

Instructions for Use		
Product:	VIRNAEx™ C € IVD	
Cat. No:	ViRNAEx 10 mL: MC-025-2020 ViRNAEx 50 mL: MC-026-2020	
Manufacturer:	MetaCell s.r.o., Videňská 1083, 142 00 Prague Czech Republic +420 732 792 355, info@virnaex.com	MetaCell
Package:	ViRNAEx 10 mL ViRNAEx 50 mL	

### Storage

**VIRNAExTM** solution should be stored at room temperature (15 - 25 °C), after opening at the temperature 4-8°C. When stored under these conditions, the solution will retain full activity until the expiration date indicated on the kit label.

#### Intended use

ViRNAEx™ -Viral Extraction solution is designed for rapid (15min) isolation of viral RNAand DNA (nucleic acids - NA) from nasopharyngeal swabs, buccal swabs, sputum and saliva using specially developed technology to remove inhibitors. The isolation is designed for use on low nucleic acid samples. Isolated NA are suitable for amplification techniques such as RT-PCR of viral NA in clinical samples.

### **Method Principle**

The principle of isolation is based on the lysis of viral particles with the aid of detergents. Isolated NA is ready for direct use in RT-PCR, RT-qPCR and other applications.

### Introduction

The SARS-CoV-2 viral RNA extraction solution **ViRNAEx™** allows rapid extraction of viral RNA intended for the qualitative detection of nucleic acid from SARS-CoV-2 virus in nasopharyngeal swab and / or nasopharyngeal lavage / aspirate samples. Samples are taken from individuals suspected of being infected with COVID -19 by their healthcare provider in a viral transmission medium (VTM). SARS-CoV-2 RNA is generally detectable in nasopharyngeal swab specimens and / or nasal lavage / aspirate specimens during the acute phase of infection.

Testing of samples using **VIRNAEx™** for the SARS-CoV-2 test run is intended for qualified personnel in certified laboratories. Positive results indicate active SARS-CoV-2 infection; clinical correlation with the patient's medical history and other diagnostic information is necessary to determine the patient's infection status.

Positive results for the presence of SARS-CoV-2 RNA do not rule out bacterial infections or coinfections with other viruses. The detected agent may not be the definitive cause of the disease. Laboratories are obliged to report all positive results to the relevant public health authorities.

Negative results do not rule out SARS-CoV-2 infection and should not be used as the sole basis for treatment or other treatment decisions for the patient. Negative results must be combined with clinical observation, patient history, and epidemiological information.

Testing with **ViRNAEx™** is intended for use by trained personnel who are experienced in performing assays using qPCR or other gene expression detection systems.



### Required material not included:

- 100 x 1,5 ml tubes with lid for lysis and sample handling.
- 100 x microtubes with lid for extracted NA storage

## **Required equipment**

Thermoblock Vortex Tubes holder Pipettes: 100 – 1000 μl

Package contents ViRNAEx 10 mL (100 isolations) ViRNAEx 50 mL (500 isolations)

or

### **Precautions**

Please wear gloves when using this product. It is recommended to work in a laminar hood (BIO hazard Class II). Avoid all skin contact with kit reagents. In case of contact, wash thoroughly with water. Do not ingest. See Safety Data Sheets for emergency procedures in case of accidental ingestion or contact, safety data sheets are available on www.virnaex.com.

# Primary sample collection, handling and storage

Collect the indicated samples according to a standard protocol in sampling tubes. Recommended swabs: Darcon swap - immerse swabs in transport medium for MicroTest ™ M4RT or MicroTest ™ M6 viruses (Termo Scientific), FLOQSwabs (Copan) – tampons immerse in UTM -(Universal Transport Medium, Copan). Other sampling kits based on polymeric materials with a transport medium for viruses.

# Do not use cotton swabs due to possible inhibition of the PCR reaction. The transport medium dilutes viruses and at low virus concentrations this dilution may cause false negative results.

Samples should be transported to the laboratory at 4 °C. They are stable for at least 72 hours after sampling at 4 °C. If you are not able to transport swabs to the laboratory at 4 °C, it is possible to transport samples at room temperature within six hours.

For sample storage longer than 72 hours, freeze the sample at -20 ° C.

### Sputum and saliva:

Collect the indicated samples according to a standard protocol in sampling tubes. In the laboratory, dilute 1:1 sputum or saliva with PBS or physiological saline solution. In case of high viscosity, you can add another portion of PBS or physiological saline solution. Then vortex the sample thoroughly. Sputum and saliva samples should be transported to the laboratory at 4 °C. For sample storage longer than 48 hours, freeze the sample at -20 °C.



# **Extraction protocol**



Please wear gloves while working with ViRNAEx<sup>TM</sup>

1. Pipette 100 μl of sample (**transport medium or diluted sputum or saliva**) into 1.5 ml tubes (not included in kit). Add 100 μl **ViRNAEx™** solution into the sample tubes (100 μl). Close the lid and vortex for 10 sec.

ViRNAEx<sup>™</sup> solution has to be mixed before the use

Note: You may add internal isolation control in this step, follow the instructions of the detection kit used, if applicable.

2. Heat the sample at 70 °C for 11 or 15 min. The incubation time is dependent on the type of the virus transfer media. If a new VTM is used, try both time variants at the beginning

3. Add 200 µl Nuclease Free Water into the preheated tube and vortex for 10 sec.

4. Diluted sample is ready to be used for qPCR, pipette the volume needed for qRT-PCR reaction.

### Performance characteristics

The kit was tested for the isolation of viral nucleic acids of coronavirus SARS-CoV-2 from clinical specimens such as buccal swabs, nasopharyngeal swabs, saliva and sputum. Subsequent testing using Real-Time PCR analyses and comparison with other commercial methods verified a sufficient yield of viral NA from the samples.

VIRNAEx<sup>™</sup> showed 100% concordance in 120 clinical samples tested for RNA viru SARS-CoV-2 presence if compared to the standardly used RNA isolation method. Reproducibility of the results reached 100%.

# Warnings and general precautions

This kit is intended for *in vitro* use only.

• As SARS-CoV-2 is a serious pathogen, please follow actual WHO recommendations for BSL2+or BSL3 laboratories!

• Lab safety gloves and respirators FFP3 are necessary for work with coronaviruses. Please work in appropriate biohazard boxes.

Keep in mind that the RNA of some viruses can also cause infection.

• Handle and dispose of all biological samples as if they could transmit infective agents. Avoid direct contact with the biological samples. Avoid splashing or spraying.

• Use vortexes in the Biohazard box only to prevent aerosol contamination.

• Dispose of all used tools, tips and work materials and clothing as potentially infectious and dispose of them in accordance with applicable regulations and recommendations for the handling of highly infectious waste.



• Keep in mind that all reagents and materials you work with may transmit infectious agents. Avoid direct contact with reagents. Waste must be disposed in accordance with adequate safety regulations. Consumables must be incinerated. Liquid wastes containing acids or bases must be neutralized before disposal.

- Wear suitable protective clothing and gloves and protect eyes/face.
- Never pipette solutions by mouth.
- Do not eat, drink, smoke or apply cosmetic products in the work areas.
- Wash hands carefully after handling samples and reagents.
- Work in standard mode of separate rooms: isolation, PCR set up, amplification, detection
- Dispose of leftover reagents and waste in compliance with adequate security measures.
- Read all the instructions provided with the kit before running the assay.
- Follow the instructions provided with the kit while running the assay.
- Do not use the kit after the expiry date.
- Only use the reagents provided and those recommended by the manufacturer.
- · Do not change recommended protocol!

# Manufacturer:

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# **Symbols**

LOT	Batch code
REF	Catalogue number
¥ **	Upper limit of temperature
淤	Keep away from sunlight
8	Do not use if package damaged
¥	Number of tests
C€	Fulfilling the requirements of European Directive 98/79/EC for in vitro medical diagnostic device
IVD	in vitro diagnostic medical device
	Manufacturer
$\square$	Use-by-date