



# Taq PCR (Protocol for ZymoTaq™ PreMix)

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#### **GUIDELINES**

ZymoTaq™ PreMix is supplied as a 2X concentrated "master mix" containing all the reagents needed to perform "hot start" PCR. The inclusion of a heat-activated, thermal-stable DNA polymerase reduces primer dimer and non-specific product formation that can occur when performing conventional PCR. This unique product is specifically designed for the amplification of bisulfite-treated DNA for methylation detection, realtime and quantitative PCR that are SYBR Green and probe based. The ZymoTaq™ PreMix yields specific amplicon formation with little or no byproducts. Simple and easy to use, just add water, primers, and template DNA to the ZymoTaq™ PreMix and then heat at 95oC for 10 minutes to initiate polymerization.

ZymoTaq™ DNA polymerase is a heat-activated, "hot start" polymerase that has 3'-terminal transferase activity. The addition of "A" overhangs to amplified DNA makes it ideal for use in TA-cloning.

Store ZymoTag™ PreMix at -20°C for up to 12 months. Avoid repeated freeze/thawing of reagents. Prolonged storage is at -80°C

#### **MATERIALS**

NAME Y	CATALOG # V	VENDOR ~
ZymoTaq™ PreMix - 50 rxns	E2003	Zymo Research
ZymoTaq™ PreMix - 200 rxns	E2004	Zymo Research

## MATERIALS TEXT

Reagent	Volume	Final concentration
ZymoTaq™ PreMix	25 μL	1X
Forward Primer (10 µM)	Variable	0.3 to 1 μM
Reverse Primer (10 µM)	Variable	0.3 to 1 μM
Template	Variable	< 200 ng/50 μL
ddH20	to 50 μL	
Total volume	50 μL	

Add all components in a 250  $\mu$ L tube making up to a 25 or 50  $\mu$ l reaction. If performing various PCR with different templates, a Master Mix is recommended to be done.

Reagent	Volume	Final Concentration
ZymoTaq™ PreMix	25 μL	1X
Forward Primer (10 µM)	Variable	0.3 to 1 μM
Reverse Primer (10 µM)	Variable	0.3 to 1 μM
Template	Variable	< 200 ng/50 μl
ddH2O	to 50 μL	
Total Volume	50 μL	

2 Gently mix the PCR reactions and transfer the tubes to a thermocycler. Thermocycling conditions for a routine PCR:

Step	Temperature	Time	Cycles
Initial Denaturation		10 minutes	1
Denaturation	94-96°C	30 sec	
Annealing	Variable	30-40 sec	30-40
Extension	72°C	30 sec - 1 minute*	
Final Extension	72°C	7 minutes	1
Hold	4°C	> 4 minutes	1

<sup>\*</sup> For  $\leq$  1kb. Add an additional 15-30 seconds to the extension time for each kb > 1 kb. Make adjustments to the temperature and/or time if necessary.

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