

## DATA SHEET



## Uracil-DNA Glycosylase (UDG)

Prevention of carry-over contaminations  
 UNG, UDG

Cat. Nº.	Amount
<input checked="" type="checkbox"/> ENZ-125S	100 units
<input type="checkbox"/> ENZ-125M	200 units
<input type="checkbox"/> ENZ-125L	1.000 units

**Unit Definition:** One unit of enzyme catalyzes the degradation of 1 µg single-stranded uracil-containing DNA at 37 °C in 60 min.

### Storage Buffer

The enzyme is supplied in: 20 mM Tris-HCl (pH 8.0), 50 mM NaCl, 1 mM EDTA, 1 mM DTT, 50 µg/ml BSA and 50% (v/v) glycerol.

### Concentration:

1 unit/µL

**For *in vitro* use only!**

### Shelf Life:

12 months

### Shipping:

Shipped on blue ice

### Storage Conditions:

Store at -20 °C

### Additional Storage Conditions:

Avoid freeze/thaw cycles.

### Description:

Uracil-DNA-glycosylase (UDG), also known as UNG is an engineered *E. coli* enzyme that catalyzes the release of uracil from single and double stranded uracil-containing DNA. The resulting abasic sites are susceptible to hydrolytic cleavage at elevated temperatures. An amount of 0.1 units UDG can completely destroy up to 200 ng dU-containing DNA in 2 min at 50°C. It is used in real-time PCR to prevent carry-over contamination of dU-containing DNA from previous reactions.

This enzyme is designed to be more susceptible to high-temperature and inactivated easily during the dUTP containing real time-PCR.

### Recommended assay:

Add 0.2 µl (0.2 units) UDG for each 50 µl of master mix and vortex thoroughly. The preparation of a master mix is crucial in quantitative PCR reactions to reduce pipetting errors. An UDG treatment of 2 min at 50°C at the onset of thermal cycling removes uracil residues from dU-containing DNA and prevents it from serving as template. UNG is easily heat-inactivated at temperatures above 65°C in the following initial denaturation step of the PCR.