

FireTaq™ DNA Polymerase

(Catalog # M1506-400; 400 Rxns; Store at -20°C)

11/20

I. Introduction:

Biovision's FireTaq™ DNA Polymerase™ is a strategically-engineered, next generation Taq Polymerase that has rapid expansion rates, robust performance and contains a proprietary antibody that blocks polymerase activity at low temperatures. FireTaq™ Hot start Polymerase allows for a convenient reaction set-up at room temperature without non-specific amplification and primer-dimer formation. With specialized reaction conditions, this polymerase provides increased processivity, yields and sensitivity while shortening reaction times by upto 70%, compared to wild-type Taq DNA Polymerase.

During the initial denaturation step, the antibody dissociates from the DNA polymerase and restores enzyme activity. This feature significantly reduces non-specific product formation that would otherwise compete for reagent availability offering higher specificity and improved yield of PCR products. FireTaq[™] has 5′-3′ polymerase and 5′-3′ exonuclease activities, lacks 3′-5′ exonuclease activity, and produces 3′-dA-tailed amplicons. PCR products made with FireTaq[™] can be used with TA cloning vectors.

II. Contents:

Components	M1506-400	Part Number
FireTaq™ DNA Polymerase	200 μl (400 rxn)	M1506-400-1
5X FireTaq™ Buffer*	2 x 1.0 ml	M1506-400-2

Buffer contains * 1.5 mM Mg²⁺

III. Key Features:

- · reduces non-specific product formation and primer-dimer formation
- Hot start DNA Polymerase
- Superior Performance
- High Sensitivity

IV. Shipping and Storage Conditions:

The kit is shipped in dry ice. All the components of the kit should be stored at -20 °C.

V. Protocol:

1. Mix individual components before use.

Component	Volume	
5X FireTaq™ Buffer	5 μl	
dNTP Mix (10 mM)	0.5 µl	
Forward Primer (10 μM)	1 µl	
Reverse Primer (10 µM)	1 µl	
Template DNA	Variable (100 ng genomic DNA)	
FireTaq™ DNA Polymerase	0.5 μl [#]	
Nuclease-free H₂O	up to 25 μl	

^{#0.5} μl of UpTaq™ DNA Polymerase is recommended for reaction volumes of 25 μl. Increase the volume to 1 μl for difficult targets or crude samples

2. Gently mix the reaction components and briefly centrifuge. Run thermocycling conditions for standard PCR.

Step	Temperature	Duration
Initial Denaturation	95 °C	10 min
25-35 cycles	95 °C	15 sec
	60 °C [†]	15 sec
	72 °C	15 sec/kb
Final Extension	72 °C	1 min

[†] FireTaq™ Buffer allows for primer annealing at 60 °C for most primers, adjust only if needed.

- 3. After PCR, maintain the reaction at 4°C or store at -20°C until use.
- 4. Analyze the amplification products by agarose gel electrophoresis.
- 5. Visualize by ethidium bromide or Safe Image™ Basic DNA Stain (Cat. No. M1193) staining.



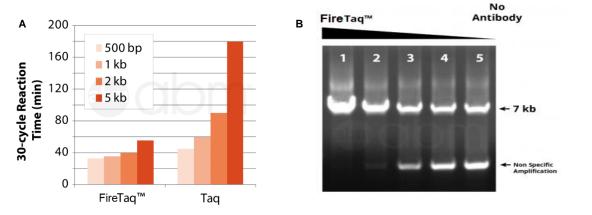


Fig A. FireTaq™ DNA Polymerase has extension speeds of 6 kb/min. Firetaq™ DNA Polymerase was used to amplify different size amplicons (500 bp, 1 kb, 2 kb and 5kb). Total reaction times are listed based on a 30-cycle program using the recommended reaction protocol for each enzyme. Fig. B. FireTaq™ DNA Polymerase eliminates non-specific amplification. FireTaq™ DNA Polymerase was used to amplify 7 kb target. The antibody concentration was decreased incrementally from Lane 1-5. Lane 1 is our FireTaq™ DNA Polymerase formulation and lane 5 does not contain antibody.

VI. Related Products:

BioVision Product Name	Cat. No.	Sizes
ExpressTaq [™] DNA Polymerase	M1504	400 Rxns
HiFidelity [™] DNA Polymerase	M1505	400 Rxns
Taq DNA Polymerase	9001	500, 2500 units
PFU DNA Polymerase	9003	500, 2500 units
Laq™ DNA Polymerase	9004	500, 2500 units

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