

# Magnetic Viral Nucleic Acid Extraction Kit (High Sensitivity) Instructions (Pre-Filled).

#### Composition

Magnetic Viral Nucleic Acid Extraction Kit	64 Preps			
Cat. No.	4012064			
Carrier RNA	310 µg			
Proteinase K	1.4 ml			
Pre-filled 96-well plates	4			
Magnetic rod sleeve	8			
Instructions	1			

#### Storage

Carrier RNA and Proteinase K can be transported at room temperature and stored at -20°C upon receipt. Other reagents can be stored at room temperature (0~30°C) and have a validity period of 2 years.

## **Technical Support**

R&D Department, Hangzhou Simgen Biotechnology Co., Ltd. technical@simgen.cn , Tel: 400-0099-857.

### Introduction

This product is designed for the extraction of a variety of viral RNA or viral DNA from 200  $\mu$ l plasma, cell-free body fluids (including plasma, serum, urine, CSF, and cell culture supernatant), viral stocks, and virus-infected tissue lysates. In the 96-well plate of the pre-filled kit, only the sample and proteinase K are added to the wells of columns 1 and 7, and the instrument can automate a series of processes such as release, adsorption, washing and elution of viral nucleic acids, and the final nucleic acids can be immediately used in PCR or RT-PCR reactions.

### Equipment and Reagents to Be Supplied by User

- 1. Pipettes and tips (to avoid contamination between samples, use RNase-free/DNase-free pipette tips with filters)
- 2. Disposable gloves and protective equipment and tissues
- 3. The Sim-300 Automatic Nucleic Acid Extractor or a similar instrument compatible with consumables.

#### **Preparation before use**

Add all proteinase K to the tube contains Carrier RNA and vortex until all the Carrier RNA is dissolved. Proteinase K with Carrier RNA should be stored at -20°C for 6 months without affecting the use effect. If not used for a long time, store the Proteinase K with Carrier RNA at -80°C.



#### Protocol

#### Sample pre-use treatment

A. Plasma, serum, cell-free body fluid, virus sample preservation fluid, virus stock solution, urine specimens fluid, cerebrospinal fluid, herpes fluid, CSF and cell culture supernatants, etc.:

Use 200 µl of sample directly to isolation and purification; if the sample volume is less than 200 µl, add PBS solution to 200 µl.

- \* Extraction of viral nucleic acids using freshly isolated or freeze-thawed samples no more than one time, whenever possible.
- B. Pharyngeal swab wash fluid, genital tract swab wash fluid, mouthwash fluid: Transfer 300 μl sample into a 1.5 ml centrifuge tube, centrifuge at 12000 rpm for 5 min, then transfer 200 μl supernatant for extraction of viral nucleic acids.
- C. Virus-infected tissue lysate:

Take 10 mg of virus-infected tissue, add liquid nitrogen to immerse the tissue for grinding, after grinding, add 300  $\mu$ l PBS solution for suspension, then transfer 200  $\mu$ l tissue suspension for isolation and purification of viral nucleic acids.

#### **D.** Stool

Add 1 ml of normal saline to a 1.5 ml centrifuge tube, take about 200 mg stool with a sterilized toothpick (if the stool are in liquid form, aspirate 200  $\mu$ l directly), add to the 1.5 ml centrifuge tube and vortex until the stool are completely dispersed. Centrifuge at 12,000 rpm for 1 min and transfer 200  $\mu$ l the top supernatant for isolation and purification of viral nucleic acids.

- **1.** Tear the aluminum foil from a 96 deep-well plate and add 20 μl of Proteinase K with Carrier RNA dissolved to each well in columns 1 and 7 of the deep-well plate.
- 2. Add 200 µl of body fluid samples to each well of column 1 and column 7 of the 96 deep-

well plate and place the 96 deep-well plate into the automatic nucleic acid extractor.

3. Insert the magnetic rod sleeve in the automatic nucleic acid extractor.

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Step	Hole	Amount	Soak	Stirring	Stirring	Descending	Bottom	Magnetism	Waiting	Stop	Plate1	Plate1	Plate2	Plate2
		of Fluid		intensity	time	magnetism	magnetism	times	time	Off/On	lyse	elute	lyse	elute
		(µL)	(s)	(level)	(s)	(s)	(s)		(s)	0/1	(°C)	(°C)	(°C)	(°C)
1	4	300	0	1	0	30	3	1	0	0	0	0	0	0
2	1	700	0	6	600	30	3	2	0	0	70	0	70	0
3	2	700	0	6	180	30	3	1	0	0	0	0	0	0
4	3	800	0	6	180	30	3	1	0	0	0	0	0	0
5	5	800	0	6	180	30	3	1	600	0	0	0	0	0
6	6	80	0	5	180	30	10	2	0	0	0	85	0	85
7	1	700	0	5	5	5	0	0	0	0	0	0	0	0

4. Follow the steps below to set up the program in the automatic nucleic acid extractor:

\*The above procedure is based on the automatic nucleic acid extractor (Cat. No. Sim-300), if used with other companies' instruments, please adjust the parameters of the program appropriately according to the characteristics of the instrument, or call 400-0099-857 for technical support.

# 5. Collect and transfer viral nucleic acids from columns 6 and 12 to a clean centrifuge tube, or directly seal the 96 deep-well plate with parafilm and store at -20 °C for later use.

\* Since the beads take away some of the eluted nucleic acid solution, the viral nucleic acid solution that can be

collected is about 50 µl.