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Manual

RT HS-PCR Mix SYBR®

Ready-to-use mix for real-time hot-start PCR with SYBR® Green.
2x concentrated.

catalog #	size
2017-100HS	200 reactions in 25 µl
2017-1000HS	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
- 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

RT HS-PCR Mix SYBR® are optimized ready to use real-time hot-start PCR mixtures with SYBR® Green.
Mixture contains all components required for qPCR except DNA template and primers.

This mix is recommended especially for difficult PCRs and multiplex PCR. StartWarm technology allows for activation of *Taq* DNA polymerase in the first steps of PCR. Long time of preliminary denaturation step is not required.

The premix 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

	2017-100HS	**2017-1000HS**	**storage**
RT HS-PCR Mix SYBR®	2 x 1.25 ml	20 x 1.25 ml	-20 °C, in darkness
ultrapure water	2 x 1.5 ml	20 x 1.5 ml	-20 °C

RT HS-PCR Mix SYBR® A/B/C composition

component	**amount**
***Taq* DNA polymerase**	0.1 U/µl
$MgCl_2$	4 mM
dNTPs	0.5 mM of each dNTP
2x reaction buffer with SYBR® Green	

Notes

- Before use all solutions should be thawed thoroughly on ice, gently mixed by inverting the tube and briefly centrifuged.
- Up to 7x 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 µl**	**25 µl**	**50 µl**
RT HS-PCR Mix SYBR®	5 µl	12.5 µl	25 µl
primer 1**	0.1-1 µM*	0.1-1 µM*	0.1-1 µM*
primer 2**	0.1-1 µM*	0.1-1 µM*	0.1-1 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
** final concentration in reaction mixture

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 °C	3-5 min
25-45 cycles	95 °C 50-68 °C 72 °C	15-30 s 30-60 s 15-60 s*

* depending on the length of PCR products
PCR product melting analysis is recommended.

Recommended ROX mixture

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

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

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