

## Quick-Start Protocol

# UDGase

UDGase (*Escherichia coli* Uracil DNA Glycosylase) (cat. nos. EN19-050 and EN19-250) catalyzes the hydrolysis of the N-glycosylic bond between uracil and sugar, leaving an apyrimidinic site in uracil-containing single-stranded or double-stranded DNA. The enzyme shows no activity on RNA or oligonucleotides. UDG should be shipped on dry ice. All components should be stored at  $-20^{\circ}\text{C}$  in a freezer without a defrost cycle.

### Further information

- Safety Data Sheets: [www.qiagen.com/safety](http://www.qiagen.com/safety)
- Technical assistance: [support.qiagen.com](http://support.qiagen.com)

### Notes before starting

- One unit is defined as the amount of enzyme that catalyzes the release of 60 pmol of uracil per minute from uracil-containing dsDNA. Activity is measured by release of [ $^3\text{H}$ ]-uracil in a 50  $\mu\text{l}$  reaction containing 0.2  $\mu\text{g}$  DNA in 30 minutes at  $37^{\circ}\text{C}$ .
- Active over a broad pH range (optimum at pH 8.0).
- The enzyme can be irreversibly inactivated by incubation at  $95^{\circ}\text{C}$  for 10 minutes.

## Procedure

### For removing the dUTP-containing DNA template from the PCR reaction

1. Add 1 U of UDGase directly into the PCR reaction mixture (usually 1  $\mu$ L of enzyme/ 20–25  $\mu$ L reaction volume).
2. Incubate for 10–30 min at 37°C.
3. Inactivate UDG at 95°C for 10 min.
4. Continue to the next round of PCR/qPCR reaction.

**Optional:** For digesting dUTP-DNA in a separate reaction, dilute 1-5  $\mu$ L of dUTP-containing DNA sample in 20  $\mu$ L of 1xUDGase Reaction Buffer and add 1 U of UDGase.

## Document Revision History

Date	Changes
08/2023	Initial release

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