

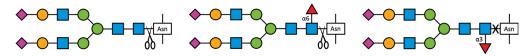
PNGase F Product Info Rev. 2023-02-23

## **PNGase F contents**

Catalog #	Description	Size	M. W.	Purity	рН	Storage
GE0101	PNGase F	4,000 units, lyophilized	37,270	> 95%	7.5-8.5 optimal	-20°C, up to 12 months
BA0501	10X Reaction Buffer 1	1 mL			7.5	4 to 25°C

This product is for research use only and not for resale or for any use in the manufacture of a therapeutic or for any diagnostic purpose.

**Product description:** This product is recombinant PNGase F (Peptide:N-Glycosidase F, EC #3.5.1.52, CAS #83534-39-8), cloned from *Elizabethkingia meningosepticum* and expressed in *Escherichia coli* with an N-terminal 6xHis tag. It catalyzes the cleavage of asparagine-linked oligosaccharides, except those containing an  $\alpha$ 1,3-linked core fucose, from glycoproteins and glycopeptides.



This product does not contain any detectable activities of proteases or other glycosidases.

**Unit definition:** The amount of PNGase F required to deglycosylate 1 nanomole (15  $\mu$ g) of denatured RNase B in 1 h at 37°C in 25  $\mu$ L 1X Reaction Buffer 1 (20 mM Tris, 50 mM NaCl, 1 mM EDTA, pH 7.5).

**Product reconstitution:** Dissolve the lyophilized product in 100  $\mu$ L molecular grade water to make a 40,000 units/mL (Cat #GE0101-4KU) or a 200,000 units/mL (Cat #GE0101-20KU) solution in 1X Reaction Buffer 1. Once reconstituted, store at 4°C for up to 10 days or -20°C for up to 6 months. Aliquoting is recommended to avoid repeated freeze-thaw cycles.

## Suggested protocol for protein deglycosylation:

- 1. Glycoprotein substrate denaturation:
  - 1.1 Mix the following components in a microfuge tube:

Glycoprotein (e.g., RNase B; user supplied) 50-500  $\mu$ g 1% SDS (user supplied) 10.0  $\mu$ L 0.5 M  $\beta$ -Mercaptoethanol or DTT (user supplied) 10.0  $\mu$ L 10X Reaction Buffer 1 (Cat #BA0501) 10.0  $\mu$ L to 100  $\mu$ L to 100  $\mu$ L final volume

1.2 Heat at 98°C for 10 min. Cool to room temperature.

- 2. PNGase F digestion:
  - 2.1 Mix the following components in a microfuge tube:

 $\begin{array}{ll} Denatured \ glycoprotein \ substrate & 2-15 \ \mu g \\ 10\% \ Triton \ X-100 \ (user \ supplied) & 2.0 \ \mu L \\ 10X \ Reaction \ Buffer \ 1 \ (Cat \ \#BA0501) & 2.5 \ \mu L \end{array}$ 

PNGase F (Cat #GE0101-4KU or GE0101-20KU) 1.0  $\mu$ L (40 or 200 units) Molecular grade water to 25  $\mu$ L final volume

- 2.2 Incubate at 37°C for 1 h.
- 2.3 Analyze by SDS-PAGE mobility shift or other method to determine the extent of deglycosylation.

**Reference:** Loo T, et al. Protein Expr Purif. 2002 Feb;24(1):90-8. PMID: 11812228.

**Note:** Reactions may be scaled up to accommodate larger amount and volume of substrate. Titration of the amount of enzyme in a reaction is recommended for each new substrate. PNGase F may remove N-glycans from native glycoproteins at higher enzyme concentration and longer incubation time.