Research use only

# 2X qPCR Master Mix-TaqMan Probe

### Product Name and Catalog Number

2X qPCR Master Mix-TaqMan Probe, Cat PCR 153, 2x1mL/5x1mL/10mL for 200/500/1000 reactions (No ROX, Low ROX or High ROX)

### Intended Use

- The 2X qPCR Master Mix is used for real-time qualitative and quantitative qPCR with TaqMan probe.
- The master mix is a premixed, 2X concentrated solution that has all the components except for gene-specific primers, probe and DNA template.

### **Kit Characterizations**

- The kit is designed for singleplex qPCR with TaqMan probe.
- This kit uses *Taq*-Probe DNA polymerase specially engineered for TaqMan probe, which increased 5' to 3' exonuclease activity produces S-shaped curve.
- The concentrations of the primers and probe are variable depending on specific assays and thermocycling protocols (Table 1).
- The preferable PCR product size is ≤150bp.
- The kit has three formulations of ROX, Low ROX or High ROX concentrations for your choice (see Table 2).

### **Kit Contents**

2X Master Mix (2x1mL/5x1mL/10mL for 100/500/1000 reactions) and an instruction for use.

### **Transportation and Storage**

The kit can be transported at  $\leq 4^{\circ}$ C for up to 3 days.

The kit should be kept stable in the dark at -20°C for  $\leq$ 24 months with  $\leq$ 10 times of freeze-thaw cycles. The kit can be stored at 4°C for a weak.

Component	Volume per 20µL	Volume per 10µL	Final concentration
2X Master Mix	10µL	5µL	1X
Primers <sup>a</sup>	Variable	Variable	Each 150- 900nM
TaqMan probe <sup>b</sup>	Variable	Variable	150-250nM
DNA template <sup>c</sup>	Variable	Variable	≤500ng human genomic DNA/20μL
H <sub>2</sub> O	Το 20μL	To 10µL	

Table 1. Setting Up a 20µL or 10µL reaction

#### Footnotes of Table 1

<sup>a</sup> The primer's T<sub>m</sub> should be designed ≥60°C, preferably between 62°C to 65°C, using primer3 software for high efficiency and specificity.

<sup>b</sup> The probe's  $T_m$  should be 8-10°C higher than the primer's  $T_m$ , preferably between 70-75°C.

<sup>c</sup> DNA templates should be extracted by a qualified silicabased kit and eluted with low EDTA TE buffer (10mM Tris-HCl, 0.1mM EDTA, pH 8.0-8.3).

#### **Applicable Instruments**

#### Table 2. Compatible instruments

qPCR Instrument	ROX required by instrument	Passive dye setup
Bio-Rad <sup>®</sup> iQ <sup>™</sup> 5, CFX96, CFX384, Opticon Roche Lightcycler <sup>®</sup> Qiagen Rotor- Gene <sup>™</sup> Eppendorf Mastercycler <sup>®</sup> Cepheid® SmartCycler <sup>®</sup>	Not recommended	Not necessary
Applied Biosystems <sup>®</sup> 7500, 7500 Fast, QuantStudio™, ViiA7™, Agilent	Low ROX (50nM final concentration)	Turn on ROX passive reference

Мх™		dye button
Applied Biosystems <sup>®</sup> 5700, 7000, 7300, 7700, 7900, 7900HT, 7900HT Fast, StepOne™, StepOnePlus™	High ROX (500nM final concentration)	Turn on ROX passive reference dye button

# Setting Up Thermal Cycling

### Table 3. Standard thermocycling protocol

Stage	Temperature	Period	Number of cycles
I	95°C	2min	1
11	95°C	10sec	
	60°C, signal acquisition	60sec	35-40

## Footnotes of Table 3

The primer concentration used is typically 0.2uM.

# Table 4. Fast thermocycling protocol

Stage	Temperature	Period	Number of cycles
I	95°C	1min	1
11	95°C	5sec	
	60°C, signal acquisition	30sec	35-40

### Footnotes of Table 4

The product size for the fast thermocycling protocol is preferred to be less than 90bp.

The primer concentration used is typically between 0.4 uM and 0.9 uM.

# **Quality control**

Not detectable DNase and RNase contaminations.

### **Related Products**

- 2X Multiplex qPCR Master Mix-TaqMan probe, Cat qPCR 156
- 2X Fast qPCR Master Mix-SYBR Green, Cat qPCR 157

# Precautions

If you order a "**No ROX**" master mix but you have an Applied Biosystems/ThermoFisher instrument, please **turn off ROX passive reference dye button** when setup assays.