

Manual

Multiplex PCR Mix Probe 4x

High specificity ready-to-use mix for real-time hot-start PCR with fluorescent probe. 4x concentrated.

catalog#	size
2017-2004PM	200 reactions in 25 μl
2017-20040PM	2000 reactions in 25 μl

For research use only.

Guarantee

 $A\&A\ Biotechnology\ provides\ guarantee\ on\ this\ product.$

The company does not guarantee correct performance of this kit in the event of:

- not adhering to the supplied protocol
- $\bullet \qquad \quad \text{use of not recommended equipment or materials}$
- use of other reagents than recommended or which are not a component of the product
- use of expired or improperly stored product or its components

Description

Multiplex PCR Mix Probe 4x is optimized for high specificity ready to use real-time hot-start PCR mixture for use with fluorescent probes. Mixture contains all components required for qPCR except DNA template, primers and probes. *Taq* DNA polymerase is blocked by monoclonal antibody.

The pre-mix formulation saves time and reduces contamination due to a reduced number of pipetting steps required for PCR set up. The mix is optimized for efficient and reproducible reaction.

Contents

	2017-2004PM	2017-20040PM	storage
Multiplex Probe 4x	2 x 0.625 ml	20 x 0.625 ml	-20 °C
ultrapure water	2 x 1.5 ml	20 x 1.5 ml	-20 °C

Multiplex Probe 4x composition

component	amount
modified Taq DNA polymerase	0.2 U/μl
MgCl ₂	20 mM
dNTPs	1 mM of each dNTP
4x reaction buffer	

Notes

- Before use all solutions should be thawed thoroughly on ice, gently mixed by inverting the tube and briefly centrifuged.
- Up to 3x repeated freeze-thaw cycles do not influence the activity of this product.

Example PCR protocol

- 1. Thaw all components of the kit on ice, gently mix by inverting the tubes and briefly centrifuge. Place the tubes on ice again.
- 2. Place PCR tubes on ice and add:

	PCR reaction volume		
component	10 μΙ	25 μΙ	50 μl
Multiplex Probe 4x	2.5 μΙ	6.5 µl	12.5 μΙ
primer 1***	0.1-1 μΜ*	0.1-1 μΜ*	0.1-1 μM*
primer 2***	0.1-1 μΜ*	0.1-1 μΜ*	0.1-1 μM*
probe***	0.05-0.1 μM**	0.05-0.1 μM**	0.05-0.1 μM**
DNA, cDNA template	10 pg-1 μg	10 pg-1 μg	10 pg-1 µg
ultrapure water	up to 10 μl	up to 25 μl	up to 50 μl

^{*} recommended for standard qPCR

- 3. Gently vortex the samples and briefly centrifuge to collect all droplets remaining on the tube walls and caps to the bottom of the tube.
- 4. Place the tubes in the thermocycler and start the PCR programme.

An example amplification profile:

step	temperature	time
initial denaturation	95℃	5 min
25-45 cycles (2 step PCR)	95 °C 58-70 °C*	15-30 s 15-60 s**

^{*} depending on the elongation of the probe and primers temperature

Recommended ROX mixture

HiROX (0.6-1 μ l per 50 μ l of total reaction volume): Applied Biosystems: 7000, 7300, 7700, 7900HT Fast, StepOne, StepOnePlus.

LowROX (0.6-1 μ l per 50 μ l of total reaction volume): Applied Biosystems: 7500, Stratagene: Mx3000P, Mx3005P, Mx4000P

^{**} amount of each probe should be optimized

^{***} final concentration in reaction mixture

^{**} depending on the length of PCR products and/or number of amplicons in the tube



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