

# PCRopsis<sup>™</sup> Reagent RVD-RT + PCRopsis<sup>™</sup> Activator

#### For Research Use Only

REF #: 78378001, 78378025, 78378100, 7831000

Store at room temperature

### **INTENDED USE**

PCRopsis™ Reagent RVD-RT is intended for viral nucleic acid extraction-free processing of liquid samples at room temperature.

#### 01 INTRODUCTION

PCRopsis™ Reagent RVD-RT, in combination with PCRopsis™ Activator, is engineered to simultaneously bind a variety of reverse transcriptase quantitative polymerase chain reaction (RT-qPCR) and PCR inhibitors found in specimens, lyse specimens and stabilize nucleic acids in a manner that's compatible with RT-qPCR / PCR. The product consists of a proprietary mixture of peptides, salts, stabilizers, buffers, sodium azide, and RVD Enhancer to achieve these tasks. Reagent RVD-RT allows for nucleic acid extraction-free sample processing without centrifugations or other sample manipulations, which may introduce errors, contaminants, and/or skew the representation of RNA fragments.

#### **02 PRODUCT SIZE**

Catalog Number	Volume
78378001	1 mL
78378025	25 mL
78378100	100 mL
783781000	1000 mL

## **03 STORAGE & STABILITY**

PCR*opsis*™ Reagent RVD-RT is shipped and stored at room temperature. The recommended storage temperature is: 4°C ~ 25°C. Bring product to room temperature if ice crystals are observed. PCR*opsis*™ Activator should be stored at 10–25°C.

## **04 TRANSPORT MEDIUM COMPATIBILITY**

#### **RECOMMENDED:**

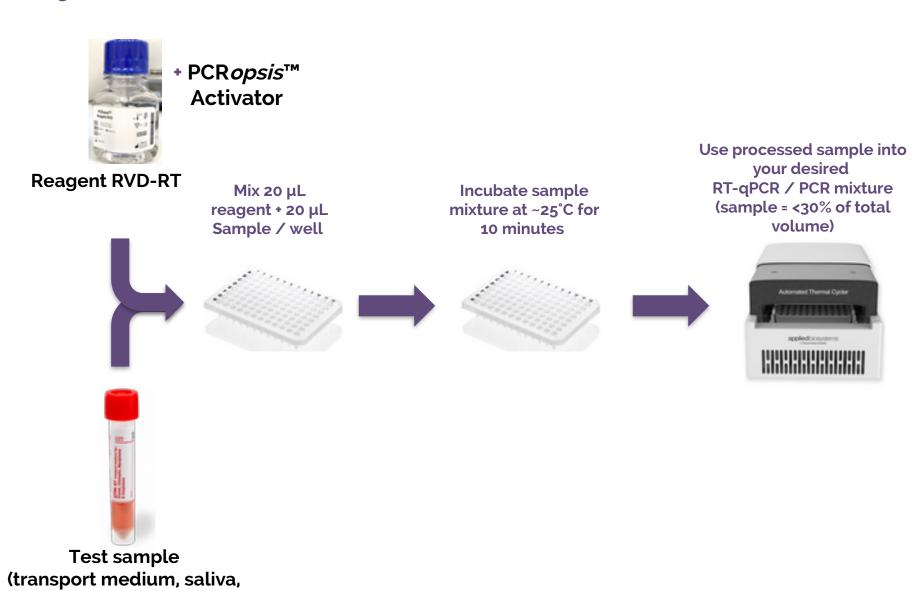
- BD™ Universal Viral Transport System (UVT)
- Quest V-C-M Medium
- MedSchenker™ Smart Transport Medium
- SORFA Viral Transport Medium
- Mediums with formulations similar to BD™ are expected to be compatible
- Phosphate buffered saline (PBS)
- CDC VTM: Hank's Balanced Salt Solution (HBSS) + 2% Fetal Bovine Serum (FBS)
- WHO VTM: Water with veal infusion broth + BSA + antibiotics

#### **NOT RECOMMENDED:**

- Mediums containing guanidinium thiocyanate, alcohols. and other enzyme inhibitors.
- Inactivating mediums

NOTE: the user must confirm the compatibility of Reagent RVD-RT with desired samples

## **05 OVERVIEW OF PROTOCOL**



centrifuged urine)

#### **06 WRITTEN PROTOCOL**

- 1. Thoroughly mix PCR*opsis*™ Reagent RVD-RT to ensure homogeneity, but avoid creating bubbles unnecessarily
  - 1. Reagent RVD-RT has a hazy, white color when homogenized and normal settlement occurs if not regularly mixed
- 2. Thoroughly mix 50 µL PCR*opsis*™ Activator with 950 µL PCR*opsis*™ Reagent RVD-RT. Called **activated Reagent RVD-RT** from here onwards
  - 1. This mixture is stable for 4 hours at room temperature and 24 hours at 4°C in the dark
- 3. Mix 1 volume of activated Reagent RVD-RT (20  $\mu$ L) with 1 volume of sample (20  $\mu$ L) in a sterile tube (0.1 ~ 0.5 mL) or 96-well PCR plate
- 4. Thoroughly pipette up & down to ensure complete mixing and then cap tube or apply plate sealer to plate to prevent evaporation
- 5. Incubate sample mixture at room temperature (~25°C) for 10 minutes
- 6. Immediately mix processed sample and use in your desired RT-qPCR / PCR procedure
  - Processed sample should represent 15%~30% of your final RT-qPCR mixture (i.e., 3~6 μL sample into a total volume of 20 μL) depending on the polymerase used
  - You might observe increasing PCR inhibition when your PCR mixture consist of >35% processed sample

#### **06 WRITTEN PROTOCOL**

### Suggested thermal cycler parameters for RT-PCR / PCR:

- 1. Reverse transcription:
  - a. 45°C for 15 minutes (extend to 30 minutes if suboptimal results observed)
  - b. 95°C for 2 minutes
- 2. PCR amplification (~40 cycles):
  - a. 95°C for 10 minutes (denaturation)
    - a. 95°C for 5 seconds
    - b. 55°C for 15 seconds
    - c. 72°C for 15 seconds
- 3. Hold: 4°C

#### NOTE:

- When amplifying RNA, a 3-step PCR amplification set-up is recommended over a 2-step PCR amplification set-up
- The suggested cycles, temperature, and heating times mentioned above may be optimized by the user as needed

# **07 STEP-BY-STEP PROTOCOL WITH FIGURES**

Step 1 Step 2 Step 3

Gently invert
Reagent RVD-RT
to ensure homogeneity

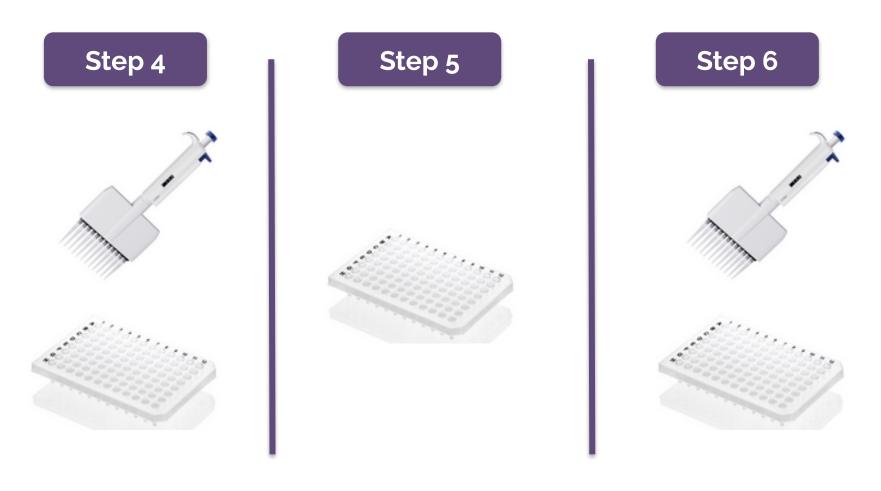
Mix 950 μL Reagent

RVD-RT + 50 μL

Activator to reservoir

Add 20 µL of reagent to wells in a 96-well PCR plate

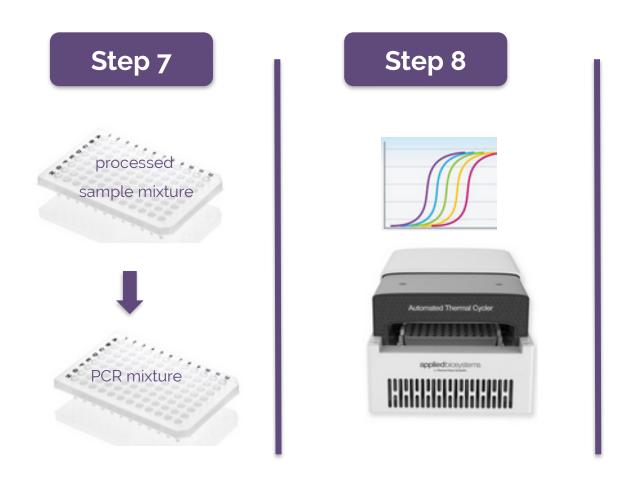
# **07 STEP-BY-STEP PROTOCOL WITH FIGURES**



Mix 20 μL of sample to wells containing 20 μL of reagent Incubate mixture at room temperature for 10 minutes

Add ~15 μL PCR mix from your desired vendor to a new plate

# **07 STEP-BY-STEP PROTOCOL WITH FIGURES**



Mix ~5 μL of processed sample mixture with your PCR mix

Detect amplification of target genes using your desired PCR equipment

#### **08 TROUBLESHOOTING & SUGGESTIONS**

- 1. Reagent RVD-RT, with Activator, is optimized for the amplification of gene targets from specimens in compatible transport mediums and may not be applicable for other applications or mediums.
- 2. Reagent RVD-RT, with Activator, may not thoroughly lyse bacteria at room temperature and a 10~15 minute heating step at 95°C may be required for optimal amplification of target genes.
- 3. For best results, use recently collected samples in compatible transport mediums that have been stored at ~4°C since collection.
- 4. Ensure that the processed sample consist of <30% of the total PCR mixture, since high concentrations of processed sample may inhibit PCR for some applications.
- 5. Take care in maintaining the sterility of your Reagent RVD-RT stock after use.
- 6. Heating of activated Reagent RVD-RT + sample mixture may be required for some specimens if suboptimal results are observed.
- 7. It's recommended to use the processed activated Reagent RVD-RT + sample mixture for downstream PCR applications within 4 hours to ensure optimal results.

## **09 CONTACT**

Contact our research team if assistance with Reagent RVD-RT and / or Activator is necessary (info@entopsis.com). We will try our best to assist with non-intended applications of this product or direct you to alternative products. Any business related questions should be directed to: Sales@PCRopsis.com.



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PROUDLY







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**NOT FOR RESALE** 

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