

TELANGANA DIAGNOSTICS

Form: TD/QSP/08-EQCAR

TITLE

EQAS CORRECTIVE ACTION FORM

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EQAS Details	
Analyte:	CBP
Month:	SEPTEMBER
Date Sample Tested:	11.10.2022

SPECIMEN HANDLING	1	61-	0
Were specimens received in an acceptable condition?	Yes 🗵	No	
Were specimens stored according to the instructions on the result forms?	Yes 🕏	No	0
Were the samples hemolyzed?	Yes 🗆	No	0
Were samples tested within the time allowed for sample stability?	Yes 🗸	No	
If applicable, were the samples reconstituted correctly?	Yes 🗆	No	0
Notes: —			
CLERICAL ERRORS	Yes 🕏	No	
Were the results transcribed onto the result forms correctly?		No	
Were the results transcribed from the result forms to the website correctly?	Yes 🗸	-	
Were the results recorded on the correct result form?	Yes 🗸	No	
Was the correct instrument/reagent/kit selected?	Yes 🗸	No	_
Were the results recorded in the correct units?	Yes 🗷	No	
Were the results on your evaluation the same as the results you reported?	Yes 🗗	No	
Notes: — QUALITY CONTROL			
Were quality control materials within the acceptable range on the date of PT testing?	Yes 🗸	No	
Verify the quality control acceptable range in use.)	Yes 🗆	No	Z
s there any indication of trending or shifting of the control results?			
Notes:			
CALIBRATION			
Were there any problems with the most recent calibration?	Yes 🗆		V
When was the last calibration performed?	27-12	-20	22
			1-17
How often is a calibration performed?	AND RESIDENCE OF THE PARTY OF T		
How often is a calibration performed? When was the last calibration verification performed?			

INSTRUMENT			
Were instrument problems noted the day the samples were tested?	Yes	No	4
Has there been any recent maintenance on the analyzer?	Yes	No	D

PREPARED & REVIEWED BY: QUALITY MANAGER: Mr.G RAVI KUMAR

APPROVED & ISSUED BY: LAB HEAD: Dr. A.CHAITANYA

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Have you contacted your analyzer manufacturer for assistance?	Yes		No	0
Notes:				
REAGENTS				
Were the reagents stored properly?	Yes	9	No	
Were the reagents expired or was the open vial stability exceeded?	Yes		No	9
Have there been any changes in reagent manufacturer or formulation?	Yes		No	9
Notes:				
		/		
				1
TESTING PERSONNEL		/		
Date of last competency assessment for testing personnel	Yes	Ø	No	
Review assay procedure and proficiency test sample preparation instructions with testing personnel to ensure that instructions were followed	Yes	9	No	0
Review with testing personnel how samples were loaded to rule out misidentification or transposition of samples.	Yes	9	No	
Notes:		/ / /		
		/		
Corrective Action: RANDOM ERROR				
LITERALL EFFOR				9 76
	Data: 1	1 "	2	22.1
Person Performing Investigation:	Date:	6-1		
ab Director: Chartanya. A.	Date			

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INVESTIGATION SUMMARY: ROOT CAUSE

Pre-analytic Phase of Testing	Analytic Phase of Testing Post-Analytic Phase of Testing	
☐ PROBLEM WITH PT SAMPLE ☐ SAMPLE PROCESSING ☐ DATA ENTRY ☐ OTHER (SPECIFY):	☐ METHODOLOGICAL PROBLEM ☐ TECHNICAL PROBLEM ☐ REAGENT PROBLEM ☐ CALIBRATOR PROBLEM ☐ OTHER (SPECIFY): ☐ OTHER (SPECIFY):	
PREVENTION Preventive action proposed		7
WE WILL M	ONITOR THE VALUES CLOSELY	
Preventive action Plan		7
	CHECK VALUES IN THE NEXT	
Responsibility		
Date Test	ng Personnel	
Date Depa	tment Technical In charge	

PREPARED & REVIEWED BY:	APPROVED & ISSUED BY:
QUALITY MANAGER: Mr.G RAVI KUMAR	LAB HEAD: Dr. A.CHAITANYA
	Chaitanya. A.

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