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Productos
Bio-Lógicos®

Innovación y Desarrollo en Biotecnología

Taq DNA Pol - Long

Cat. no. EC11

Storage: -20℃

Concentration: 2.5 U/μl

Product Size

Product Components	EC1101	EC1102
***Taq* DNA Polymerase Long**	250 U	500 U
10× *Taq* Long Buffer	1.8 ml	1.8 ml
10× *Taq* Long Buffer	1.8 ml	1.8 ml

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http://www.pb-l.com.ar

Introduction

Long*Taq*DNA Polymerase is anindependently developed thermo-stable DNA polymerase with3'-5'exonuclease activity. It possesses high amplification efficiency and high fidelity. Provided with two kinds of buffer, Long *Taq*DNA Polymerasecould amplify varied templates. To simple templates, it's good for 40 kb; to complex templates asGC-rich and repeated sequences,it's good for 15 kb.The PCR products can be used directly in TA-cloning procedures.If required of high cloning efficiency, please purify, add A and then make T/A-cloning.

Unit Definition

One unit of Long TaqDNA Polymerase is defined as the amount that incorporates10 nmol of dNTPs into acid-insoluble material within 30 minat 74°Cwith activated salmon sperm DNA as the template-primer.

Storage Buffer

20mM Tris-HCl(pH8.0),0.1mM EDTA,1mM DTT,100mM KCl,Stabilizers,50% Glycerol.

10×Long *Taq*Buffer

Provided with two kinds of buffer, 10× Long*Taq*Buffer Iand10× Long *Taq*Buffer II. Please use Buffer I first, if it fails to amplify the template, usebuffer II.

Applications

PCR amplification oflongDNA fragments, and complex templates as GC-rich and repeated sequences, *e.g.*,gene map construction, sequencing, molecular geneticsresearch.

Example

Note: The following example only for reference, user must set up optimal reaction system according to different reaction conditions such as different templates or primers *etc*.

1. To 50 μl PCR reaction system: 1 kb fragment of human genomic DNA was amplified (If use different reaction system, please proportionally increase or decrease the amount of reaction components referring to this system).

Template	< 1 μg
Primer 1(10 μM)	1 μl
Primer 2(10 μM)	1 μl
10× *Taq* Platinum Buffer	5 μl
dNTP Mixture(2.5 mM)	4 μl
Long Taq (2.5 U/μl)	0.5-1 μl
ddH_2O	up to 50 μl

2. PCR cycle set-up□

94°C 5 min
94°C 30 sec
55°C 30 sec] 30 cycles
72°C 2 min
72°C 5 min

3. Result detection: Load 5 μl PCR products to agrose gel for PCR detecting.