

Exo-Minus Klenow DNA Polymerase (D355A, E357A)

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Manual

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1. Introduction

Exo-Minus Klenow DNA Polymerase is a DNA-dependent DNA polymerase that lacks both of the $5'\rightarrow 3'$ and $3'\rightarrow 5'$ exonuclease activities of *E. coli* DNA Polymerase I¹ from which it is derived. This N-terminal truncation of DNA Polymerase I has two mutations (D355A and E357A).

2. Product designations and kit components

Product	Kit size	Catalogue number	Reagent description	Part number	Volume
Exo-Minus Klenow DNA Polymerase (D355A, E357A)	1,000 units	KL11101K	Exo-Minus Klenow DNA Polymerase (D355A, E357A) (10 U/μL)	E0141-10D1	100 μL
			Klenow exo- DNA Pol. 10X Buffer	SS000070-D1	5 mL

3. Product specifications

Storage: Store only at -20 °C in a freezer without a defrost cycle.

Storage buffer: Exo-Minus Klenow DNA Polymerase (D355A and E357A) is supplied in a 50% glycerol solution containing 50 mM Tris-HCl (pH 7.5), 100 mM NaCl, 1.0 mM dithiothreitol (DTT), 0.1 mM EDTA and 0.1% Triton X-100 (Rohm & Haas).

Specific activity: Exo-Minus Klenow DNA Polymerase has a specific activity of >1 × 10⁴ units/mg. **Unit definition:** One unit converts 10 nmol of dNTPs into acid-insoluble material in 30 minutes at 37 °C under standard assay conditions.

10X Reaction Buffer: 0.2 M Tris-HCl (pH 7.5), 50 mM MgCl₂ and 5.0 mM DTT.

Quality Control: Exo-Minus Klenow DNA Polymerase (D355A and E357A) is function-tested in a reaction containing 20 mM Tris-HCl (pH 7.5), 5 mM MgCl $_2$, 0.5 mM DTT, 10 μ g of denatured activated calf thymus DNA, 2.5 μ M each dNTP and varying amounts of Exo-Minus Klenow DNA Polymerase.

Contaminating activity assays: Exo-Minus Klenow DNA Polymerase (D355A and E357A) is free of detectable endo- and exonuclease and RNase activities.

4. Reference

1. Lehman IR (1981) The Enzymes Academic Press 14, 16.

5. Further support

If you require any further support, please do not hesitate to contact our Technical Support Team: techsupport@lgcgroup.com.



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