

## Unmodified Taq-*FORCE*<sup>™</sup> DNA Polymerase

Product No.: BTAQ0500		BTAQ2500	
<i>System Includes:</i>		<i>System Includes:</i>	
1 x 500 U	Unmodified Taq- <i>FORCE</i> <sup>™</sup> DNA Polymerase (5 U/μl)	5 x 500 U	Unmodified Taq- <i>FORCE</i> <sup>™</sup> DNA Polymerase (5 U/μl)
1 x 1.2 ml	10X NH <sub>4</sub> Reaction Buffer	5 x 1.2 ml	10X NH <sub>4</sub> Reaction Buffer
1 x 1.2 ml	MgCl <sub>2</sub>	5 x 1.2 ml	MgCl <sub>2</sub>

## Unmodified Taq-*FORCE*<sup>™</sup> DNA Polymerase (High Concentration)

Product No.: BTAQ0250H	
<i>System Includes:</i>	
5 x 500 U	Unmodified Taq- <i>FORCE</i> <sup>™</sup> DNA Polymerase (15 U/μl)
5 x 1.2 ml	10X NH <sub>4</sub> Reaction Buffer
5 x 1.2 ml	MgCl <sub>2</sub>

**Unmodified Taq-*FORCE*<sup>™</sup> DNA Polymerase** is the native enzyme isolated from *Thermus aquaticus* YT-1 as described (Kaledin, A.S., et al. (1980) *Biokhimiia* 45:644-651). Unmodified Taq-*FORCE*<sup>™</sup> DNA polymerase produces products that have an A overhang, and are suitable for cloning into T-vectors.

**Unit Definition:** One unit is the amount of enzyme that will incorporate 10 nmoles of dNTPs into an acid-insoluble form in 30 minutes at 72°C under the following assay conditions: 25 mM TAPS, pH 9.3 (25°C); 50 mM KCl; 2 mM MgCl<sub>2</sub>; 0.2 mM each dATP, dGTP, dTTP and 0.1 mM radiolabeled dCTP; 0.25 mg/ml activated salmon sperm DNA; 1 mM β-mercaptoethanol.

**Storage Buffer:** Enzyme is supplied in 20 mM Tris-HCl (pH 7.5), 100 mM NaCl, 0.1 mM EDTA, 2 mM DTT, 50% glycerol, 0.1% Tween-20.

**Storage Conditions:** -20°C. *DO NOT STORE IN A FROST-FREE FREEZER.*

**Quality Control:** Endonuclease, nickase, or exonuclease activities were not detected after 8 hours incubation, respectively, of 1 μg each of lambda, pBR322, or *Hind* III - digested lambda DNA at 72°C in the presence of 5 units of Unmodified Taq-*FORCE*<sup>™</sup> DNA Polymerase.

**10X NH<sub>4</sub> Reaction Buffer:** 160 mM (NH<sub>4</sub>)<sub>2</sub>SO<sub>4</sub>, 670 mM Tris-HCl (pH 8.8 at 25°C), 0.1% Tween-20.

**Mg<sup>++</sup> Stock Solution:** 50 mM MgCl<sub>2</sub>

**Reaction Conditions:** The optimal conditions (incubation time, temperatures, concentration of enzyme, template DNA, primers, MgCl<sub>2</sub>) depend on the system and must be determined individually. **IMPORTANT: Spin vials briefly before use.**

<u>Component</u>	<u>Volume</u>	<u>Final Concentration</u>
10X NH <sub>4</sub> Reaction Buffer	5 μl	1X
DNTPs Pre-Mixed (Cat. #DNTP10)	4 μl	0.2 mM
MgCl <sub>2</sub>	variable	0.5 – 4 mM
Primer	variable	0.1 – 1.0 μM (each)
Unmodified Taq- <i>FORCE</i> <sup>™</sup> DNA Polymerase	variable	0.01 – 0.05 U/μl
Template DNA	variable	variable
Sterile H <sub>2</sub> O	variable	-----
Final Volume	50 μl	-----

FOR RESEARCH USE ONLY

Note: Some applications in which this product can be used may be covered by patents issued and applicable in the United States and certain other countries. Purchase of this product does not convey a license to perform any patented process.