

# Titan HotTaq Probe qPCR Universal Mix

| Cat. No. | Pack Size            |
|----------|----------------------|
| BT11004  | 1 ml (250 reactions) |

## **Application:**

- DNA/LNA hydrolysis probe-based assays
- · Detection and quantification of DNA and cDNA targets
- Profiling gene expression
- Microbial detection
- Viral load determination

#### **Description:**

Titan HotTaq Probe qPCR Universal Mix is optimized for real-time quantitative PCR assays and contains all the components necessary to perform singleplex or duplex qPCR, with the exception of template, primers, and probes. The qPCR Mix contains optimized components and Titan HotTaq DNA polymerase supplied in a proprietary reaction buffer that enables efficient amplification of regular and GC-rich targets. Titan HotTaq Probe qPCR Universal Mix is optimized for DNA/LNA hydrolysis probes based on the 5' flap endonuclease activity. Titan HotTaq DNA polymerase is activated by a 10 min incubation step at 95°C. This prevents extension of nonspecifically annealed primers and primer-dimers formed at low temperatures during qPCR setup.

# **Reagent Composition:**

- Titan HotTaq DNA polymerase
- Titan HotTaq Probe qPCR Universal buffer
- 15 mM MqCl2

1x PCR solution – 3 mM MgCl2

• dNTPs, including dUTP. The mix allows UNG treatment to prevent carryover contamination from previous runs.

**IMPORTANT:** UNG is not included in the Titan HotTaq Probe qPCR Universal Mix and must be purchased separately.

Internal reference based on ROX dye.

For multiplex application: if ROX dye is used as one of the fluorophores, internal reference might interfere with the signal – a version without ROX is available upon request.

## In a separate vial:

100% DMSO is included in the kit in a separate vial.

# Addition of 100 % DMSO:

| Final conc. Of DMSO | 10 μl/rxn |  |
|---------------------|-----------|--|
| 2.5%                | 0.25 μΙ   |  |
| 5 %                 | 0.5 μΙ    |  |
| 10%                 | 1 µl      |  |

DMSO is recommended as a PCR additive for templates with high GC content. In some cases, DMSO is also required to relax secondary structures. While testing it is recommended to include one sample with additional 2,5 % DMSO to test if it improves the results. For further DMSO optimization the concentration can be raised in 2,5% increments up to 10% based on following table. Volumes are given per reaction depending on final volume of reaction mix. The highest DMSO concentration recommended is 10% which should be used for all templates with GC content over 70%.



#### Benefits:

- Increased sensitivity and specificity for a wide range of templates, including AT-rich, GC-rich and regular cDNA and gDNA.
- Suitable for singleplex and duplex assays.
- Reaction set-up at room temperature the mix is stable at ambient temperature for one month.
- Benchtop stability for 48 hours for pre-assembled reactions.

Wide instrument compatibility: suitable for qPCR cyclers regardless of ROX requirements (except capillary).

# **Shipping and Storage conditions:**

Routine storage: -20°C. Shipping and temporary storage for up to 1 month at room temperature has no detrimental effects on the quality of Titan HotTaq Probe qPCR Universal Mix.

# Recommended qPCR reaction mix:

| Component                                 | Volume      | Final conc. |  |
|---|-------------|-------------|--|
| 5 x Titan HotTaq Probe qPCR Capillary Mix | 4 μΙ        | 1x          |  |
| Primer Forward (10 pmol/µl)               | 0.4-0.8 μΙ  | 200-400 nM  |  |
| Primer Reverse (10 pmol/µl)               | 0.4-0.8 μΙ  | 200-400 nM  |  |
| Probe                                     | x µl        | 100-250 nM  |  |
| Optional UNG (Uracil-N_glycosylase)       | x μl        | x U/ μl     |  |
| Optional 100% DMSO                        | variable    | Up to 10%   |  |
| DNA template                              | variable*   | variable*   |  |
| H₂O PCR grade                             | up to 20 μl |             |  |
| Total                                     | 20 μΙ       |             |  |

<sup>\*</sup>concentration of cDNA 0.1/ µl- 10ng/ µl or gDNA 10 pg/ µl- 4ng/ µl.

## Recommended qPCR cycles:

| Cycle step               | Temp. | Time   | Cycles |  |
|--------------------------|-------|--------|--------|--|
| Optional UNG treatment** | 50°C  | 2 min  | 1      |  |
| Initial activation***    | 95°C  | 10 min | 1      |  |
| Denaturation             | 95°C  | 15-20s | 40     |  |
| Annealing/ Elongation    | 60°C  | 60 s   | 40     |  |

#### **IMPORTANT:**

# Safety warnings and precautions:

For *in vitro* use only

This product and its components should be handled only by persons trained in laboratory techniques. It is advisable to wear suitable protective clothing, such as laboratory overalls, gloves and safety glasses. Care should be taken to avoid contact with skin or eyes. In case of contact with skin or eyes, wash immediately with water.

Some applications this product is used in may require a license which is not provided by the purchase of this product. Users should obtain the license if required.

<sup>\*\*</sup> Add UNG treatment step ONLY if UNG enzyme is added in the reaction mix for carryover contamination removal.

<sup>\*\*\*</sup> To activate the polymerase, include an incubation step at 95°C for 10 minutes at the beginning of the qPCR cycle.