



# Exonuclease I (Exo I)

**Product Number: EN01**

---

## Shipping and Storage

-20°C.

## Components

Component	EN01	EN01
	4000U	20000U
Exonuclease, 20 U/μl	200μl	5×200μl
10×Exo I Reacting Buffer	1mL	5×1mL

## Description

This product is derived from recombinant E.coli strain, carrying the Exo I gene, with exonuclease activity of hydrolyzing single-stranded DNA from the 3'-5' direction, which can gradually release deoxyribonucleic acid 5' monophosphate and leave a complete 5' end dinucleotide. This product is mainly used to degrade digestion primers after PCR amplification, and has no activity for double-stranded DNA and 3' OH terminal DNA strands closed by phosphoryl or acetyl groups.

## Unit definition

The amount of enzyme required to catalyze the release of 10 nmol of soluble nucleotide within 30 minutes at 37°C was defined as 1 unit of activity (U).

## Quality Control

The purity was higher than 95% after SDS-PAGE electrophoresis and Coomassie bright blue staining. The addition of BSA can ensure the stability of enzyme.

## Protocol

The following is an example of cleaning PCR products before sequencing. The reaction removes single-stranded primers and degrades unpaired nucleotides.

1. Mix PCR products with Exonuclease I as shown in the table below.

Reagent	Volume
PCR Product	4.9μl
Exonuclease I	0.5μl
10×Exo I Reaction Buffer	0.6μl

2. Mix and incubate at 37°C for 30 minutes.
3. 80°C incubation for 20 minutes can deactivate.