

## Taq DNA Polymerase with Standard Buffer (+ dNTPs)

<b>Product code</b>	02-001 200 U      02-001-5 200 U x 5																
<b>Size</b>	200 U																
<b>Storage</b>	-20°C																
<b>Concentration</b>	5 U/μl																
<b>Product Description</b>	<i>Thermus aquaticus</i> DNA polymerase (Taq DNA polymerase) was expressed in <i>E. coli</i> in large quantities and highly purified. The enzyme has thermostable DNA polymerase activity and the MW is 94 kDa. This enzyme is suitable for PCR reactions; capable of amplifying DNA with various primers.																
<b>Definition of activity</b>	One unit is defined as the amount of enzyme that can incorporate 10nmols of total dNTPs into an acid-insoluble material in 30 minutes at 74°C when activated salmon sperm DNA was used as template / primer.																
<b>Purity</b>	Greater than 95% purity as determined by SDS-PAGE (CBB staining) (Fig.1) The absence of endonucleases and exonucleases was confirmed.																
<b>PCR Test</b>	Good amplification result was obtained in PCR reaction using λDNA as a template (Fig.2)																
<b>Components</b>	<b>Taq DNA polymerase (5U/μl):</b> 20 mM Tris-HCl (pH 8.0), 100 mM KCl, 0.1 mM EDTA, 1 mM DTT, 50% glycerol, 0.5% Tween20, 0.5% Igepal CA-630 (02-Taq 40μl) <b>10 x Standard Buffer (Taq):</b> 100 mM Tris-HCl (pH 8.3), 500 mM KCl, 15 mM MgCl <sub>2</sub> (02-Tsd 1.0ml) <b>2.5 mM (each) dNTPs</b> (02-Dnt 640 μl)																
<b>Application</b>	<ol style="list-style-type: none"> <li>1. High-throughput PCR</li> <li>2. Colony PCR</li> <li>3. Incorporation of dUTP, dITP, and fluorescence-labeled nucleotides</li> <li>4. Primer extension</li> <li>5. Addition of a single nucleotide (adenosine) at the 3'-blunt ends</li> </ol> <table border="1" style="margin-top: 10px;"> <tr> <td colspan="2"><b>General composition of PCR reaction mixture (total 50μl)</b></td> </tr> <tr> <td>Taq DNA polymerase (5 U/μl) (02-Taq)</td> <td>※0.25 μl</td> </tr> <tr> <td>10x Standard Buffer (Taq) (02-Tsd)</td> <td>5 μl</td> </tr> <tr> <td>2.5mM (each) dNTPs (02-Dnt)</td> <td>4 μl</td> </tr> <tr> <td>Template</td> <td>&lt;500 ng</td> </tr> <tr> <td>Primer 1</td> <td>0.2~1.0 mM (final conc.)</td> </tr> <tr> <td>Primer 2</td> <td>0.2~1.0 mM (final conc.)</td> </tr> <tr> <td>Sterile distilled water</td> <td>up to 50 μl</td> </tr> </table> <p>※ Use of excess amount is not recommended.</p>	<b>General composition of PCR reaction mixture (total 50μl)</b>		Taq DNA polymerase (5 U/μl) (02-Taq)	※0.25 μl	10x Standard Buffer (Taq) (02-Tsd)	5 μl	2.5mM (each) dNTPs (02-Dnt)	4 μl	Template	<500 ng	Primer 1	0.2~1.0 mM (final conc.)	Primer 2	0.2~1.0 mM (final conc.)	Sterile distilled water	up to 50 μl
<b>General composition of PCR reaction mixture (total 50μl)</b>																	
Taq DNA polymerase (5 U/μl) (02-Taq)	※0.25 μl																
10x Standard Buffer (Taq) (02-Tsd)	5 μl																
2.5mM (each) dNTPs (02-Dnt)	4 μl																
Template	<500 ng																
Primer 1	0.2~1.0 mM (final conc.)																
Primer 2	0.2~1.0 mM (final conc.)																
Sterile distilled water	up to 50 μl																
<b>Related product</b>	02-011 Taq DNA polymerase with Standard Buffer (-dNTPs) 02-021 Pfu DNA polymerase with Standard Buffer (+dNTPs)																
Please note: All products are FOR RESEARCH USE ONLY. NOT FOR USE IN DIAGNOSTIC PROCEDURES. NOT FOR MILITARY USE.																	

Data Images: 02-001 Taq DNA Polymerase with Standard Buffer (+ dNTPs)

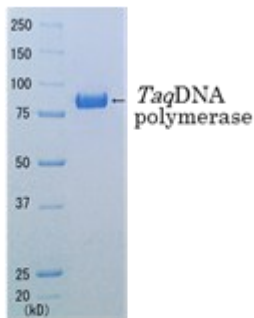
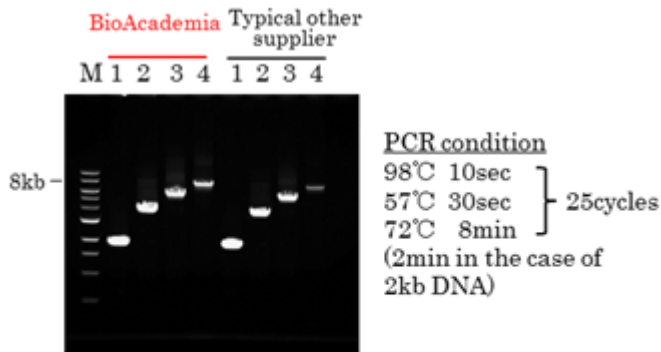


Fig.1 SDS-PAGE of *Taq* DNA polymerase



M : marker,  
 lane 1 : 2 kb, lane 2 : 4 kb,  
 lane 3 : 6 kb, lane 4 : 8 kb.

Fig.2 Amplification of  $\lambda$  DNA