

#### **Product Details**

PNGase F is the most effective enzymatic method for removing almost all N-linked oligosaccharides from glycoproteins. PNGase F is an amidase, which cleaves between the innermost GlcNAc and asparagine residues of high mannose, hybrid, and complex oligosaccharides.

# **Application**

- Leaves N-glycan core oligosaccharides intact and suitable for further analysis
- Non-recombinant with no detectable endoglycosidase F1, F2 or F3 contamination

#### **Unit Definition**

One unit is defined as the amount of enzyme required to remove > 95% of the carbohydrate from 10  $\mu g$  of denatured RNase B in 1 hour at 37°C in a total reaction volume of 20  $\mu l$ .

## **Quility Control**

Less than 1.0 EU per µg by the LAL method.

### **Purity**

>90% as determined by SDS-PAGE.

## Formulation

Supplied as  $0.2 \mu m$  filtered solution in 20 Mm Tris,  $50 \mu m$  NaCl,  $5 \mu m$  EDTA, pH7.5 with glycerol as protectant.

Contact us for customized product form or formulation.

## **Shipping**

This product is supplied and shipped with dry ice, please inquire the shipping cost.

## Storage

This product is stable after storage at:

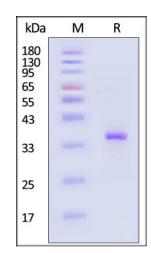
- The product MUST be stored at -70°C or lower upon receipt.
- -70°C for 12 months under sterile conditions.

#### **Notes**

To deglycosylate a native glycoprotein, longer incubation time as well as more enzyme may be required.

PNGase F will not cleave N-linked glycans containing core α1-3 Fucose.

### **SDS-PAGE**



PNGase F (500U/ul) on SDS-PAGE under reducing (R) condition. The gel was stained with Coomassie Blue. The purity of the protein is greater than 90% (With <u>Star Ribbon Pre-stained Protein Marker</u>).

### Clinical and Translational Updates

Please contact us via TechSupport@acrobiosystems.com if you have any question on this product.