Express Viral RNA Purification Kit (BW-VR6532)

Contents

Kit Contents	2
Introduction	2
Storage and Stability	2
Before Starting	3
Important	3
Materials not Supplied	3
Safety Information	3
Protocol (For extracting viral RNA from infected specimen)	4
Limited Use and Warranty	5

Kit Contents

Catalog#	BW-VR6532-00	BW-VR6532-01	BW-VR6532-02
Preps	10	50	250
Buffer LYE	8 mL	40 mL	200 mL
L Solution	24 μL	120 μL	600 μL
Proteinase K	320 μL	1600 μL	8 mL
RNA Wash Buffer*	3 mL	15 mL	75 mL
Buffer MKB	7 mL	35 mL	175 mL
DEPC-Treated ddH ₂ O	1 mL	3 mL	15 mL
MV RNA Micro Columns	10	50	250
User manual	1	1	1

^{*}Add 12 mL (BW-VR6532-00), 60 mL (BW-VR6532-01) or 300 mL (BW-VR6532-02) 96-100% ethanol to each RNA Wash Buffer bottle before use.

Introduction

Purpose: This procedure describes the extraction of viral RNA including COVID-19 and other RNA virus infected specimen for real-time RT-PCR detection of COVID-19 in respiratory specimens and sera.

Protocol use limitations: The RNA extraction protocol described here has not been validated for platforms or chemistries using clinical samples.

Acceptable Specimens

- Respiratory specimens including: nasopharyngeal or oropharyngeal aspirates or washes, nasopharyngeal or oropharyngeal swabs, broncheoalveolar lavage, tracheal aspirates, and sputum.
 Swab specimens should be collected only on swabs with a synthetic tip with aluminum or plastic shafts. Swabs with calcium alginate or cotton tips with wooden shafts are not acceptable.
 - Serum, plasma, or other body fluid.

Storage and Stability

The guaranteed shelf life is 12 months from the date of production. L Solution and Proteinase K

store at -20°C. All other components can be stored at room temperature (15-25°C).

Before Starting

Prepare all components and get all necessary materials ready by examining this instruction booklet and become familiar with each step.

Wear appropriate personal protective equipment (e.g. gowns, gloves, eye protection) when working with clinical specimens. Specimen processing should be performed in a certified class III biological safety cabinet following biosafety level 3 or higher guidelines.

Important

Calculate and aliquot amount of Buffer LYE to be used in a clean tube and add 3 μL β-mercaptoethanol (β-Me) or 10 μL DTT (1 M) to 600 μL Buffer LYE. Add 2 μL of L Solution per 600 μL of Buffer LYE/β-Me or Buffer LYE /DTT. Buffer LYE contains β-Me or DTT can be stored at room temperature up to 1 month.

Add 12 mL (BW-VR6532-00), 60 mL (BW-VR6532-01) or 300 mL (BW-VR6532-02) 96-100% ethanol to each RNA Wash Buffer bottle before use.

Materials not Supplied

• Tabletop microcentrifuge.

© β-mercaptoethanol or DTT.

© 96-100% ethanol.

Safety Information

Buffer LYE and Buffer MKB contains chaotropic salts, which may form reactive compounds when combines with bleach. Do not add bleach or acidic solutions directly to the preparation waste, wear gloves and protective eyewear when handling.

Protocol (For extracting viral RNA from infected specimen)

 Add 30 μL Proteinase K to 300 μL sample. Mix well. Add 600 μL Buffer LYE-L solution by vortexting for 5 seconds.

Note: The sample range 200-400 μL.

- 2. Incubate the sample at 30°C for 10 minutes.
- 3. Transfer the lysate to a **MV RNA Micro Column** and spin at 12,000 rpm for 1 minute. Discard the 2 mL Collection Tube with the flow-through and put the column back to a new collection tube.
- 4. Add 600 μL Buffer MKB to the column and centrifuge at 12,000 rpm for 1 minute. Discard the flow-through.
- Add 600 μL RNA Wash Buffer to the column and centrifuge at 12,000 rpm for 30 seconds.
 Discard the flow-through.
- 6. Repeat step 5.
- 7. Centrifuge the empty column, with the lid open, at 12,000 rpm for 2 minutes.

Note: It is critical to remove residual ethanol for optimal elution.

- 8. Place the column to a **1.5 mL RNase-free Microfuge Tube**, add **35-50 μL DEPC-Treated ddH₂O** to the column and centrifuge at 12,000 rpm for 1 minute. Store the purified RNA at -20°C.
- 9. Optional: Add the eluent back to the column for a second elution.

Note: The first elution normally yield 60-70% of the DNA/RNA while the second elution yield another 20-30% of the DNA/RNA bound to the column.

Limited Use and Warranty

This product is intended for *in vitro* research use only. Not for use in human.

This product is warranted to perform as described in its labeling and in BEIWO's literature when used in accordance with instructions. No other warranties of any kind expressed or implied, including, without limitation, implied warranties of merchantability or fitness for a particular purpose, are provided by BEIWO. BEIWO's sole obligation and purchaser's exclusive remedy for breach of this warranty shall be, at the option of BEIWO, to replace the products, BEIWO shall have no liability for any direct, indirect, consequential, or incidental damage arising out of the use, the results of use, or the inability to use it product.

For technical support or learn more product information, please contact us or visit our website.



Contact Us: 400-115-2855

www.beiwobiomedical.com

Customer Support:

market@beiwobiomedical.com

Technical Support:

tech@beiwobiomedical.com