



Neu-QAP

(an EXTERNAL QUALITY ASSURANCE PROGRAMME)

PARTICIPANT FINAL ASSESSMENT REPORT (SUMMARY)

PT SCHEME:SPECIAL CHEMISTRY



PC-1047

Cycle No	C3
Ref.No.	NEUQAP109
Sample ID	NEUQAP SBIO SERUM/2021/7/C3/S7

Report Date : 10/08/2021

Sample : JUL 2021

All Methods

	Parameters	Labs (n)	Lab Value	Unit	Assigned value	SDPA	Um	Z-Score	Remarks
1	T3	18	0.99	ng/mL	0.33	0.20	0.058	3.38	Unsatisfactory
2	T4	20	0.82	mcg/dl	1.19	0.54	0.151	0.69	Good
3	TSH	22	0.20	mIU/mL	0.24	0.04	0.012	0.83	Good
4	HbA1c	20	5.60	%	5.59	0.28	0.079	0.04	Good
5	FSH	11	Not Recvd.	mIU/mL	0.49	0.11			N/A
6	PROLACTIN	12	Not Recvd.	ng/mL	1.47	0.18			N/A
7	CPK	12	Not Recvd.	U/L	53.29	4.10			N/A
8	LDH(pyruvate to Lactate)	5	Not Recvd.	U/L	28.80	1.75			N/A
9	MAGNESIUM	11	Not Recvd.	mg/dL	2.12	0.19			N/A
10	PHOSPHORUS	16	Not Recvd.	mg/dL	4.04	0.19	0.060		N/A
11	LH	12	Not Recvd.	mIU/mL	0.60	0.07			N/A
12	Iron	9	Not Recvd.	mcg/dl	26.32	5.19			N/A
13	TIBC	6	Not Recvd.	mcg/dl	59.80	9.78			N/A
14	AMYLASE	12	Not Recvd.	U/L	127.06	6.69			N/A
15	Apo A	4	Not Recvd.	mg/dL	13.50	2.01			N/A
16	Apo B	5	Not Recvd.	mg/dL	12.60	4.76			N/A
17	LDL- Cholesterol	5	Not Recvd.	mg/dL	16.46	1.00			N/A
18	LDH(Lactate to Pyruvate)	9	Not Recvd.	U/L	30.81	2.13			N/A

Authorised Signatory

Dr. Sujay Prasad
Technical Manager and Program coordinator

* denotes adjusted SDPA as per ISO 13528:2015(E)
SDPA - Standard Deviation for Proficiency Assessment

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This report is for use only by the intended participant.



An external quality assurance program from
NEUBERGER ANAND ACADEMY OF LABORATORY MEDICINE PVT.LTD.

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RCA FORM OF ILC / PT / EQAS OUTLIERS

Department:

Date:

1. Proficiency test exception for: T3
2. Proficiency test provider: NEU-QAP
3. Proficiency test analyte group:
4. Cause for PT exception: ① This is the first-PT Analyt(M) with NEU-QAP as per NABL(MELT) program
5. How did the section investigate the cause: ② The external PT provider BEORAD-IT Analyt(M) is not satisfactory and reports of this have been attached.
6. What was the status of the internal QC on the day PT initially analysed: passed
7. Category into which the cause will fit into:
 - A. Method
 - B. Technical
 - C. Clerical
 - D. Problem with PT material Other
 - ① E. No explanation after investigation
8. Evidence that the problem was corrected successfully: Attached
9. Specific corrections taken to prevent the recurrence if possible:
10. Signatures :
 - Quality manager: [Signature]
 - Pathologist:

③ PT with NEU-QAP under observation ^{→ parameter}

④ External comparison satisfactory reports attached

Troubleshooting Guidelines

Method	Technical	Clerical	Problem with PT material	No explanation after investigation
Equipment function checks	Misinterpretation / Wrong identification / Wrong labeling	Transcription error	Leaked / broken vial / not fit for analysis	Use this choice only when the investigation has yielded no satisfactory explanation
Scheduled maintenance not carried out or out of acceptable range	Dilution error / incorrect pipetting	Registration of wrong method or method change is not updated	Bacterial contamination	
Problem with data processing functions	Time delay between reconstitution and analysis		Perceived survey bias / inappropriate target value	
Faulty standard or other reagent	Calculation error			
Incorrect calibration	Analysis accepted in nonlinear range		Unstable material	
Carry over from previous specimen	Analysis done even though controls were out of range or controls not assayed		Matrix effect incompatible with method	
Result close to the detection limit of method	QC data within acceptable limits but showed trend suggestive of problem with the assay		No comparable peer group	
			Acceptable range too low	
	Sample mix up		Late shipment	
Other method related problem	Other technical problem		Improper package and temperature control	

Note: When all identifiable sources of error have been excluded, a single unacceptable result may be attributed to random error, particularly when the result of repeat analysis is acceptable. In such cases, no corrective action should be taken; as such an action may actually increase the probability of a future unacceptable result.

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Lab 164803
PRIMA DIAGNOSTICS
#93, 1ST MAIN ROAD, 1ST PHASE, SECTOR A,
BANGALORE
INDIA

Sample Summary Report
Clinical Chemistry (Monthly) Program

Cycle 19
Jul 2020 – Jul 2021
Sample No: 12
Sample Date: 28 Jun 21
Lot No: 211800



Instrument: SENA CORE – ST-100/100B/100C /200

Analyte	Unit	Result	Mean	Z-score	RMZ	Comparator
✓ Chloride	mmol/L	111.9	113	-0.27	-0.07	Peer
✓ Potassium	mmol/L	5.45	5.86	-1.58	-0.90	Peer
✓ Sodium	mmol/L	145.3	150	-1.20	-0.57	Peer

Instrument: VITROS Microslide Series

Analyte	Unit	Result	Mean	Z-score	RMZ	Comparator
✓ Albumin	g/dL	4.8	5.07	-1.76	-0.28	Peer
✓ Alkaline Phosphatase	U/L	345	354	-0.48	0.76	Peer
✓ ALT (ALAT/GPT)	U/L	170	172	-0.32	-0.21	Peer
✓ AST/GOT	U/L	361	343	1.45	0.76	Peer
✓ Bilirubin, Direct	mg/dL	1.8	1.77	0.08	-0.80	Peer
✓ Bilirubin, Indirect/BU	mg/dL	6.1	6.17	-0.27	1.37	Peer
✓ Bilirubin, Total	mg/dL	7.9	8.01	-0.43	-0.10	Peer
✓ Calcium	mg/dL	14.4	15.2	-1.62	0.58	Peer
✓ Cholesterol, HDL	mg/dL	70	72.5	-0.55	0.40	Peer
✓ Cholesterol, Total	mg/dL	358	374	-0.79	0.17	Peer
✓ Creatinine	mg/dL	1.29	1.33	-0.88	0.37	Peer
✓ G-Glutamyltransferase	U/L	449	435	0.98	1.40	Peer
✓ Glucose	mg/dL	132	137	-1.50	-0.71	Peer
∇ Protein, Total	g/dL	7.2	7.62	-2.45	-0.83	Peer
✓ Triglycerides	mg/dL	213	223	-1.23	-0.44	Peer
✓ Urea	mg/dL	170.7	171	-0.04	0.09	Peer
✓ Uric Acid	mg/dL	8.8	8.84	-0.18	-0.79	Peer

Instrument: VITROS Microwell Series

Analyte	Unit	Result	Mean	Z-score	RMZ	Comparator
✓ T3, Total	ng/mL	2.41	2.27	1.27	0.43	Peer
✓ T4, Total	µg/dL	12.3	12.1	0.34	-0.06	Peer
✓ TSH	µIU/mL	19.5	20.0	-0.52	0.41	Peer

Legend: No Warnings Missing Result Late Results $2.0 \leq |Z\text{-score}| < 3.0$ $|Z\text{-score}| \geq 3.0$
 * Amended Result (per participant's request) Non-robust determination of Mean and SD

Problem Classification: _____

Corrective Action: _____

Reviewed by: _____

Date: _____