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Ref. No.:- F.02(83)/RMSCL/S&S/RNA Extraction (Manual) Kit./NIB-15/2020/395 Dated: 2) 10/2020

Corrigendum - II

Subject: - Amendment in Date/Specifications/Conditions

Ref.:- NIB No. F.02(83)/RMSCL/S&S/RNA Extraction (Manual) Kit./NIB-15/2020/354

Dated: 09.10.2020 (Technical bid opening due on dated -22.10.2020)

S. No	Existing date / technical specification		
1.	F.02(83)/RMSCL/S&S/RNA Extraction (Manual) Kit.		
	/NIB-15/2020/354 Dated:-09.10.2020		
ļ	Last date and time of submission	22.10.2020 at	
	of online bids	3.00 PM	
	EMD, Tender fees, RISL fees	22.10.2020 at	
	through challan and Physically	3.00 PM	
	Date and time of opening of	22.10.2020 at	
	Online technical bids	04.00 PM	
2.	LIST OF SPECIFICATION OF VIRAL RNA EXTRACTION		
	(MANUAL) KITS		
	1	ilica membrane	
	column/magnetic bead-based to	chnology ,should	
	be optimized for use with biologi		
	free samples such as serum, plasn culture medium allowing extrac		
	from these samples.	HOIT OF VIEW MINA	
	• The Viral Extracted using this ki	t should be used	
	for downsteam application like		
	time PCR	, .	
	 The Process of extraction using 		
	either centrifugation/vacuum	based/magnetic	
	bead based.		
	 Kit should extract Viral RNA between 40μl-80μl. 	clution volume	
	• Carrier RNA Should be used in t	ha kit to conture	
	maximum amount of the Viral R		
	and carrier RNA should help vira		
	from degradation by RNases.		
ŀ	• Time per extraction should be 30-	60 Min.	
	 Yield of the Viral RNA should be>9 	0% recovery.	
		have necessary	
	components to prevent microb	oial growth and	
	contamination with RNAas.		
	• The extraction kit should be a	ble to work on	
	manualPerfor. Each kit should co magnetic stands to process at lea		
	a time. Firm to quote accessor	si 40 samples at	
	perform the test.	res required to	
-	The Batch of every kit will be valid	dated as per the	

guidelines of ICMR. Extraction of Viral RNA from

(20) SARS-CoV-2 positive and ten(10) SARS-CoV-2

Last date and time of submission 26.10.2020 at of online bids 3.00 PM

Amended date / technical specification

F.02(83)/RMSCL/S&S/ RNA Extraction (Manual) Kit.

/NIB-15/2020/354 Dated:-09.10.2020

EMD, Tender fees, RISL fees 26.10.2020 at through challan and Physically 3.00 PM Date and time of opening of 26.10.2020 at Online technical bids 04.00 PM

LIST OF SPECIFICATION OF VIRAL RNA EXTRACTION (MANUAL) KITS

- Kit should work with silica membrane column/magnetic bead-based technology ,should be optimized for use with biological fluids and cellfree samples such as serum, plasma, swabs and cell culture medium allowing extraction of Viral RNA from these samples.
- The Viral Extracted using this kit should be used for downsteam application like PCR, qPCR, Real time
- The Process of extraction using the kit should be either centrifugation/vacuum based/magnetic bead based.
- Kit should extract Viral RNA clution volume between 40µl-80µl.
- Carrier RNA Should be used in the kit to capture maximum amount of the Viral RNA from sample and carrier RNA should help viral RNA to escape from degradation by RNases.
- Time per extraction should be 30-60 Min.
- Yield of the Viral RNA should be>90% recovery.
- The Elution buffer, should have necessary components to prevent microbial growth and contamination with RNAas.
- The extraction kit should be able to work on manualPerfor. Each kit should come with enough magnetic stands to process at least 48 samples at a time. Firm to quote accessories required to perform the test.
- The Batch of every kit will be validated as per the guidelines of ICMR, Extraction of Viral RNA from (20) SARS-CoV-2 positive and ten(10) SARS-CoV-2





S. No Existing date / technical specification negative samples according to the manufacturer's instruction. • Testing of extracted RNA by targeting SARS-CoV-2 genes along with human RNAseP or any other human housekeeping gene as an internal control (IC) in real time PCR to assess overall RNA extraction efficiency and consistency. • The kit performance would be considered satisfactory if atleast 95% concordance among positive samples and atleast 90% concordance among negative samples. • More than 95% samples showed amplification in internal control. • Validation of kits can be done in the presence of expert from company. Amended date / technical specification negative samples according to the manufacturer's instruction. • Testing of extracted RNA by targeting SARS-CoV-2 genes along with human RNAseP or any other human housekeeping gene as an internal control (IC) in real time PCR to assess overall RNA extraction efficiency and consistency. • The kit performance would be considered satisfactory if atleast 95% concordance among positive samples and atleast 90% concordance among negative samples. • More than 95% samples showed amplification in internal control. • Validation of kits can be done in the presence of expert from company.	negative samples according to the manufacturer's instruction. Testing of extracted RNA by targeting SARS-CoV-2 genes along with human RNAseP or any other human housekeeping gene as an internal control (IC) in real time PCR to assess overall RNA extraction efficiency and consistency. The kit performance would be considered	•
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96 Samples can be processed in one Run.	 among negative samples. More than 95% samples showed amplification in internal control. Validation of kits can be done in the presence of expert from company. 	instruction. Testing of extracted RNA by targeting SARS-CoV-2 genes along with human RNAseP or any other human housekeeping gene as an internal control (IC) in real time PCR to assess overail RNA extraction efficiency and consistency. The kit performance would be considered satisfactory if atleast 95% concordance among positive samples and atleast 90% concordance among negative samples. More than 95% samples showed amplification in internal control. Validation of kits can be done in the presence of expert from company. Magnetic Stands should be in such number so that

Note:-

- It may be noted that if any further amendments are issued then a corrigendum will be published and informed.
- Rest of the terms and conditions will remain the same.

(Shriniwas Meena) **Executive Director (Proc.)**

RMSCL