Medix Biochemica

Product Manual Cat. No: #1101

Taq DNA Polymerase Hot-Start

Description

Taq DNA Polymerase Hot-Start is supplied together with the 10x Taq reaction buffer. The reaction buffer has been specifically designed for optimal PCR performance and polymerase activity. Taq DNA Polymerase Hot-Start can also be used for real-time cycling, when adding a suitable real-time dye or a fluorescent probe.

Applications include standard PCR, real-time-PCR (addition of suitable dye or probe required), primer extension reactions, TA cloning, 3'A-tailing of blunt ends, and screening / high-throughput PCRs.

Kit components

| Component | S pack* | L pack* |
|---------------------------------|-------------|-----------------|
| Taq DNA Polymerase Hot-Start | 1 x 80 µL | 2 x 400 µL |
| 10x Taq reaction buffer | 2 x 1.25 mL | 13 x 1.25 mL |

*Other pack sizes, bulk orders and customization are available upon request.

Storage and shipment

Transport with cool packs. The reagents should be stored at -20°C upon arrival. The reagents are stable until the expiration date if stored correctly.

Reaction master mix set-up

The recommended master mix set-up for a 50 μL reaction volume is shown in the table below.

| Reagent | Volume (µL) | Final concentration |
|--|--------------------------|-----------------------|
| Taq DNA Polymerase Hot- Start (5 U/µL) | 0.25 | 1.25 U/rxn |
| 10x Taq reaction buffer | 5 | 1x |
| ∞Forward primer (10 μM) | 1 | 0.2 μM (0.05–1 μM) |
| ∞Reverse primer (10 μM) | 1 | 0.2 μM (0.05–1 μM) |
| dNTPs (2 nM) | 5 | 200 µM |
| Template / Sample extract | x | <1000 ng* DNA |
| Nuclease-free water | Up to 50 µL final volume | |

Keep all components on ice.

Spin down and mix all solutions carefully before use.

 ∞ Primers should ideally have a GC content of 40–60% typically.

*Suggested template concentration should be about 1 ng - 1000 ng (genomic DNA) or 1 pg - 1 ng (plasmid/viral DNA) per reaction.

Instrument and program set-up

| Cycles | Steps | Temperature | Time |
|--------|-------------------------|-------------|-------------------|
| 1 | Initial denaturation | 95°C | 2 min |
| 25–40 | Denaturation | 95°C | 15 sec |
| | Annealing* | 54–72°C | 30 sec |
| | Extension | 72°C | 1 min /1000 bp |

*Typically, the annealing temperature is about 3–5°C below the calculated melting temperature of the primers used.



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Technical information and support

For technical enquiries or assay development support, please contact us via e-mail at: mdx@medixbiochemica.com.

Additional information and technical resources are available on our website at: info.medixbiochemica.com/resources.



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