



PNGase F (Peptide-N-Glycosidase F)

Peptide-N4-(acetyl- β -glucosaminy1)-asparagine amidase,
N-Glycosidase F

Source

recombinant gene from *Elizabethkingia miricola* in *E. Coli*

Catalog Number

E-RPNG01	60 μ l	0.3 U
E-RPNG01-20	20 μ l	0.1 U
E-RPNG01-200	200 μ l	1.0 U

EC 3.5.1.52

Recommended Reagents

included with E-RPNG01:

- 1 vial: 5x Reaction Buffer pH 7 - 400 μ l
- 1 vial: Denaturation Solution - 200 μ l
2% SDS/ 1 M β -mercaptoethanol
- 1 vial: 15% Triton X-100 - 200 μ l

Specific Activity \geq 25 U/mg

Activity 5 U/ml

Formulation

The enzyme is provided as a sterile-filtered solution in
20 mM Tris-HCl (pH 7.5).

Molecular Weight approximately 35 kD.

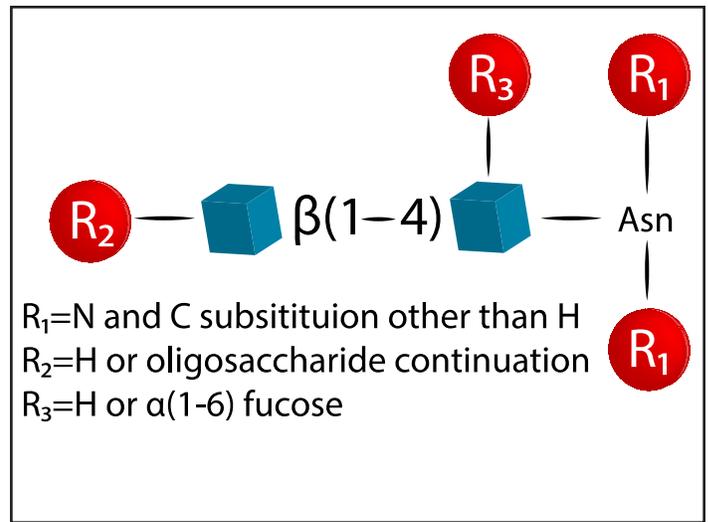
pH optimum: 7.5, active over the range 6-10.

Storage

Store enzyme at 4°C. Do not freeze.

Specific Activity

One unit of PNGase F activity is defined as the amount of enzyme required to catalyze the release of N-linked oligosaccharides from 1 micromole of RNase B in 1 minute at 37°C, pH 7.5. Cleavage is monitored by SDS-PAGE (cleaved RNase B migrates faster).



Specificity

QA-Bio™ PNGase F cleaves asparagine-linked (N-linked) oligosaccharides from glycoproteins. PNGase F deaminates asparagine to aspartic acid, leaving the oligosaccharides intact.

Denaturation increases the rate of cleavage up to 100x. Most native proteins can still be completely N-deglycosylated but incubation time must be increased. PNGase F will remain active under incubation conditions for at least 72 hours.

PNGase F will not remove oligosaccharides containing Alpha-(1,3)-linked core fucose commonly found on plant glycoproteins; for this purpose, use peptide N-glycosidase A.

Stability

Several days exposure to ambient temperatures will not reduce activity. Stable at least 12 months when stored properly.

Quality & Purity

QA-Bio PNGase F is tested for contaminating protease as follows: 10 μ g of denatured BSA is incubated at 37°C for 24 hours with 2 μ l of enzyme. SDS-PAGE analysis of the treated BSA shows no evidence of degradation.

The production host strain has been extensively tested and does not produce any detectable glycosidases.

Directions for use

1. Add up to 200µg of glycoprotein to an Eppendorf tube. Adjust to 35 µl final volume with de-ionized water.
2. Add 10 µl 5x Reaction Buffer 7.5 and 2.5 µl of Denaturation Solution. Heat at 100°C for 5 minutes.
3. Cool. Add 2.5 µl of Triton X-100 and mix.
NOTE: Failure to add Triton X-100 will result in a 3-fold reduction of PNGase F activity.
4. Add 2.0 µl of PNGase F to the reaction. Incubate 3 hours at 37°C.

If SDS or heat denaturation is omitted, increase incubation time to at least 24 hours.

Monitor cleavage by SDS-PAGE.

References:

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Preparation of deglycosylated egg white avidin. Appl Biochem Biotech 53: 1-9 (1995)
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- Tarentino A.L. and T.H. Plummer. Enzymatic deglycosylation of asparagine-linked glycans: purification, properties, and specificity of oligosaccharide-cleaving enzymes from *Flavobacterium meningosepticum*. Meth Enzymol 230:44-57 (1994)
- Trimble R.B. and A.L. Tarentino. Identification of distinct endoglycosidase (endo) activities in *Flavobacterium meningosepticum*: endo F1, endo F2 and endo F3. Endo F1 and endo H hydrolyze only high mannose and hybrid glycans. J Biol Chem 266:1646-1651 (1991).
- Taga, E. M., A. Waheed and R. L. Van Etten. Structural and chemical characterization of a homogeneous peptide N-glycosidase from almond. Biochemistry 23:815-22 (1984).

Warranties and liabilities

QA-Bio, Inc warrants that the above product conforms to the specifications described herein. Should the product fail for reasons other than through misuse QA-Bio will, at its option, replace free of charge or refund the purchase price. This warranty is exclusive and QA-Bio makes no other warrants, expressed or implied, including any implied conditions or warranties of merchantability or fitness for any particular purpose. QA-Bio shall not be liable for any incidental, consequential or contingent damages.

This product is intended for *in vitro* research only.

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