

Taq DNA Polymerase with Standard Buffer (- dNTPs)

Product code	02-011 200 U 02-011-5 200 U x 5																
Size	200 U																
Storage	-20°C																
Concentration	5 U/μl																
Product Description	<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.																
Definition of activity	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.																
Purity	Greater than 95% purity as determined by SDS-PAGE (CBB staining) (Fig.1) The absence of endonucleases and exonucleases was confirmed.																
PCR Test	Good amplification result was obtained in PCR reaction using λDNA as a template (Fig.2)																
Components	Taq DNA polymerase (5U/μl): 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) 10 x Standard Buffer (Taq): 100 mM Tris-HCl (pH 8.3), 500 mM KCl, 15 mM MgCl ₂ (02-Tsd 1.0ml)																
Application	<ol style="list-style-type: none"> 1. High-throughput PCR 2. Colony PCR 3. Incorporation of dUTP, dITP, and fluorescence-labeled nucleotides 4. Primer extension 5. Addition of a single nucleotide (adenosine) at the 3'-blunt ends <table border="1" style="margin-top: 10px;"> <tr> <td colspan="2">General composition of PCR reaction mixture (total 50μl)</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><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>	General composition of PCR reaction mixture (total 50μl)		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
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Related product	02-001 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: 02-011 Taq DNA Polymerase with Standard Buffer (- dNTPs)

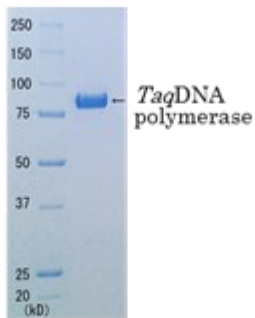
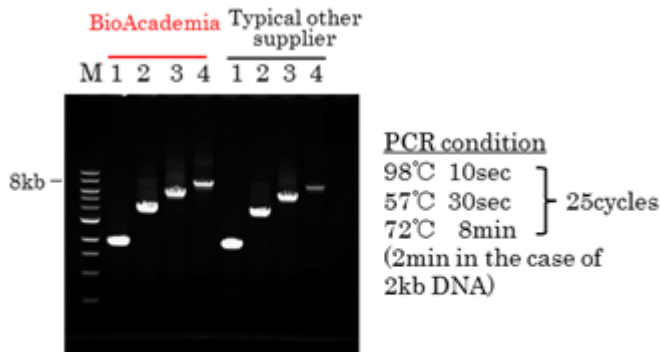


Fig.1 SDS-PAGE of *Taq* DNA polymerase



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

Fig.2 Amplification of λ DNA