# Quick Guide / Protocol

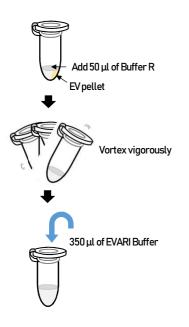
### Protocol: EV RNA Extraction

#### Revision Number: 2.2

Store at room temp.

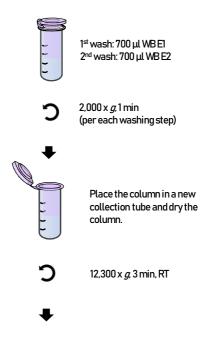
### Step1

Resuspend the EV containing pellet with 50  $\mu l$  of Buffer R. Then, thoroughly mix 350  $\mu l$  of EVARI Buffer by vortexing.



## Step 4

Discard the flow-through and wash the column.



Transfer the column into a new microcentrifuge tube

40 µl of Elution Buffer

12,300 x g, 2 min, RT

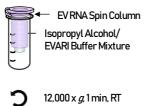
## Step 2

Add 350  $\mu l$  of isopropyl alcohol and mix vigorously. (\*Optional : Add Spike RNA if applicable)



## Step 3

Pipet up the mixture of 700  $\mu$ l, including any precipitate, into Small RNA Spin Column in a 2 ml collection tube. Close the lid and centrifuge at 12,000 x g for 1 min at room temperature.



Kit Contents

Step 5

and elute RNA.

XENO-EVARI KIT (Cat. No. EVARI-765)	Amount
EVARI Buffer	20 ml
Washing Buffer El (WB El)	8 ml
Washing Buffer E2 (WB E2)	8 ml
Elution Buffer	4 ml
Buffer R	4 ml
EV RNA Spin Column	50 ea
Collection Tube	50 ea