

<b>Catalog number</b>	C15034-1ML		
<b>Set package</b>	Cat.	Name	Amount
	C15034-1ML	StablePlus™ 2X Fluorescent RT-LAMP Master Mix	1 mL
		50X LAMP Fluorescent Dye	40 µL
<b>Description</b>	<p>Croyez StablePlus™ 2X Fluorescent RT-LAMP Master Mix is an optimized master mix for reverse-transcription loop-mediated 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. The StablePlus™ version contains nucleic acid stabilizing agent to protect the amplified products.</p>		
<b>Storage</b>	Stored at -20°C. Protect from light. <b>Avoid repeated freeze/thaw cycles.</b>		

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:**

- 10X LAMP primer mix

**Manuel**

Component	10X concentration	Final concentration
<b>FIP</b>	16 µM	1.6 µM
<b>BIP</b>	16 µM	1.6 µM
<b>F3</b>	2 µM	0.2 µM
<b>B3</b>	2 µM	0.2 µM
<b>LOOP F</b>	8 µM	0.8 µM
<b>LOOP B</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
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<b>StablePlus™ 2X Fluorescent RT-LAMP Master Mix</b>	12.5 µL	1X
<b>10X LAMP primer mix</b>	2.5 µL	1X
<b>50X LAMP Fluorescent Dye</b>	0.5 µ	1X
<b>RNA template</b>	1-2 µL	variable
<b>Nuclease-Free H<sub>2</sub>O</b>	X µL	-
<b>Total reaction volume</b>	25 µL	-

\* See Usage Notes for additional guidelines on primer/template preparation.

3. 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® or FAM channels.

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**Usage 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.

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*For Research Use Only.*