



Bio-Star qPCR-Mastermix SYBR High-ROX (2x)

Features:

- Enzyme with hot start capability increases reaction specificity and sensitivity
- DFS-Taq PLUS DNA polymerase activation requires not more than 5 min heating
- High selectivity and reaction yield
- The mix is colored for easy pipetting
- Reduced preparation time

Applications:

- Real-time PCR with intercalating dye SYBR Green I and HIGH-ROX content
- Conventional PCR
- High-throughput PCR
- Genotyping
- For detection of bacterial DNA

Platforms:

(ABI) 7000, 7300, 7700, 7900, 7900HT, and StepOnePlus.

Description:

Bio-Star qPCR-Mastermix SYBR HIGH-ROX (2x) is developed for quantitative real-time PCR with fluorescent dye SYBR Green I. The Mastermix contains all components, except template and primers, for successful PCR.

- The mix is optimized for efficient and reproducible hot-start real-time PCR of genomic, plasmid and viral DNA samples.
- The solution contains substances that increase half-life and processivity of DFS-Taq PLUS DNA polymerase by enhancing its stability during PCR.
- smart components that influence primer annealing temperature and characteristics of template melting thus enabling to increase the specificity of PCR and use templates with complicated structure.
- DFS-Taq Plus DNA polymerase is inactive at room temperature because of monoclonal antibodies.
- The inert dye allows control when using multi-well plates. Use of the kit saves time and minimizes contamination risk due to reduced number of pipetting steps.

Components and Mixture

2X Bio-Star qPCR-Mastermix SYBR HIGH-ROX contains:

- 100 mM Tris-HCl (pH 8.5 at 25 °C), 100 mM KCl, 0.4 mM each of ultrapure dNTPs, 3 mM MgCl₂, 0.06 U/μl DFS-Taq DNA Polymerase, 0.025% Tween 20, stabilizers and enhancers, SYBR Green I, 900 nM ROX, inert-dye, which does not interfere the PCR in a real-time Cycler
- Tube of MgCl₂ 100 mM
- Water Mol.Bio Grade 2 x 1,25 ml

Storage and transportation: at -20 °C; not more than 50 thawing-freezing cycles. Shipping with blue-ice or at room temperature

Storage terms: up to 24 months

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Amplification protocol:

1. Defrost the reaction mixture and stir thoroughly.
2. Add the following components into the thin-wall PCR tubes considering the final volume of a reaction mixture equal to 50 µl:

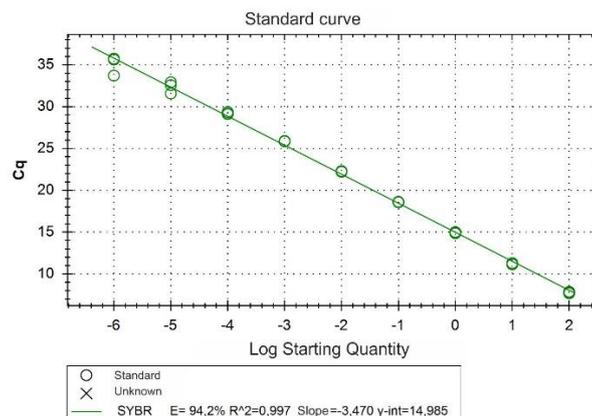
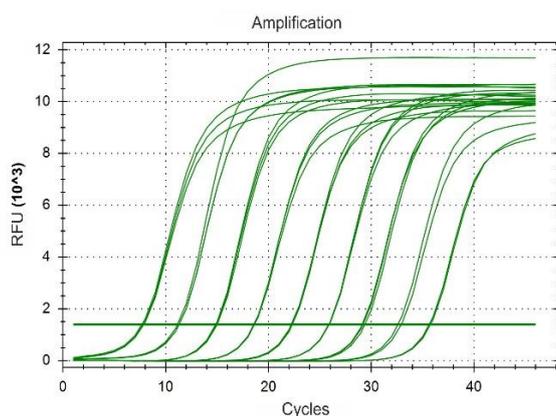
Component	Volume	Final concentration
Bio-Star qPCR-Mastermix SYBR HIGH-ROX(2x)	25	1x
Forward primer	variable	0.1 – 600 nM
Reverse primer	variable	0.1 – 600 nM
DNA template	variable	10 pg – 1 µg
Sterile water	up to 50 µl	

3. Gently vortex and remove droplets by centrifugation.

Cycler Protocol:

Step	Temperature, °C	Incubation time	Number of cycles
Preliminary denaturation	95	1-5 min	1
Denaturation	95	15 sec	25 - 40
Annealing	50 - 68	10-30 sec	
Elongation	58 - 72	30-60 sec	
Melting curve (recommended)			1

Amplification curves and standard curve obtained during real-time PCR using Bio-Star qPCR-Mastermix SYBR HIGH-ROXHIGH-ROX(2x)



NTC – negative control.

Amplification of 18s mRNA of gene fragment in 10-fold serial dilution of cDNA (10 fg – 1 µg). Amplicon length is 120 bp. Reaction was performed at CFX 96 touch BIO-Rad thermal cycler. Amplification curves and standard curve show area of system linearity

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Activities:

Exodeoxyribonuclease activity

DNA was stable after incubation of 1 µg fragment of phage lambda DNA in the presence of 25 µl of **Bio-Star qPCR-Mastermix SYBR HIGH-ROX (2x)** in 50 µl reaction solution at 37 °C and 70 °C for 4 h.

Ribonuclease activity

Absence of ribonuclease activity was confirmed after incubation of 1 µg of 5'-[P³²]-labeled RNA fragment in the presence of 25 µl of **Bio-Star qPCR-Mastermix SYBR HIGH-ROX(2x)** in 50 µl reaction solution at 37 °C for 4 h.

DFS-Taq Plus DNA Polymerase features

Recombinant DFS-Taq Plus DNA polymerase possesses 5'→3' DNA-dependent polymerase activity and 5'→3' exonuclease activity of native Taq DNA Polymerase from *Thermus aquaticus*. The rate of DNA synthesis by Taq polymerase depends on the complexity of DNA template and is approximately 1 kbp/min. Recombinant HS-Taq DNA Polymerase is ideal for conventional PCR and real-time PCR. Free of E-coli DNA

SYBR Green I

SYBR Green I is a fluorescent intercalating dye for quantitative and qualitative detection of PCR products during real-time PCR. SYBR Green I provides easy and economical way for detection and quantitative assessment of PCR products during real-time PCR without a need of using specific fluorescent probes. During amplification, SYBR Green I dye penetrates into the minor groove of DNA products and emits stronger fluorescent signal than unbound dye. Absorption and emission maxima of SYBR Green I are 494 nm and 521 nm, respectively, which enables to use it for every real-time PCR platform existing to date.

Inert dye

The inert dye included in **Bio-Star qPCR-Mastermix SYBR HIGH-ROX (2x)** does not reduce PCR efficiency; it facilitates monitoring of multi-well plate pipetting. Absorption maximum of a HIGH-ROX dye is 615 nm.

Frequently ask Questions

Recommendations for avoiding contamination during PCR

- Preparation of DNA samples, preparation of reaction solutions, amplification and analysis of PCR products should be carried out in different territorial areas.
- Prepare reaction solutions in PCR laminar flow cabinet equipped with UV lamp.
- Use new pair of gloves when purifying DNA and preparing mixtures and solutions.
- Use reagents designed specifically for PCR. Use pipette tips with integrated aerosol filter when preparing DNA samples and reaction solutions.
- For verification of the absence of contamination, prepare a mixture sample without DNA template (negative control).

Recommendations for primer selection

For primer design, use well-established programs like Primer3 <http://biotools.umassmed.edu/bioapps/primer3> www.cgi and follow the basic principles:

- Primer length usually falls in the range of 18 – 30 nucleotides.
- Melting temperature is 58-65 °C. Difference in melting temperatures (T_m) of the two primers shouldn't exceed 3 °C.
- Recommended length of amplicon for real-time PCR is 50 – 200 bp.
- Optimal GC composition of the primers is 40 – 60%. Theoretically, G and C nucleotides should be evenly distributed over the whole length or primer.
- Avoid the presence of ≥3 G or C nucleotides at the 3' terminus of primer in order to prevent risk of nonspecific annealing.
- If possible, primer should end with G or C nucleotide at 3' end.
- Avoid using primers with self-complementary regions, primers complementary to each other and primer repeats for preventing formation of hairpin structures and primer-dimers.
- Make sure that there are no unwanted complementary regions between primers and DNA template.
- Check your primers using BLAST.

Components of reaction solution

DNA template

Optimal amount of DNA template per 50 µl reaction solution is 0.01 – 1 ng in case of using plasmid or phage DNA and 0.1 – 1 µg in case of genomic DNA. Higher amounts of template increase the risk of formation of non-specific amplification products, low amounts of template reduce accuracy of amplification. All conventional techniques of DNA purification can be applied for the preparation of a studied sample. It should be mentioned that trace amounts of certain agents used for the isolation and purification of DNA, such as phenol, EDTA and proteinase K, can

inhibit

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DNA polymerase. Precipitation and repeated washing with 70% ethanol usually removes trace contaminants from DNA sample.

Primers

Recommended concentrations of PCR primers are in the range of 0.1 – 0.6 μ M. Excessive concentration of primers increases the chance of non-specific binding to the template and formation of alternative PCR products.

For degenerated primers and primers utilized for PCR of long fragments, we recommend using higher concentrations in the range of 0.3 – 1 μ M.

Mg²⁺ concentration

Change in concentration of Mg²⁺ ions can have a significant impact on PCR efficiency and specificity. These ions are necessary for the performance of *Taq* DNA polymerase. They also bind with deoxyribonucleotides in the ratio 2:1. Therefore, additional optimization of Mg²⁺ concentration may be necessary if dNTP concentration is altered in the reaction solution. The recommended concentration for Mg²⁺ is 1-5 mM. In case if Mg²⁺ concentration is too small, the yield of PCR product will be reduced. On the other hand, the formation of non-specific products and decreased PCR specificity can take place at high concentration of Mg²⁺ is observed the appearance of nonspecific PCR products and decrease precision. If DNA sample DNA contains EDTA or other substances, chelating metals, concentration of Mg²⁺ should be increased proportionally in PCR mixture (binding to EDTA occurs at the ratio 1:1).

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Characteristics of amplification steps

Initial DNA denaturation and enzyme activation

It is very important to achieve complete denaturation of DNA template at the beginning of PCR which provides its efficient use in the first amplification cycle. If GC composition of the template is 50% or less, initial denaturation at 95 °C for 1-3 min will be enough.

Denaturation

Standard time of denaturation per cycle is considered to be 30 seconds at 95 °C. For GC-rich DNA templates this step can be extended to 3-4 minutes.

Primer annealing

Annealing temperature for primers should be 5 °C lower than their melting temperature (T_m). Conventional annealing time is 30 sec. In case if accumulation of non-specific PCR products takes place, the annealing temperature should be optimized by stepwise 1-2 °C increase.

Number of cycles

If there is less than 10 copies of DNA template available per reaction, then efficient amplification requires not less than 40 cycles. A total of 25 – 35 cycles is enough for higher amount of template.

Elongation

Optimal efficiency of *Taq* DNA polymerase is observed in the temperature range of 70 – 75 °C. The rate of synthesis by *Taq* DNA polymerase ranges from 30 to 60 bp per second depending on template complexity. In the case of using long templates (>2 kbp), it is recommended to estimate elongation time based on the ratio 1 min/kb.

Final elongation

When the last cycle is finished, it is recommended to incubate PCR solution for further 5 – 15 min at 72 °C for complete synthesis of the products. If PCR product is to be further cloned into TA vector, final elongation should be extended to 30 min in order to achieve maximal efficiency of formation of 3'-dA ends of PCR products.

Ordering Information:

Cat.-no	Description	Amount
Y240	BIO-Star Mastermix (2x) with SybrGreen High-ROX	100 rcs (2,5 ml)
Y240L	BIO-Star Mastermix (2x) with SybrGreen High-ROX	5x100 rcs (12,5 ml)
Y240XL	BIO-Star Mastermix (2x) with SybrGreen High-ROX	10x100 rcs (25 ml)

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