

# Beh-taq PCR Master Mix (2X) - 100 Reactions

# **Product Insert**

# Description:

Beh-taq PCR Master Mix (2X) is a ready-to-use Beh-taq PCR Master Mix (2X) solution that contains a PCR internal control which can be detected by HEX/VIC channel in a real-time PCR machine. By detecting the internal control users can validate the DNA template quality, thereby preventing any false negatives in the PCR results. The user needs only to add template, target TaqMan primer/probe mix and water to set up the Beh-taq real-time PCR.

# PCR Control:

Beh-taq PCR Master Mix (2X) contains PCR control primers/probe (HEX/VIC) and PCR control template. The PCR control reaction in the TaqMan 2x PCR Master Mix is optimized to not interfere with target amplification. The fluorescence of the target probe should not be HEX/VIC.

## Advantages:

- Convenience and time savings
- Cost efficient
- · High sensitivity
- Avoid false negatives due to template quality

# Applications:

- Routine TagMan PCR
- · Sensitive detection with internal control

#### Reagents supplied:

Beh-tag PCR Master Mix (2X) (3 Vials, 100 Reactions)

### Storage Conditions:

Beh-taq PCR Master Mix (2X) should be stored at -20°C. For everyday use aliquots can be stored at 4°C for up to 3 months. Beh-taq PCR Master Mix (2X) is stable for multiple freeze-thaw cycles. When stored at the proper temperature this reagent is stable for at least 1 year.

### Tips for Performing PCR Reactions:

Polymerase Chain Reaction (PCR) is a powerful method used to amplify specific DNA transcripts using multiple cycles containing denaturation and annealing/extension steps. Successful PCR relies on various factors, and it is important to keep a number of points in mind when performing PCR:

- 1. Using high quality, purified DNA templates greatly enhances the success of PCR.
- Clean, disposable gloves should be worn at all times when handling reagents, samples, pipettes, disposable tubes, etc. It is recommended that gloves are changed frequently to avoid contamination.
- There should be designated solutions, tips, tubes, pipettes, etc. for PCR only.
- 4. Optimize the template amount: up to 1 μg genomic DNA and 10 pg-100 ng for cDNA or Plasmid.

## Procedure

# Reaction Setup Table

TaqMan PCR Reaction Mixture	Single 20 µL Rxn	10 Rxn + 1 Rxn **
Beh-taq PCR Master Mix (2X)	10 μL	110 μL
Target Primer/Probe Mix*	2 μL	22 µL
Template DNA	2 - 5 µL	2 - 5 µL / rxn
Nuclease-Free Water	Up to 20 µL	Up to 220 μL

<sup>\*</sup> Suggested concentration of primer (F and R) and probe is 2.5 µM. The fluorescence of the target probe should not be HEX/VIC.

- Dispense 10 uL of Beh-tag PCR Master Mix (2X)into the PCR tube.
- Add DNA template (up to 1 μg genomic DNA and 100 ng -10 pg for cDNA or Plasmid) and Target Primer/Probe Mix to the PCR tube as shown in the Reaction Setup Table.
- 3. Add nuclease-free water to bring the total volume to 20  $\mu$ L.
- 4. Mix the PCR mixture thoroughly and spin down briefly.
- Place the PCR tubes into the PCR machine and carry out the PCR according to the Suggested TagMan PCR Program shown in the table below.

# Suggested Beh-taq PCR Cycle Conditions

PCR Cycle	Step	Temperature	Duration
Cycle 1	Initial Denaturation	95°C	3 min
Cycle 2 (40X)	Denaturation	95°C	15 sec
	Annealing / Extension	60°C	30 sec

<sup>\*\*</sup> Experienced User Protocol for Reaction Preparation for Multiple Samples