Protocol for <u>Zika virus</u> (ZIKV) RT-qPCR positive control using <u>ZIKV Armored RNA</u> (adapted to <u>TaqMan platform</u>)

For RUO (Research Use Only)

Vial content

ZIKV Armored RNA (ArRNA) is a positive control to be used in combination with RT-qPCR assay <u>Lyoph-P&P</u> (lyophilized primers and probe) <u>ZIKV Duo</u> for the detection of Zika virus (ZIKV). The target RNA sequence is encapsidated through an *in vitro* protocol to provide a virus-like particle; therefore, the ArRNA must be extracted (such as virus supernatant or a clinical sample) before RT-qPCR.

Caution

Vials containing ArRNA must be stored at -20°C in the dark after reception. Stable for at least 6 months under the described conditions.

Instructions

This protocol is adapted to the following kit and extraction/RT-qPCR platforms:

- EZ1 Virus Mini Kit v2.1 (QIAGEN) on EZ1 extraction platform (QIAGEN)
- SuperScript™ III Platinum™ One-Step qRT-PCR Kit (ThermoFisher)
- CFX (BIORAD) RT-qPCR thermal cycler

Use of other kits or extraction/RT-qPCR platforms may require adaptation of the protocol.

I. Rehydration of the ArRNA

- Write the date on the vial before opening.
- Resuspend ArRNA as follows to obtain the stock solution (these steps are critical to ensure adequate homogenization):
 - ✓ Add 500µL of nuclease-free water
 - ✓ Homogenize by pipetting 10 to 20 times the volume up and down in the glass vial
 - ✓ Incubate rehydrated ArRNA at room temperature for 10 minutes
 - ✓ Perform a second series of 10-times pipetting

II. <u>Preparation of the working solution</u>

- Extract 400μL of the stock solution on EZ1 platform (QIAGEN) using EZ1 Virus Mini Kit v2.1
- Choose 60μL as elution volume
- Dilute 60μL of eluates in 440μL of PBS to obtain working solution
- If the positive control is used several days after extraction, the working solution should be aliquoted to avoid several freezing/thawing cycles, and be stored at -80°C for optimal stability.

III. RT-qPCR

- Prepare PCR mix using RT-qPCR assay Lyoph-P&P (lyophilized primers and probe), previously rehydrated as explained in the SoP 'Protocol for Real-Time RT-PCR assay with Lyoph-P&P (adapted to TaqMan platform)' (Table 1)
- Perform RT-qPCR with the cycling program in Table 2

Table 1. Preparation of the PCR mix.

	Volume/PCR test (μL)	
2X Reaction mix	12.5	
Rehydrated Lyoph-P&P	7	
SSIII/Taq enzyme mix	0.5	
Total (1 PCR test)	20	
Template RNA (H ₂ O for Ctrl-) 5		
Final volume	25	

Table 2. Cycling program.

50°C	15min	
95°C	2min	
95°C	15sec	45 cycles
60°C	45 cycle	

Using this protocol, a vial contains material for 100 PCR runs.