

# Leadgene<sup>®</sup> 2X Fluorescent RT-LAMP Master Mix

Cat no. LDG0025RF

# **Product Overview**

### Package component

Item	Content
2X Fluorescent RT-LAMP	1 vial (1 mL)
Master Mix	
50X LAMP Fluorescent Dye	1 vial (40 μL)

### Description

Leadgene<sup>®</sup> 2X Fluorescent RT-LAMP Master Mix is an optimized master mix for reverse-transcription loopmediated isothermal amplification (RT-LAMP) reactions. This product is a dual enzyme system, providing a rapid and sensitive detection in one pot. A fluorescent dye is also supplied with the kit. The LAMP reactions can be monitored through real-time fluorescence detection.

### Storage and Stability

Stored at -20°C. Protect from light. Avoid repeated freeze/thaw cycles.

### Procedure

The following procedure is a general guideline for RT-LAMP reaction. To maintain an RNase-free environment, always wear disposable gloves, and use laboratory consumables and water of nuclease-free grade during the whole experiment course.

# RT-LAMP reaction set-up:

## 1. 10X LAMP primer mix

Component	10X concentration	Final concentration	
FIP	16 µM	1.6 µM	
BIP	16 µM	1.6 µM	
F3	2 µM	0.2 μΜ	

# Product Information & Manual

Information of other products is available at: www.leadgenebio.com

B3	2 µM	0.2 μΜ	
LOOP F	8 µM	0.8 µM	
LOOP B	8 µM	0.8 µM	

 An overview of the reaction set-up is listed in the table below. Place all required reagents on ice.
Distribute appropriate volumes into each tube before adding template.

Component	Amount	Final concentration
2X Fluorescent	12.5 μL	1X
RT-LAMP Master Mix		
10X LAMP primer mix	2.5 μL	1X
50X LAMP Fluorescent Dye	0.5 μL	1X
Nuclease-Free H <sub>2</sub> O	XμL	-
RNA template	1-2 μL	variable
Total reaction volume	25 μL	-

 Add target RNA template to the detection tube. Gently mix the reaction thoroughly to achieve uniform distribution and avoid making bubbles.

- 4. Incubate at 65°C for 30-60 min.
- 5. After LAMP reaction complete, the enzyme can be inactivated by heating at 80°C for 10 min.
- 6. For real-time detection, collect fluorescent data using the SYBR<sup>®</sup> or FAM channels.

## Important notes

### **Primer concentration**

Primer concentration can be titrated between 0.25X – 1X if undesired background signal appeared.

### Reaction mixture preparation

Fluorescent dye should be freshly added to the reaction mixture.

# Disclaimer

This product is for research use only and is not intended for diagnostic use.





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