

# Data sheet

## Taq DNA Polymerase Master Mix (2X)

Cat. No: P0035  
2x 1.25 mL  
Ready-to-use

### Introduction

**Taq DNA Polymerase Master Mix (2X)** is an optimized ready-to-use master mix that contains all PCR reaction components: dNTPs, PCR buffer,  $Mg^{2+}$  and **Taq DNA polymerase**. Only primers and template need to be added.

The convenient 2x master mix formulation saves time and eliminates the risk of contamination due to a reduced number of pipetting steps required.

### Features

- Ready-to-use
- Adds extra nucleotides (preferentially adenine) without template at 3' ends leaving 3' overhangs PCR fragments. This fact allows the popular TA-cloning or GC cloning.

### Applications

- Design for medium or high throughput applications (e.g. colony screening)
- PCR fragments amplification for TA or GC cloning
- High-throughput PCR

### Unit definition

One unit is defined as the amount of enzyme required to catalyse the incorporation of 10 nanomoles of dNTPs into acid-insoluble material in 30 minutes at 74°C.

### Unit assay conditions

**Enzyme activity is assayed in the following mixture:** 25mM Tris-HCl pH 9.0 at 25°C, 50mM KCl, 2mM  $MgCl_2$ , 0.1mg/mL gelatine, 200  $\mu$ M de dATP, dGTP, dTTP, 100 $\mu$ M[ $\alpha$ 32-P]dCTP (0.05 $\mu$ Ci/nmol) and 12.5  $\mu$ g activated salmon sperm DNA.

### Concentration:

Buffer PCR 2X; dNTPs 0.4 mM each dNTP (dATP, dCTP, dGTP and dTTP); 4 mM  $MgCl_2$ ; Taq DNA polymerase 0.1 U/ $\mu$ L and Glycerol 4%.

### Quality Certifications

- ✓ Functionally tested in PCR.
- ✓ Undetected bacterial DNA (by PCR).
- ✓ Undetectable nucleases activity (endo-, exo, and ribo-).

**Storage:** Store at -20°C.

*(Continued on reverse side)*

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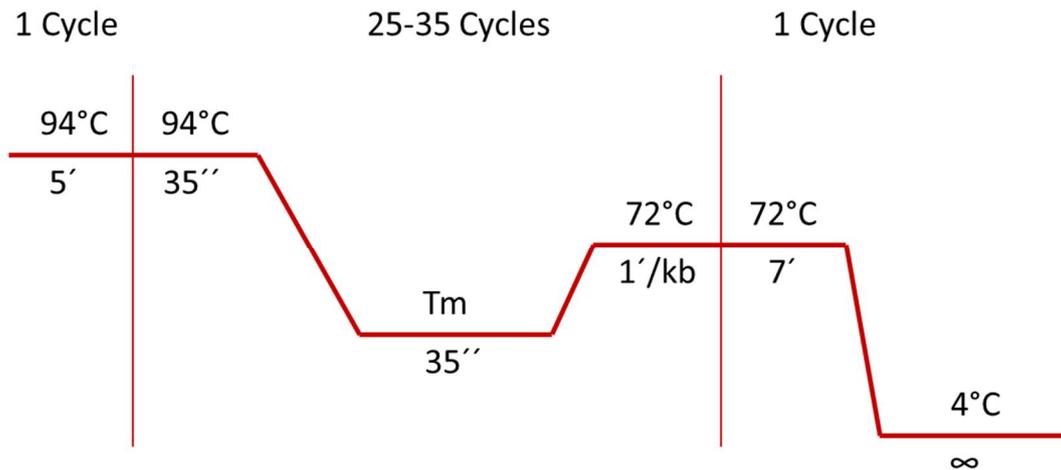
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## Recommended PCR assay (20 µl assay)

The following protocol can be used as a starting point for reaction optimization. The optimal conditions will vary from reaction to reaction and are dependent on the template/primers used.

Taq DNA Polymerase2X Master Mix	10µl (1X)
Forward Primer (15µM)	1µl (0.75 pmol/µL)
Reverse Primer (15µM)	1µl (0.75 pmol/µL)
Template DNA PCR grade H <sub>2</sub> O	Plasmide: 30-75ng; gDNA: 100-500ng up to 20 µl

**Cycling instructions:** 94°C 5:00, 25-30x (94°C 0:35, T<sub>m</sub> 0:35, 72°C 1'/kb), 72°C 7:00, 4°C ∞



## PRODUCT USE LIMITATION

This product is developed, designed and sold exclusively for research purposes and in vitro use only. The product was not tested for use in diagnostics or for drug development, nor is it suitable for administration to humans or animals. Please refer to [www.canvaxbiotech.com](http://www.canvaxbiotech.com) for Material Safety Data Sheet of the product.

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