





PC-1034

CMC VIROEQAS

By National Reference Laboratory (NRL)
Under National AIDS Control Organisation (NACO)
PT Unit, Department of Clinical Virology, Christian Medical College, Vellore-632004, Tamil Nadu Email: viroeqas@cmcvellore.ac.in Phone: 0416-2283455

PANEL: HIV Serology

Laboratory ID: SAICTCKLKOT009

Distribution Round: Round -2, 2022-2023

Panel dispatch date: 12 October 2022

Result received date: 03 November 2022

SL NO	SAMPLE ID	INTENDED RESULT	YOUR RESULT	MATCHING YES / NO	% MATCHING
1	S022201	Positive for HIV-1	Positive for HIV-1	YES	100
2	S022202	Positive for HIV-1	Positive for HIV-1	YES	100
3	S022203	Negative	Negative	YES	100
4	S022204	Negative	Negative	YES	100

REMARKS: PERFORMANCE IS SATISFACTORY

Comments:

Participants are scored based on the qualitative result.

Turnaround time: The time taken to report your results does not form part of your performance assessment.

ICTCs under all SRLs who reported all analyzed specimens accurately

Marker	Number of ICTCs with all four	ICTCs with concordant result from all four	
	specimen results	specimens (%)	
HIV antibodies	153	100	

PT Unit, Department of Clinical Virology, Christian Medical College, Vellore The data in this CMCVIROEQAS reports are confidential.







PC-1034

CMC VIROEQAS

By National Reference Laboratory (NRL) Under National AIDS Control Organisation (NACO) of Clinical Virology, Christian Medical College, Vellore 632

PT Unit, Department of Clinical Virology, Christian Medical College, Vellore-632004, Tamil Nadu Email: viroeqas@cmcvellore.ac.in Phone: 0416-2283455

ICTCs who reported all analyzed specimens accurately under respective SRLs

SRL	Number of ICTC with all four specimen results	ICTCs with concordant result from all four specimens (%)
Alappuzha	25	100
Kottayam	24	100
Kozhikode	37	100
Thiruvananthapuram	29	100
Thrissur	38	100

Assigned Value:

Assigned value is determined using multiple assays/multiple testing of the same analyte.

The proficiency testing material is a pooled lyophilized plasma sample prepared by following institutional proficiency testing standard operating protocol (PT-SOP) by trained staff. None of the work related to testing, preparation and packaging of sample is subcontracted. The homogeneity of the PT material is determined using multiple testing of multiple aliquots. The stability of the testing material is determined by assessing the reactivity of the specimen till the closing date.

Enquires: For queries, please contact CMCVIROEQAS coordinator at email: nrl.cmc@gmail.com

Name of CMCVIROEQAS coordinator

Signature

Dr. Rajesh Kannangai

Department of Clinical Virology

Report Dispatch Date: 22 November 2022

Report authorized by: CMCVIROEQAS Coordinator

END OF REPORT

PT Unit, Department of Clinical Virology, Christian Medical College, Vellore The data in this CMCVIROEQAS reports are confidential.