

TAQuest[™] qPCR Master Mix for TaqMan Probes *High ROX*

Catalog number: 17286, 17287 Unit size: 1 mL, 5 mL

Component	Storage	Amount (Cat No. 17286)	Amount (Cat No. 17287)
TAQuest™ qPCR Master Mix for TaqMan Probes	Freeze (< -15 °C), Minimize light exposure	1 mL	5 mL
High ROX			

OVERVIEW

TAQuest[™] qPCR Master Mix for TaqMan Probes is a ready-to-use 2X solution optimized for qPCR and 2-step RT-qPCR compatible with TaqMan gene expression assays. The master mix provides all of the essential components including our proprietary TAQuest[™] Hot Start Taq DNA Polymerase enzyme and dNTPs in an optimized PCR buffer, except the template, primers and probes. The Hot Start Taq DNA polymerase allows you to set up a PCR reaction at room temperature, thus minimizing non-specific product formation. The optimized composition ensures PCR specificity and sensitivity with all sample types such as genomic, plasmid, viral and cDNA templates. TAQuest[™] qPCR Master Mix for TaqMan Probes has been designed to be used for duplex reactions using internal positive controls. This master mix contains a high amount of ROX reference dye.

KEY PARAMETERS

qPCR

Instrument specification(s)

ication(s) Filter based on probes

SAMPLE EXPERIMENTAL PROTOCOL

The following protocol can be used as a guideline.

Note Thaw the TAQuest[™] qPCR Master Mix for TaqMan Probes *High ROX* at room temperature. Vortex qPCR Master Mix thoroughly before use.

- 1. Prepare one of the following reaction mixes as indicated in Table 1.
- 2. Carefully mix the reagents with a gentle vortex followed by a brief centrifuge.
- 3. Set up the plate in the qPCR instrument and run as indicated in Table 2.

Table 1. Reagents composition per well for each reaction

Components	Volume (25 µ L/reaction)	Volume (50 µ L/reaction)	Final Conc.	
TAQuest™ qPCR Master Mix for TaqMan Probes *High ROX*	12.5 µL	25 µL	1X	
Upstream primer, 10 µM	0.25-2.5 µL	0.5-5.0 μL	0.1-1.0 μM	
Downstream primer, 10 µM	0.25-2.5 μL	0.5-5.0 μL	0.1-1.0 µM	
TaqMan Probes, 10 µM	0.25-0.625 µL	0.5-1.25 µL	100-250 nM	
DNA template	1-5 µL	1-5 µL	Optimized conc.	
Nuclease-Free Water	25 µL	50 µL		

Table 2. Thermal cycling parameters

Parameter	Polymerase Activation	PCR (30-40 cycles)		
	Hold	Denature	Anneal	Extend
Temperature	95 °C	95 °C	55-65 °C	68-72 °C
Time (m:ss)	0:20	0:30	1:00	1:00

DISCLAIMER

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