

MagListo™Viral DNA/RNA Extraction Kit For ExiPrep™96 Lite

Before You Begin

- 1) Completely dissolve one vial of **Proteinase K** in 1,250 μ l of nuclease-free water. For short term storage, dissolved Proteinase K should be stored at 4°C. For long term storage, it is recommended to aliquot the enzyme into separate tubes and store at -20°C.
- 2) Completely dissolve Poly(A) in 500 µl of nuclease-free water. Mix dissolved Poly(A) solution into VB Buffer. Shake it thoroughly.
- 3) Add correct amount of absolute ethanol to VWM1 Buffer.

Sample Preparation and Loading the Kit to the Instrument

- Apply 200 μl of Serum, Plasma, or CSF sample to a new 96-well dome plate.
 (Note) Serum and Plasma can be used below 200 μl.
- 2) Add 200 µl of VB Buffer to the sample.
- 3) Add 10 µl of Proteinase K to the sample.
- 4) Aliquot the solution from MagListo™ Viral DNA/RNA Extraction Kit to each of the new 96-well dome plate using multichannel pipette

Cartridge No.	Solution	Volume
1	Sample + VB Buffer + Proteinase K	Up to 410 μl
2	Magnetic Nano Bead solution	200 μΙ
3	VWM1 Buffer	700 µl
4	VWM1 Buffer	700 µl
<u> </u>	RWA2 Buffer	700 µl
6	80% Ethanol	700 µl
7	ER Buffer	100 μΙ

- 5) Press the 'Plate' Button on the instrument.
- 6) Place the Magnetic Rod Cover to the Magnetic Rod.
- 7) Place the plate onto the proper position of the base plate.
- 8) Press the 'Standard Protocol' Button and select 'K-3624/K-3617 Viral DNA/RNA (V1.0)'.
- 9) Press the 'Run' Button to start the selected protocol.
- 10) When the equipment stops as the message 'Pause...Input Ethanol' pops up, open the door and take out 96-well dome plate (Cartridge ①) containing lysate.
- 11) Add **400 µl** of **100% Ethanol** to 96-well dome plate (Cartridge ①) containing lysate using multichannel pipette.
- 12) After adding ethanol, press the 'Check' Button to resume the extraction.
- 13) When the extraction of nucleic acid is complete, collect the final nucleic acid in about 100 μ l of ER Buffer from the 96-well dome plate (Cartridge \Im).

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