

GENETAQ™ Green PCR Master Mix

Store at -20°C

Cat. No.
PGK108-A/B

Pack Size
200 Rxns

Ordering Information

Component	PGK107-A	PGK107-B
GENETAQ Green PCR Master Mix (2X)	4 × 1.25 mL	20 × 1.25 mL
Water, nuclease-free	4 × 1.25 mL	20 × 1.25 mL

Description

Puregene GENETAQ Green PCR Master Mix (2X) is a ready-to-use solution containing Taq DNA polymerase, optimized GENETAQ Green buffer, MgCl₂ and dNTPs. The Master Mix is supplemented with two tracking dyes and a density reagent that allows for direct of the PCR product onto gel. The Dye in the master mix do not interfere with PCR performance & are compatible with downstream application applications such as DNA sequencing, ligation & restriction Digestion. This pre-mixed formulation saves time and reduces contamination due to a reduced number of pipetting steps required for PCR set up. The master mix retains all features of Taq DNA polymerase. It is capable of robust amplification up to 6 kb from genomic DNA and up to 20 kb from viral DNA.

Applications

- Routine PCR amplification of DNA fragments up to 6 kb from genomic DNA and up to 20 kb from viral DNA.
- RT-PCR.
- Genotyping.
- Generation of PCR products for TA cloning.

GENETAQ Green PCR Master Mix (2X) composition

GENETAQ Green DNA polymerase is supplied in 2X GENETAQ Green buffer, dATP, dCTP, dGTP and dTTP, 0.4 mM each, and 4 mM MgCl₂. DreamTaq buffer is a proprietary formulation optimized for robust performance in PCR. It contains a density reagent & two Dyes for monitoring electrophoresis progress: The blue dye migrates with 3-5 kb DNA Fragment in a 1% agarose gel & the yellow dye migrates faster than 10 bp in 1% agarose gel. The dyes have absorption peak at 424 nm & 615 nm.

PROTOCOL

1. Gently vortex and briefly centrifuge GENETAQ Green PCR Master Mix (2X) after thawing.
2. Place a thin-walled PCR tube on ice and add the following components for each 50 µL reaction:

GENETAQ Green PCR Master Mix (2X)	25 µL
Forward primer	0.1-1.0 µM
Reverse primer	0.1-1.0 µM
Template DNA	10 pg – 1 µg
Water, nuclease-free to	50 µL
Total volume	50 µL

3. Gently vortex the samples and spin down.
4. When using a thermal cycler that does not contain a heated lid, overlay the reaction mixture with 25 µL of mineral oil.
5. Perform PCR using the recommended thermal cycling conditions outlined below:

Step	Temp. °C	Time	Number
			of cycles
Initial denaturation	95	1-3 min	1
Denaturation	95	30 s	
Annealing	T _m -5	30 s	25-40
Extension*	72	1 min	
Final extension	72	5-15 min	1

* The recommended extension step is 1 min for PCR products up to 2 kb. For longer products, the extension time should be prolonged by 1 min/kb

GUIDELINES FOR PREVENTING CONTAMINATION OF PCR REACTION

During PCR more than 10 million copies of template DNA are generated. Therefore, care must be taken to avoid contamination with other templates and amplicons that may be present in the laboratory environment. General recommendations to lower the risk of contamination are as follows:

- Prepare your DNA sample, set up the PCR mixture, perform thermal cycling and analyze PCR products in separate areas.
- Set up PCR mixtures in a laminar flow cabinet equipped with an UV lamp.
- Wear fresh gloves for DNA purification and reaction set up.
- Use reagent containers dedicated for PCR. Use positive displacement pipettes, or pipette tips with aerosol filters to prepare DNA samples and perform PCR set up.
- Always perform “no template control” (NTC) reactions to check for contamination.



GENETIX BRAND

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GUIDELINES FOR PRIMER DESIGN

Follow the general recommendations for PCR primer design as outlined below:

- PCR primers are generally 15-30 nucleotides long.
- Differences in melting temperatures (T_m) between the two primers should not exceed 5°C.
- Optimal GC content of the primer is 40-60%. Ideally, C and G nucleotides should be distributed uniformly along the primer.
- Avoid placing more than three G or C nucleotides at the 3'-end to lower the risk of non-specific priming.
- If possible, the primer should terminate with a G or C at the 3'-end.
- Avoid self-complementary primer regions, complementarities between the primers and direct primer repeats to prevent hairpin formation and primer dimerization.
- Check for possible sites of undesired complementary between primers and template DNA.
- When designing degenerate primers, place at least 3 conserved nucleotides at the 3'-end.

Estimation of primer melting temperature

For primers containing less than 25 nucleotides, the approx. melting temperature (T_m) can be calculated using the following equation:

$$T_m = 4(G + C) + 2(A + T),$$

where G, C, A, T represent the number of respective nucleotides in the primer.

If the primer contains more than 25 nucleotides we recommend using specialized computer to account for interactions of adjacent bases, effect of salt concentration, etc.

COMPONENTS OF THE REACTION MIXTURE

Template DNA

Optimal amounts of template DNA for a 50 µL reaction volume are 0.01-1 ng for both plasmid and phage DNA, and 0.1-1 µg for genomic DNA. Higher amounts of template increase the risk of generation of non-specific PCR products. Lower amounts of template reduce the accuracy of the amplification.

All routine DNA purification methods can be used to prepare the template. Trace amounts of certain agents used for DNA purification, such as phenol, EDTA and proteinase K, can inhibit DNA polymerases. Ethanol precipitation and repeated washes of the DNA pellet with 70% ethanol usually removes trace contaminants from DNA samples.

Primers

The recommended concentration range of the PCR primers is 0.1-1 M. Excessive primer concentrations increase the probability of mispriming and generation of non-specific PCR products. For degenerate primers and primers used for long PCR we recommend higher primer concentrations in the range of 0.3-1 µM.

CYCLING PARAMETERS

Initial DNA denaturation and enzyme activation It is essential to completely denature the template DNA at the beginning of PCR to ensure efficient utilization of the template during the first amplification cycle. If the GC content of the template is 50% or less, an initial 1-3 min denaturation at 95°C is sufficient.

Denaturation

A DNA denaturation time of 30 seconds per cycle at 95°C is normally sufficient. For GC-rich DNA templates, this step can be prolonged to 3-4 min.

Primer annealing

The annealing temperature should be 5°C lower than the melting temperature (T_m) of the primers. Annealing for 30 seconds is normally sufficient. If non-specific PCR products appear, the annealing temperature should be optimized stepwise in 1-2°C increments.

Extension

The optimal extension temperature for Taq DNA polymerase is 70-75°C. The recommended extension step is 1 min/kb at 72°C for PCR products up to 2 kb. For larger products, the extension time should be prolonged by 1 min/kb. For amplification of longer templates (>6 kb) a reduction of the extension temperature to 68°C is required to avoid enzyme inactivation during prolonged extension times.

Number of cycles

If less than 10 copies of the template are present in the reaction, about 40 cycles are required. For higher template amounts, 25-35 cycles are sufficient.

Final extension

After the last cycle, it is recommended to incubate the PCR mixture at 72°C for an additional 5-15 min to fill-in any possible incomplete reaction products. If the PCR product has to be cloned into a TA vector, the final extension step may be prolonged to 30 min to ensure the highest efficiency of 3'-dA tailing of the PCR product. If the PCR product has to be used for cloning the final extension step can be omitted.



CERTIFICATE OF ANALYSIS

Endodeoxyribonuclease Assay

No detectable conversion of covalently closed circular DNA to nicked DNA was observed after incubation of 25 µL of DreamTaq PCR Master Mix (2X) with 1 µg of pUC19 DNA in 50 µL of reaction mixture for 4 hours at 37°C.

Exodeoxyribonuclease Assay

No detectable degradation of lambda DNA/HindIII fragments was observed after incubation of 25 µL of GENETAQ Green PCR Master Mix (2X) with 1 µg of digested DNA in 50 µL of reaction mixture for 4 hours at 37°C.

Ribonuclease Assay

Less than 0.5% of the total radioactivity was released into trichloroacetic acid-soluble fraction after incubation of 25 µL GENETAQ Green PCR Master Mix (2X) with 1 µg of [3H]-RNA in 50 L of reaction mixture for 4 hours at 37°C.

Functional Assay

GENETAQ Green PCR Master Mix (2X) was tested for amplification of 20 kb DNA fragment.

