

(Department of Pathology)

Root cause analysis of outlier parameter for EQAS performance

Format No: SSL/FM/90

SL No	Check Points .	Observa	tions		
Name of	Parameter (s): Unic Acid (SDI-10-46)	Month: F	eboury 1023		
	O/Sample No/Distribution No: Feb, 2023	Year: 2	(023		
Lycie No	ALERRORS Datof testing-16/02/2023				
	d d				
1.	Transcription error (may be pre- or post-analytical factors)		NIL		
2.	Wrong method has been registered for analysis or method change not updated		MIL		
метно	DOLOGICAL PROBLEM				
3.	Instrument function checks (e.g., temperatures, blan readings, pressures) not performed as necessary, or result not within acceptable range.	k S	NIL		
4.	Scheduled instrument maintenance not performe	d	NA		
5.	appropriately.  Incorrect instrument calibration.		MA		
6.	Standards or reagents improperly reconstituted and stored or inadvertently used beyond expiration date.	d,	NA		
7.	Instrument probes misaligned.		NIL		
8.	Problem with instrument data processing functions. The laboratory may need to contact the manufacturer to evaluate such problems.	te	MIL		
9.	Problem in manufacture of reagents / standards, or wi instrument settings specified by manufacturer	th	NIL		
10.	Carry-over from previous specimen.		MIL		
11.	Automatic pipettor not calibrated to acceptable precision ar accuracy.		MIL		
12.	Imprecision from result being close to detection limit method.		MIL		
13.	QC material not run within expiration date, or improper	·ly	MIL		
14.	QC material not run at relevant analyte concentration		MIL		
15.	Result not within reportable range (linearity) for instrume / reagent system.	nt	MIL		



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		Format No: SS
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16.	Obstruction of instrument tubing / orifice by clot or protein.	NIL
17.	Incorrect incubation times.	NIL
	11	MIL
ECHNIC	CAL PROBLEM	
18.	EQA material improperly reconstituted.	NIL
19.	Testing delayed after reconstitution of EQA material (with problem from evaporation or deterioration).	NIL
20.	Sample not placed in proper order on instrument.	MIL
21.	Result released despite unacceptable QC data.	on the testing delie 16/02/23 the Control Yall for un got out side trang
22.	QC data within acceptable limits but showed trend suggestive of problem with the assay.	It shows thread but sample was tested with taking connective act
23.	Inappropriate quality control limits / rules. If the acceptable QC range is too wide, the probability increases that a result will fall within the acceptable QC range yet exceed acceptable limits for EQA.	NIL
24.	Manual pipetting / diluting performed inaccurately, at an incorrect temperature or with incorrect diluent.	MIL
25.	Calculation error or result reported using too few significant digits.	NIL
26.	Secondary specimen tubes incorrectly labeled.	NIL
PROBL	EM WITH PROFICIENCY TESTING MATERIALS	
27.	PT sample with appropriate matrix to that as prescribed by the equipment manufacturer for testing of samples.	NIL
28.	Non-homogenous test material	MIL
29.	Haemolysis on an immune-haemtology program samples.	NIL
PROBL	EM WITH EVALUATION OF RESULTS BY THE PT PROVIDER	
30.	Peer group not appropriate.	NA
31.	Inappropriate target value	NA



## 🕲 इपल्डोच्च SURAHA SPECIALITY LABORATORY

(Department of Pathology)
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SL No	Check Points	Observations				
	1.	<b>.</b>				
32.	Incorrect data entry by PT provider	MH				



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SL No

**Check Points** 

Observations

#### CONCLUSION:

done after checking all points and it has been observed but the outlier of unic acid it was due to the mesult meleased despite unamer--table ac data (Troend) suggested. The parameters Shall be Send for ILC to NABL accomedited 1ab. and parameter shall be strictly monitors in the eaas cycle. next

Root Cause Analysis done by: Saswah' Sengupta

Remarks: UA will be moniton in next cycle. Ith

Reviewed By: Sanwah' Sengupta



## (Department of Pathology) EQAS Corrective Action Details

	Format No: SSL/FM/						
Sl.			Cycle No/	Parameter	Remarks & Corrective Action Taken	Reviewed By	
No		Testing Date	Sample No	Outlier			
1	CMC	16/02/2023	Feb, 2023	Unic Acid	ILC has been done, Afflen	Sanwahi Sengupta	
	Vellowe				Contractive action 2 score		
			i		found 0.70 and the parameters will be monitored in next	Sanwahi Sengupta	
				6			





PATIENT NAME: EQAS-II REF. DOCTOR: DR. SURAHA SPECIALITY LABORATORY

CODE/NAME & ADDRESS: C000131225

SRISHTI DIAGNOSTICS 30, RASH BEHARI AVENUE,

KOLKATA 700026 9123818736 ACCESSION NO: 0031WC012022

PATIENT ID : EQASU630609200

CLIENT PATIENT ID: ABHA NO : AGE/SEX : 1 Days Male

DRAWN : 15/03/2023 19:11:05 RECEIVED : 16/03/2023 14:13:25 REPORTED : 16/03/2023 16:39:04

Test Report Status Final Results Biological Reference Interval Units

#### **BIOCHEMISTRY**

#### URIC ACID, SERUM

URIC ACID

METHOD : URICASE

6.5

3.5 - 7.2

mg/dL

Interpretation(s)

URIC ACID, SERUM-Causes of Increased levels:-Dietary (HighProteinIntake, ProlongedFasting, Rapid weight loss), Gout, Lesch nyhansyndrome, Type 2 DM, Metabolic syndrome

Causes of decreased levels-Low Zinc intake, OCP, MultipleSclerosis

\*\*End Of Report\*\*

Please visit www.srlworld.com for related Test Information for this accession

#### **CONDITIONS OF LABORATORY TESTING & REPORTING**

- 1. It is presumed that the test sample belongs to the patient named or identified in the test requisition form.
- All Tests are performed and reported as per the turnaround time stated in the SRL Directory of services (DOS).
- SRL confirms that all tests have been performed or assayed with highest quality standards, clinical safety & technical integrity.
- 4. A requested test might not be performed if:
- a. Specimen received is insufficient or inappropriate specimen quality is unsatisfactory
  - b. Incorrect specimen type
- Request for testing is withdrawn by the ordering doctor or patient
- d. There is a discrepancy between the label on the specimen container and the name on the test requisition form

- SRL confirms that all tests have been performed or assayed with highest quality standards, clinical safety & technical integrity.
- Laboratory results should not be interpreted in isolation; it must be correlated with clinical information and be interpreted by registered medical practitioners only to determine final diagnosis.
- Test results may vary based on time of collection, physiological condition of the patient, current medication or nutritional and dietary changes. Please consult your doctor or call us for any clarification.
- 8. Test results cannot be used for Medico legal purposes.
- In case of queries please call customer care (91115 91115) within 48 hours of the report.

#### SRL Limited

Fortis Hospital, Sector 62, Phase VIII, Mohali 160062

chaitalily.

Dr. Chaitali Ray, PHD Senior Biochemist cum Management Representative





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View Details

View Report



SRL Ltd

P S SRIJAN TECH PARK BUILDING, DN-52, UNIT NO. 2, GROUND FLOOR, SECTOR V, SALT LAKE, KOLKATA, 700091

WEST BENGAL, INDIA

Tel: 9111591115, Fax: 30203412 CIN - U74899PB1995PLC045956 Email: customercare.saltlake@srl.in



Lab Name SURAHA SPECIALITY LABORATORY Details About RobustAnalysis

Lab No 16610 Detail About Monthly Summary

Month February Detail about SDI

Year 2023

Chemistry II

Constituent

Group

Click on the analyte to view Graphical Data

All Analyser Result

Print

Date of Result Entered: 18/02/2023

Date of Report Published: 08/03/2023

SI.N	o Analyte	Method / Principle	Analyzer Name	No of	DV	Partic	ipants	Your	SDI	U
31.19	o Analyte	Name	Analyzer Name	Participants	DV	CV	SD	Value	301	U
1	GLUCOSE II	GOD-POD II	Agape	494	119.76	7.78	9.32	126.5 mg/dL	0.72	0.84
2	UREA II	Urease UV / GLDH II	Agape	413	125.79	9.60	12.08	149.2 mg/dL	1.94	1.19
3	CREATININE II	Jaffes Kinetic- Alkaline Picrate II	Agape	358	4.19	9.72	0.41	4.64 mg/dL	1.11	0.04
4	T.BILIRUBIN II	Diazonium Salt ( Colorimetric ) / Jendrassik II	Agape	446	1.58	21.85	0.34	1.87 mg/dL	0.84	0.03
5	T-PROTEIN II	Biuret - Colorimetric	Agape	314	4.68	8.32	0.39	4.73 g/dL	0.13	0.04
6	ALBUMIN II	BCG - Colorimetric II	Agape	335	2.99	10.41	0.31	3.49 g/dL	1.61	0.03
7	URIC ACID II	Enzymatic / Uricase Colorimetric II	Agape	384	7.16	10.39	0.74	14.9 mg/dL	10.46	0.08
8	CHOLESTROLII	CHOD-PAP II	Agape	488	96.79	12.52	12.11	104.8 mg/dL	0.66	1.10
9	TRIGLYCERIDE II	GPO-PAP / Enzymatic Colorimetric / End Point II	Agape	328	84.47	11.21	9.47	86.1 mg/dL	0.17	1.05

SDI Range	Interpretation
Within -1.00 to +1.00	Excellent.
Within $\pm 1.01$ to $\pm 2.00$	Good.
Within ±2.01 to ±2.99	Accept with caution. Warning Signal.
Beyond ±3.0	Unacceptable performance. Action Signal.

#### Links

Sanwahi Sengupta

Modify Address	Modify Methods	Submit Lab Results	View Lab Monthly Summary
Yearly Summary	Change Password	Home	Sign-out

### SURAHA SPECIALITY LABORATORY (DEPARTMENT OF PATHOLOGY)

Laboratory 1 : SURAHA SPECIALITY LABORATORY

Laboratory 2 : SRL DIAGNOSTICS

Laboratory	3	:									
SI. No	Date of Testing	ID No	Parameter	Lab 1	Lab 2	Lab 3	Mean	SD	Z-Score	Corrective action ( ≥ 2 Z-score)	Reviewed By
1	16-02-2023		u.acid	14.9	6.5		10.7	5.939697	0.707107		Sanuah' Sengupta

Z- Score = ( Lab 1 - Mean)/SD