



2×Ultra Sybr qPCR Mix

Product Number: PCM33

Shipping and Storage

-20°C; if used frequently, store at 2-8°C to avoid repeated freezing and thawing.

Kit Components

Component	PCM33 1mL	PCM33 5mL	PCM33 25mL
2×Ultra Sybr qPCR Mix	1ml	5×1ml	5×(5×1 ml)
ddH ₂ O	1ml	5×1ml	5×(5×1 ml)

Product Introduction

The 2×Ultra Sybr qPCR Mix is a premixed system for real-time fluorescence quantitative PCR (SYBR Green I), and the concentration is 2×. It contains Golden Star Taq DNA Polymerase, PCR Buffer, dNTPs, SYBR Green I Fluorescent Dye, and Mg²⁺. The operation is simple and convenient. This product is mainly used for the detection of genomic DNA target sequences and cDNA target sequences after RNA reverse transcription.

This product contains the fluorescent dye SYBR Green I which can bind with all double-strand DNA, so that the product can be used for the detection of different target sequences without the need for the synthesis of specific labeled probes. The Golden Star Taq DNA Polymerase in the mixture is a chemically-modified, new efficient hot-start enzyme that does not have polymerase activity at room temperature which prevents non-specific amplification efficiently, and it is activated by incubation at 95°C for 10 minutes. The combination of a unique PCR buffer system and a hot-start enzyme effectively inhibits non-specific PCR amplification and significantly increases the amplification efficiency of PCR.

This product is suitable for fluorescent qPCR instruments that do not require ROX as a calibration dye, such as Roche LightCycler 480, Roche LightCycler 96, Bio-rad iCycler iQ, iQ5, and CFX96.

Features

1. This product uses a new high-performance hot start enzyme (Golden Star Taq DNA Polymerase) and a unique PCR buffer system. This product significantly improves the PCR amplification efficiency and has high sensitivity and specificity.
2. This product is suitable for quantitative PCR detection and can accurately quantify and detect the target gene.

Note

1. Mix gently before use, avoid foaming, and use after brief centrifugation.
2. This product contains SYBR Green I fluorescent dye. Avoid strong light irradiation when storing this product or preparing PCR reaction solution.
3. Avoid repeated freezing and thawing of this product. Repeated freezing and thawing may comprise product performance.
4. This product cannot be used for qPCR using probes.
5. When preparing the reaction solution, use new or noncontaminated tips and centrifuge tubes to prevent contamination.

Protocol

The following protocol is an example of conventional PCR reaction system and condition. The actual protocol should be improved and optimized based on the template, primer structure and the size of the target.

1. PCR reaction system:

Reagent	50µl	Final Conc.
2×Ultra Sybr qPCR Mix	25µl	1x

Forward Primer, 10 uM	1µl	0.2µM
Reverse Primer, 10 uM	1µl	0.2µM
DNA template	2µl	
ddH ₂ O	Up to 5µl	

Note: 1) Usually 0.2µM of primer concentration gives better results, and the final concentration of primers should be between 0.1 and 1.0µM.

2) Usually the amount of DNA template is 10-100ng for genomic DNA or 1-10ng for cDNA. Template can be gradient diluted to optimize.

3) The recommended reaction volume is 50µl, and the reaction volume can also be scaled up or down according to actual experimental requirements.

2. PCR reaction program:

Note! The pre-denaturation reaction of this product must be completed at 95°C for 10 minutes!

It is recommended to use two-step PCR reaction program. This program uses ABI7500 qPCR machine as an example. If a good result cannot be obtained due to the low T_m of the primers, try a three-step PCR program.

Procedure	Temperature	Time	
Pre-denaturation	95°C	10min	
Denaturation	95°C	15s	} 35-40cycles
Annealing/Extension	60°C	1min	
Melting curve analysis			
	95°C	15s	
	60°C	1min	
	95°C	15s	
	60°C	15s	

Note: 1) The hot-start enzyme used in this product must be pre-denatured at 95°C for 10 minutes to activate the enzyme.

2) The annealing temperature should be between 60-64°C. If there is non-specific reaction, increase the annealing temperature.

3) This program uses qPCR machine as an example. The melting curve analysis should be set according to the procedure recommended by qPCR instrument used.

Optimization of reaction conditions

When optimizing the qPCR reaction conditions, different aspects such as the concentration of the primer, the annealing temperature, and the extension time should be considered, to improve the reaction specificity and amplification efficiency.

1. The experimental system with high reaction specificity and high amplification efficiency should be as the following conditions:

1.1. High specificity: no non-specific amplification such as primer dimers for negative control; No other amplification beyond the target fragment.

1.2. High amplification efficiency: Low Ct value; Amplification efficiency of PCR is high, close to the theoretical value of 100%.

2. Methods of optimization of reaction conditions:

2.1. Primer concentration: Usually 0.2µM of primer concentration gives better results, and the final concentration of primers should be between 0.1 and 1.0µM. To increase the specificity of the reaction, decrease the concentration of the -5'-primer; To increase the amplification efficiency, increase the concentration of the primer.

2.2. Annealing temperature: It is recommended to use two-step PCR, and set the annealing temperature as 60°C. To increase the specificity, increase the annealing temperature, which should be between 60-64°C. If a good result cannot be obtained due to the low T_m of the primers, try a three-step PCR program. The annealing temperature of the three-step PCR program should be between 56°C and 64°C.



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2.3. Extension time: It is recommended to use two-step PCR and set the extension time as 1 minute. To increase amplification efficiency, increase the extension time, or try three-step PCR.

Note! The pre-denaturation reaction of this product must be completed at 95°C for 10 minutes!