

Manual

Exonuclease I

Recombinant enzyme that degrades single-stranded DNA in the 3'-5'. Concentration 5 U/ μ l.

catalog #	size
1020-1	1000 U
1020-5	5000 U

For research use only.

Guarantee

A&A Biotechnology provides guarantee on this product.
The company does not guarantee correct performance of this kit in the event of:

- not adhering to the supplied protocol
- use of not recommended equipment or materials
- use of other reagents than recommended or which are not a component of the product
- use of expired or improperly stored product or its components



Description

Exonuclease I removal of nucleotides from ssDNA in the 3'-5' direction.

Application

- removal of ssDNA with a hydroxyl group at the 3-end
- removing primer residues in the mixture after DNA amplification
- when used simultaneously with alkaline phosphatase, removes primers and nucleotides

Contents

	1020-1	1020-50	storage
Exonuclease I	1000 U	5000 U	-20 °C
storage buffer: 10 mM Tris-HCl, pH 7.5, 100 mM NaCl, 0,5 mM EDTA, 5 mM 2-mercaptoethanol, 100 µg/ml BSA, 50% glycerol (v/v)			
Exonuclease reaction buffer	1.5 ml	5 x 1.5 ml	-20 °C
10x reaction buffer: 500 mM Tris-HCl, pH 8,0, 50 mM MgCl ₂			

Unit definition

1 U of enzyme catalysis the release of 10 nmole of acid-soluble nucleotide in 30 min at 37 °C under standard reaction conditions.

Protocol

1. Thaw and mix all components and add:

component	reaction volume
Exonuclease reaction buffer	10 μ l
Exonuclease I	1 μ l
DNA sample	1-8 μ l
Sterile water	up to 10 μ l

2. Incubate for 30 min at 37 °C.
3. Enzyme inactivation: incubate for 20 min at 80 °C.



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