

Uracil DNA Glycosylase (UDG)

Shipping: On Dry/Blue Ice

Catalog number

Batch No.: See vial

BIO-27044: 500u

Store at -20°C

meridian BIOSCIENCE™

Storage and stability:

Uracil DNA Glycosylase is shipped on dry/blue ice. On arrival store at -20 °C for optimum stability.

Expiry:

When stored under the recommended conditions and handled correctly, full activity of the kit is retained until the expiry date on the outer box label.

Unit Definition: One unit is the amount of enzyme that catalyses the release of 60 pmol of uracil per minute from double stranded, uracil-containing DNA. Activity was measured by release of [³H]-uracil in a 50 mL reaction mix containing 0.2 mg DNA (10⁴ -10⁵ cpm/mg) in 30 minutes at 37 °C. Safety precautions: Please refer to the material safety data sheet for further information.

Notes:

For research or further manufacturing use only.

Description

Uracil DNA Glycosylase (UDG) catalyzes the release of uracil from uracil-containing single or double-stranded DNA, but not from RNA or oligonucleotides (6 or fewer bases). UDG is active over a broad pH range with an optimum at pH 8.0, does not require a divalent cation, and is inhibited by high ionic strength (>200 mM).

Meridian UDG is purified to SDS-PAGE purity and is free of endonucleases, exonucleases, nickases and RNases.

Source: Recombinant *E. coli* strain carrying the over-expressed modified gene of Uracil DNA Glycosylase from *E. coli*.

Features

- Removal of uracil from uracil-containing DNA
- Novel temperature sensitive mutant irreversibly inactive after heating

Applications

- UDG treatment of uracil-containing DNA to prevent amplification by DNA Polymerases
- Used to investigate features of protein-DNA interactions

Components

Product Name	500 Units
Uracil DNA Glycosylase	500 µL
10x Reaction Buffer	1.2 mL

Associated Products

Product Name	Pack Size	Cat. No.
dUTP Mix 50 mM	25 µmol (1 x 500 µL)	BIO-39041
dUTP 100 mM	25 µmol (1 x 500 µL)	BIO-39035
IMMOLASE	250 Units	BIO-21046

Reagent Specifications:

10x Reaction Buffer (1 mL): 0.25 M Tris-Cl, 1 mM EDTA, 10 mM DTT, pH 8.0.

Citations:

1. Rashtchian, A., *et al.* PCR methods and applications **2(2)**, 124-130 (1992).
2. Stivers, J.T. & Drohat, A.C. Archives of Biochemistry and Biophysics **396(1)**, 1-9 (2001).

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