



## **ADS™ *Taq* DNA Polymerase User Manual**

### **Product catalog numbers**

Catalog Number	Unit Size	Components
101004	400 units (80 $\mu$ L)	<i>Taq</i> DNA polymerase 400 units, 80 $\mu$ L; 5x PCR Green Buffer 3 mL
101020	2000 units (400 $\mu$ L)	<i>Taq</i> DNA polymerase 2000 units, 400 $\mu$ L; 5x Green Buffer 18 mL

### **Product description**

*Taq* DNA polymerase is frequently used in PCR amplification of DNA fragments. The cost-effective ADS *Taq* DNA polymerase has been routinely used for generation of PCR templates for DNA sequencing.

### **Product storage conditions**

Store the *Taq* DNA polymerase and 5x Green Buffer at -20 °C upon arrival.

### **PCR reaction setup**

Components	25 $\mu$ L reaction	50 $\mu$ L reaction	Final concentration or note
5x Green PCR Buffer	5 $\mu$ L	10 $\mu$ L	Make it 1x Green buffer
10 mM dNTPs	0.5 $\mu$ L	1.0 $\mu$ L	200 $\mu$ M
Primer 1 and 2 (10 $\mu$ M each)	0.5 $\mu$ L each	1.0 $\mu$ L each	0.2 $\mu$ M
DNA template	Varies based on DNA complexity	Varies based on DNA complexity	High-complexity DNA (genomic DNA) at 100 ng to 1 $\mu$ g DNA per 50 $\mu$ L reaction; Low complexity DNA (plasmids) at 1 pg to 10 ng DNA per 50 $\mu$ L reaction
<i>Taq</i> DNA polymerase	0.125 $\mu$ L	0.25 $\mu$ L	1.25 U per 50 $\mu$ L reaction
H <sub>2</sub> O	To final 20 $\mu$ L	To final 50 $\mu$ L	Add water last up to the final volume
Total Volume	20 $\mu$ L	50 $\mu$ L	

Vortex the reaction mixture and do a quick spin in a bench centrifuge. Program and run the PCR reaction before purifying the PCR product using the ADS PCR Cleaning beads.