



CHRISTIAN MEDICAL COLLEGE

DEPARTMENT OF CLINICAL BIOCHEMISTRY

CMC EXTERNAL QUALITY ASSURANCE SCHEME

MONTHLY SUMMARY REPORT - NOVEMBER 2023



PC-1024

Lab Name **MEDITRUST DIAGNOSTIC CENTRE** Lab No **18608**
 Constituent Group **Chemistry II** Date of Result Entered : **20/11/2023**
 PT item **Lyophilized human serum based** Date of Report Published : **05/12/2023**

Sl.No	Analyte	Method / Principle Name	Analyzer Name	No of Participants	DV	Participants		Your Value	SDI	U
						CV	SD			
1	GLUCOSE II	GOD-POD II	Any Analyser (Automation / Semi Automation)	1594	238.88	9.59	22.91	238 mg/dL	-0.04	1.15
2	UREA II	Urease UV / GLDH II	Any Analyser (Automation / Semi Automation)	1098	67.14	11.47	7.70	69 mg/dL	0.24	0.46
3	CREATININE II	Jaffes Kinetic-Alkaline Picrate II	Any Analyser (Automation / Semi Automation)	1210	4.52	12.57	0.57	4.3 mg/dL	-0.39	0.03
4	T.BILIRUBIN II	Diazonium Salt (Colorimetric) / Jendrassik II	Any Analyser (Automation / Semi Automation)	1211	1.15	23.45	0.27	2.63 mg/dL	5.48	0.02
5	T-PROTEIN II	Biuret - Colorimetric II	Any Analyser (Automation / Semi Automation)	1305	4.77	13.33	0.64	5.79 g/dL	1.60	0.04
6	ALBUMIN II	BCG - Colorimetric II	Any Analyser (Automation / Semi Automation)	1267	2.90	11.42	0.33	3.46 g/dL	1.69	0.02
7	URIC ACID II	Enzymatic / Uricase Colorimetric II	Any Analyser (Automation / Semi Automation)	1323	8.13	12.01	0.98	9.2 mg/dL	1.10	0.05
8	CHOLESTROL II	CHOD-PAP II	Any Analyser (Automation / Semi Automation)	1480	98.77	11.12	10.98	129 mg/dL	2.75	0.57
9	TRIGLYCERIDE II	GPO-PAP / Enzymatic Colorimetric / End Point II	Any Analyser (Automation / Semi Automation)	1379	88.73	18.66	16.56	138 mg/dL	2.98	0.89

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

LAB ADDRESS :

MEDITRUST DIAGNOSTIC CENTRE
439, ANUKUL CHANDRA ROAD
GARIA
WEST BENGAL 700084

Coordinator Contact Details:
 Email: clinqc@cmcvellore.ac.in
 Contact Number: 0416-2283102

Pamela Christudoss
 Dr. Pamela Christudoss
 CMC EQAS Coordinator
 Christian Medical College, Vellore

Homogeneity and Stability of the sample is passed.

Data in CMC EQAS reports is confidential

CMC EQAS does not sub contract any components

***** End of Report *****



INV. No. MTR-INV-L-533/2023(22660)
Patient Name **Ms. DEBANJANA DEBNATH**
Age/Gen 25 Years | Female
Referred By **Dr. Hospital**
Source HOIP

Barcode No 68728
Invoice Generated 15/12/2023 10:09 AM
Sample Received 15/12/2023 10:09 AM
Report Generated 15/12/2023 11:22 AM

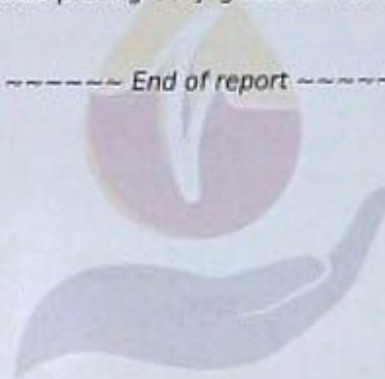



Report Of Biochemistry Examination

Investigation	Result	Unit(s)	Reference Range
BILIRUBIN (TOTAL AND DIRECT)			
Serum Bilirubin (Total) Method (Jendrassik & Grof Method)	0.81	mg/dL	Upto 1.0
Serum Bilirubin (Conj) Method (Jendrassik & Grof Method)	0.20	mg/dL	Upto 0.25
Serum Bilirubin (Uncon) Method (Calculated)	0.61	mg/dL	<0.75

1. In newborns with CB levels ≥ 0.5 mg/dL and < 2 mg/dL, the infection must be ruled out, and the newborn should be observed. In newborns with levels ≥ 2 mg/dL, a more in-depth assessment of the hepatobiliary system is indicated. (interpreting conjugated bilirubin level in newborns,[Epub 2010 Nov 12])

----- End of report -----




Dr. Aniruddha Saha
MBBS, MD
Consultant Pathologist



NAME:	Ms. DEBJANA DEBNATH	UHID No.:	488648
Patient ID:	012312150452	Collection DATE:	15/Dec/2023 07:35 PM
Referred BY:	Dr. HOSPITAL	Received DATE/TIME:	15/Dec/2023 07:39 PM
Age/Gender:	25 Y/Female	Approved DATE/TIME:	15/Dec/2023 08:15 PM
LabID:	1390498	CLIENT Grp.:	MEDITRUST DIAGNOSTICS
SpecimenType:	Serum	Report STATUS:	Final Report

DEPARTMENT OF BIOCHEMISTRY

Test Description	Observed Value	Unit	Method	Biological Ref. Interval
BILIRUBIN (DIRECT And INDIRECT)*				
Bilirubin Total	0.76	mg/dL	Diazo	0.20 - 1.20
Bilirubin, Direct	0.21	mg/dL	Diazo	0.00-0.30
Bilirubin, Indirect	0.55	mg/dL	Calculated	0.20-0.70

*** End Of Report ***

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Dr. Amrita Ghosh
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Dr. Soumick Sarkar
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MEDITRUST DIAGNOSTIC CENTRE
(Department of Pathology)

Root cause analysis of outlier parameter for EQAS performance

Format No: MDC/FM/92

SL No	Check Points	Observations
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Name of Parameter (s): T.BIL	Month: NOVEMBER
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Cycle No / Sample No/ Distribution No: NOV, 2023	Year: 2023
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CLERICAL ERRORS

1.	Transcription error (may be pre- or post-analytical factors)	- NIL -
2.	Wrong method has been registered for analysis or method change not updated	- NIL -

METHODOLOGICAL PROBLEM

3.	Instrument function checks (e.g., temperatures, blank readings, pressures) not performed as necessary, or results not within acceptable range.	- NIL -
4.	Scheduled instrument maintenance not performed appropriately.	- NIL -
5.	Incorrect instrument calibration.	- NIL -
6.	Standards or reagents improperly reconstituted and stored, or inadvertently used beyond expiration date.	- NIL -
7.	Instrument probes misaligned.	- NIL -
8.	Problem with instrument data processing functions. The laboratory may need to contact the manufacturer to evaluate such problems.	- NIL -
9.	Problem in manufacture of reagents / standards, or with instrument settings specified by manufacturer	- NIL -
10.	Carry-over from previous specimen.	- NIL -
11.	Automatic pipettor not calibrated to acceptable precision and accuracy.	- NIL -
12.	Imprecision from result being close to detection limit of method.	- NIL -
13.	QC material not run within expiration date, or improperly stored	- NIL -
14.	QC material not run at relevant analyte concentration	- NIL -
15.	Result not within reportable range (linearity) for instrument / reagent system.	- NIL -

MEDITRUST DIAGNOSTIC CENTRE
(Department of Pathology)

Root cause analysis of outlier parameter for EQAS performance

Format No: MDC/FM/92

SL No	Check Points	Observations
16.	Obstruction of instrument tubing / orifice by clot or protein.	- NA -
17.	Incorrect incubation times.	- NA -
TECHNICAL 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.	- NIL -
21.	Result released despite unacceptable QC data.	- NIL -
22.	QC data within acceptable limits but showed trend suggestive of problem with the assay.	It shows some trend on that date.
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.	- NIL -
25.	Calculation error or result reported using too few significant digits.	- NIL -
26.	Secondary specimen tubes incorrectly labeled.	- NIL -
PROBLEM 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	- NIL -
29.	Haemolysis on an immune-haemtology program samples.	- NIL -
PROBLEM WITH EVALUATION OF RESULTS BY THE PT PROVIDER		
30.	Peer group not appropriate.	- NA -
31.	Inappropriate target value	- NA -



MEDITRUST DIAGNOSTIC CENTRE
(Department of Pathology)

Root cause analysis of outlier parameter for EQAS performance

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SL No	Check Points	Observations
32.	Incorrect data entry by PT provider	- NA -



MEDITRUST DIAGNOSTIC CENTRE
(Department of Pathology)

Root cause analysis of outlier parameter for EQAS performance

Format No: MDC/FM/92

SL No	Check Points	Observations
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CONCLUSION:

T. Bill got outlier item may be due to random error as no ILE had been done as a corrective action with NABL accredited lab and parameter shall be strictly monitored in the next EQAS cycle.

Root Cause Analysis done by: *Guj*

Remarks: Parameter monitored in the next eqas.

Reviewed By: *Guj*



MEDITRUST DIAGNOSTIC CENTRE
(Department of Pathology)
EQAS Corrective Action Details

Format No: MDC/FM/93

Sl. No	EQAS Agency	Sample Testing Date	Cycle No/ Sample No	Parameter Outlier	Remarks & Corrective Action Taken	Reviewed By
1.	eme vellore	18/11/23	November, 2023	T.Bilirubin	ITC has been done after corrective action Z score found 0.70 and the parameter will be monitored in next cycle	
2.	eme vellore	18/11/23	November, 2023	Cholesterol (warning)	REA done and it has been observed as random error. Parameter will be monitored next cycle. Sample has been re-run.	
3.	eme vellore	18/11/23	November, 2023	Triglyceride (warning)	REA done and it has been observed as random error. Parameter will be monitored next cycle. Sample has been re-run.	

MEDITRUST DIAGNOSTIC CENTRE
(Department of Pathology)

Laboratory 1 : DISA
Laboratory 2 :
Laboratory` 3 :

Sl. No	Date of Testing	ID No	Parameter	MDC	Lab 1	Lab 2	Lab 3	Mean	SD	Z-Score
1	15.12.2023		T.BILIRUBIN	0.81	0.76			0.785	0.035355	0.707107

$$\text{Z-Score} = \frac{(\text{Lab 1} - \text{Mean})}{\text{SD}}$$

Corrective action (≥ 2 Z-score)	Reviewed By
NA	