedure not followed			✓
Improper preparation of reagents			✓
Reagents open stability not within acceptable range			✓
Faulty standards run			✓
Unacceptable IQC results (Run accepted in non-linear range/though controls were out of range)			✓
Incorrect interpretation of microscopic examinations			✓
Incorrect dilution or pipetting error			✓
Incorrect staining or interpretation			✓
Time delay between reconstitution and analysis			✓
Media preparation related			✓
Antibiotic disc potency			✓
Any other procedural problem			✓

A response of "Yes" to any of these questions may indicate a procedural error. These errors indicate inappropriate operation of equipment or performance of a method.

A review of the instructions provided with the proficiency testing material and/or review of laboratory procedures may be required.

Analytical	Yes	No	N/A
Most recent calibration unacceptable or not within established stability limits at the time of PT			✓
Instrument repaired or replaced recently at time of PT run			✓
Review of past PT results indicate unevenly distributed data or a bias			✓
Intended result not within the measuring range for the instrument			✓
Was instrument maintenance performed on schedule?	✓	✓	✓
Review of records indicate there was related instrument/test problems noted prior to or after the PT was performed			✓
Any other analytical problem?			✓

A response of "Yes" to any of these questions may indicate an analytical error. These types of errors could indicate a failure to follow recommended instrument maintenance and calibration.

<b>Specimen handling</b>	<b>Yes</b>	<b>No</b>	<b>N/A</b>
Survey specimens not reconstituted as indicated in the Kit Instructions			✓
Survey specimens not stored as indicated in the Kit Instructions			✓
Special instructions provided in the Kit Instructions not performed as indicated			✓
Correct tests not performed on the correct vial of proficiency testing material			✓
Survey specimen mix-up			✓
Any other problem related to specimen handling?			✓

A response of "Yes" to any of these questions may indicate a specimen handling error. These types of errors could indicate a failure to read the material provided with the Surveys materials.

<b>PT Material</b>	<b>Yes</b>	<b>No</b>	<b>N/A</b>
Late shipment			✓
Hemolysed sample			✓
Bacterial contamination			✓
Perceived survey bias			✓
Poor growth in culture			✓
Unstable PT material			✓
Matrix effect incompatible with method			✓
No comparable peer group			✓
Inappropriate peer group based on method reported on result form			✓
Acceptable range too low			✓
Any other problem?			✓

A response of "Yes" to any of these questions may indicate a problem with the PT material. If a delay in receipt of material in the laboratory is an issue, ensure timely receipt of Surveys after arrival in the institution. If you believe your result was compared to an inappropriate peer group, verify the method reported on the result form. Contact PT provider for additional information if needed.

<b>Miscellaneous</b>	<b>Yes</b>	<b>No</b>	<b>N/A</b>
Was analysis performed by technically competent personnel? (Check technical competence records)	✓		
Were all patient reports satisfactory on day of run? (i.e. there were no patient complaints)	✓		
Were the results of rerun (where possible) the same?	✓		
Any other actions (please specify)			✓

## SUMMARY REPORT

## Root-cause analysis

UNIT issue found.

## Corrective Action proposed

unit changed as requested.



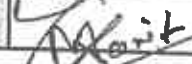
## Corrective Action taken (To be filled within 30 days of receipt of EQAS result)

Unit changed done after that EQAS TLC result found within the  $\pm 2sd$  limits.

## Evidence that problem is solved after corrective action (To be attached)

Attaching instrument printout, EQAS Report.

<b>Preventive Action (if any) proposed:</b> Result will review after or before submitting on the portal by the Lab incharge, Lab head or Lab director.
<b>Conclusion and future plan if no evidence found:</b> Result will review after or before submitting on the portal by the Lab incharge, manager and Lab head or Director.

Recorded by (Sign / Date)	 22-5-2023
Reviewed by Lab/Medical Director (Sign / Date)	 22-5-2023
Reviewed by Head - QA (Sign / Date)	 22-5-2023

  
 22/05/2023



Measure



AS



Database



Print

Sample ID:

HOBC SAMPLE 3 Patient ID:

Date:

10/05/2023 09:47 Mode:

WBC:	1.50	LL	10 <sup>3</sup> /μL	[5.00	-	10.00]	
NEU:	0.00	LL	10 <sup>3</sup> /μL	[2.00	-	7.50]	DIFF
LYM:	0.69	L	10 <sup>3</sup> /μL	[1.30	-	4.00]	
MON:	0.06	LL	10 <sup>3</sup> /μL	[0.15	-	0.75]	
EO:	0.07		10 <sup>3</sup> /μL	[0.00	-	0.50]	
BAS:	1.33	HH	10 <sup>3</sup> /μL	[0.00	-	0.15]	
NEU%:	0.0	LL	%	[40.0	-	75.0]	
LYM%:	45.7	H	%	[21.0	-	40.0]	
MON%:	3.7		%	[3.0	-	7.0]	BAS
EO%:	4.6		%	[0.0	-	5.0]	
BAS%:	88.6	HH	%	[0.0	-	1.5]	
RBC:	1.03	LL	10 <sup>3</sup> /μL	[4.00	-	5.50]	
HGB:	3.1	LL	g/dL	[12.0	-	17.0]	
HCT:	12.0	LL	%	[36.0	-	52.0]	
MCV:	116.7	H	fL	[78.0	-	95.0]	
MCH:	29.7		pg	[27.0	-	32.0]	
MCHC:	25.4	L	g/dL	[30.0	-	35.0]	RBC
RDWsd:	100.1	H	fL	[45.0	-	59.0]	
RDWcv:	—		%	[0.0	-	16.0]	
PLT:	27	LL	10 <sup>3</sup> /μL	[150	-	400]	
PCT:	0.02		%				
MPV:	6.0	L	fL	[8.0	-	15.0]	
PDWsd:	15.7		fL				
PDWcv:	40.0		%				PLT
PLCR:	19.13		%				
PLCC:	5		10 <sup>3</sup> /μL				

Warning flags

ADT

Interpretive flags

Leukopenia?,  
Neutropenia?, Bas

Morphology flags

Down

Up

Measure

AS

Database

Print

Sample ID:

HCBC SAMPLE 04 Patient ID:

Date:

10/05/2023 09:50 Mode:

WBC:	8.48		$10^3/\mu\text{L}$	[5.00 - 10.00]	
NEU:	0.00	LL	$10^3/\mu\text{L}$	[2.00 - 7.50]	DIFF
LYM:	1.72		$10^3/\mu\text{L}$	[1.30 - 4.00]	
MON:	0.47		$10^3/\mu\text{L}$	[0.15 - 0.70]	
EO:	0.25		$10^3/\mu\text{L}$	[0.00 - 0.50]	
BAS:	—		$10^3/\mu\text{L}$	[0.00 - 0.15]	
NEU%:	0.0	LL	%	[40.0 - 75.0]	
LYM%:	20.3	L	%	[21.0 - 43.0]	
MON%:	5.6		%	[3.0 - 7.0]	BAS
EO%:	3.0		%	[0.0 - 5.0]	
BAS%:	—		%	[0.0 - 1.5]	
RBC:	2.15	L	$10^6/\mu\text{L}$	[4.00 - 5.5]	
HGB:	7.7	L	g/dL	[12.0 - 17.0]	
HCT:	28.3	L	%	[36.0 - 52.0]	
MCV:	131.5	H	fL	[76.0 - 96.0]	
MCH:	35.9	H	pg	[27.0 - 32.0]	
MCHC:	27.3	L	g/dL	[30.0 - 36.0]	RBC
RDWsd:	100.2	H	fL	[46.0 - 59.0]	
RDWcv:	—		%	[10.0 - 14.0]	
PLT:	244		$10^3/\mu\text{L}$	[150 - 400]	
PCT:	0.16		%		
MPV:	7.5	L	fL	[8.0 - 15.0]	
PDWsd:	15.2		fL		
PDWcv:	38.0		%		
PLCR:	19.38		%		
PLCC:	41		$10^3/\mu\text{L}$		

System flags

Interpretive flags

Morphological flags

ADT

Neutropenia?  
Anemia? Macrocy

G

Down

Up

11.0

10

10

10

