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DATA SHEET

Version: 02
Revision date: 26/05/2023

1. Identification

Product name **HigherPurity™ Viral RNA/DNA Extraction Kit**

Cat. No **AN0605 (100 reactions)**
Cat. No **AN0605-XL (250 reactions)**

2. Description

HigherPurity™ Viral RNA/DNA Extraction Kit is designed for the rapid simultaneous purification of viral DNA and/or RNA from cell –free samples such as serum, plasma, urine, cell free body fluids, cell culture supernatants and rinse liquid from swabs samples.
The kit is based in nucleic acid ability to bind silica in the presence of high concentrations of chaotropic salts. The viral RNA/DNA molecules bind to the silica-based
media and impurities such as proteins and nucleases are removed by thorough washing with Wash Buffer. The RNA/DNA is then eluted in sterile, RNase free water. The isolated viral RNA/DNA is ready to use and should be stored at - 80˚C.
The procedure can be used for isolation of viral RNA/DNA from a broad range of viruses. However, performance cannot be guaranteed for every virus species and must be validated by the costumer. The amount of purified viral nucleic acid depends on the sample type, the virus titer, sample source, transport, storage, and age.
The Kit also includes carrier RNA that improves binding and recovery of low-concentrated viral RNA/DNA.

3. Composition

Item	Quantity	
	AN0605	AN0605-XL
Minispin-columns	100	250
Collection tubes (2 ml)	200	500
Carrier RNA (Lyophilized)*	8 x 0.2 mg	4 x 1 mg
Proteinase K (Lyophilizate)**	100 mg	250 mg
BLY-Buffer	40 ml	100 ml
WB1 Buffer***	33 ml	90 ml
WB2 Buffer***	20 ml	50 ml
RNase-free Water	15 ml	30 ml

* Add appropriate volume RNase-free Water (included with the kit) to lyophilized Carrier RNA to obtain 4 µg/µL Carrier RNA stock solution. Prepare aliquots of 100 µl and store -20°C. For 25 purifications, thaw one vial of 100 µl Carrier RNA and mix thoroughly with 10 ml BLY-Buffer. Mark the label of the bottle to indicate that Carrier RNA was added.

**Dissolve Proteinase K in water to obtain a 20 mg/mL stock solution. The Proteinase K solution can be stored for several days at 2–8 ˚C. For longer-term storage, the unused portion of the solution may be stored in aliquots at –20 ˚C until needed. This product as supplied is stable at room temperature.

**Add the volume ethanol (96%-100%) specified [Not included] to WB1 and WB2 Buffer prior to initial use (see bottle label for volume). After ethanol has been added, mark the bottle to indicate that this step has been completed.

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4. Storage

Store the Carrier RNA(Lyophilized) and Proteinase K (Lyophilized) at -20°C and all other components at room temperature (+15 to +25 ˚C). If any kit reagent forms a precipitate, warm at 55–65 ˚C until the precipitate dissolves and allow cooling to room temperature before use.

5. Applications

The purified viral DNA/RNA is suitable for use in RT-PCR and RTq-PCR and can be used for:

- Viral load monitoring
- Viral detection
- Viral genotyping

6. Quality Control

The quality of **HigherPurity™ Viral RNA/DNA Extraction Kit** is tested on a lot-to-lot basis by isolating viral RNA/DNA from a 200µl serum sample.

7. Further information

Product Use Limitations	This product is developed, designed, and sold exclusively only for research purposes use. The product was not tested for use in diagnostics or for drug development, nor is it suitable for administration to humans or animals.
Safety Information	When working with chemicals, always wear a suitable lab coat, disposable gloves, and protective goggles. For more information, please consult the appropriate material safety data sheets (MSDS). These are available online in convenient and compact PDF format at www.canvaxbio.com where you can find, view, and print the MSDS for each CANVAX kit.

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DETAILED PROTOCOL

1. Transfer 200 µL of sample (plasma, serum, urine, body fluids or cell cultured supernatant) into a microcentrifuge tube (not provided).

• ***Nasopharyngeal swab (NP) /oropharyngeal swab (OP):*** *If the swab is delivered in a transport media suitable for nucleic acid virus stabilization, transfer 200 µl directly into microcentrifuge tube. If you get a swab without transport media, place the swab with into microcentrifuge tube containing PBS and incubate for 15 minutes at room temperature. Afterwards shake the swab vigorously, squeeze it and remove the swab. Use a 200 µl aliquot of the liquid for viral RNA/DNA extraction.*

2. Add 200 µL **BLY-Buffer** (**containing Carrier RNA**) and mix by vortexing vigorously for 20 seconds.

• ***Carrier RNA*** *enhances binding of nucleic acid viral to the silica membrane and reduces the risk of degradation.*

• ***Internal Extraction Control:*** *When performing RNA/DNA extraction, it is often advantageous to have an exogenous source of RNA/DNA template that is spiked into the lysis buffer (BLY-Buffer). This control RNA/DNA is then co-purified with the sample and can be detected as a positive control for the extraction process.*

DO NOT add the internal control directly to the biological sample as this will lead to degradation and a loss in signal strength.

3. Add 50 µL Proteinase K solution and mix immediately.

4. Incubate the mix for 10 minutes at 70°C.

5. Add 100 µL **BLY-Buffer** and mix.

6. Place **Minispin-column** in a 2 ml Collection tube and transfer lysed sample. Centrifuge at 8000xg for 1 minute.
Discard the flow-through.

• For successful nucleic acid purification, it is important to obtain a homogeneous, clear, and non-viscous sample before loading onto the Minispin- Column.

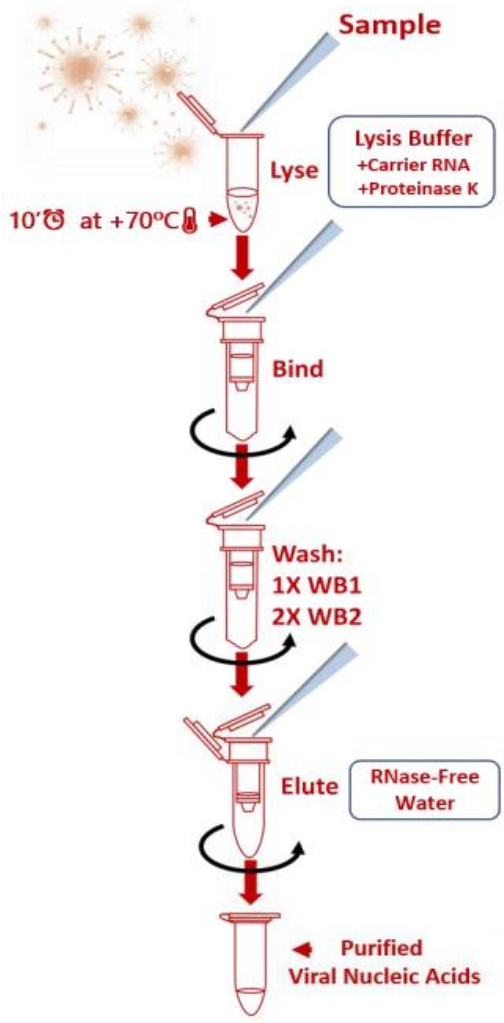
7. Place the **Minispin-column** in a new Collection tube and add 500 µL of **WB1 Buffer**. Centrifuge at 8000xg for 1 minute. Discard the flow-through.

8. Place the **Minispin-column** in the *same Collection tube* and add 500 µL of **WB2 Buffer**. Centrifuge at 8000xg for 1 minute. Discard the flow-through.

9. Place the **Minispin-column** in the *same Collection tube* and add 500 µL of **WB2 Buffer**. Centrifuge at full speed for 1 minute. Discard the flow-through.

10. Centrifuge at full speed for an additional 3 min to dry the **Minispin-column**.

• *This step will avoid the residual liquid to inhibit subsequent enzymatic reactions.*



11. Place the **Minispin-column** into a new, labelled 1.5 microcentrifuge tube (not provided) and pipet 50 µL **RNase-free Water** directly into the membrane. Close the cap and incubate for 2 minutes at room temperature.

12. Centrifuge at full speed for 1 minute to elute. The eluate contains viral DNA and/ or viral RNA. After extraction place the Elution Tube on ice. For long time storage place the nucleic acids at –70˚C.

• *Final eluates contain viral Nucleic acid and Carrier RNA; therefore, it is not possible to quantify the nucleic acids isolated with the kit by photometric or fluorometric methods.*