

Endo S Product Info Rev. 2023-04-19

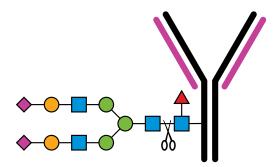
## **Endo S contents**

Catalog #	Description	Size	M. W.	Purity	рН	Storage
GE0801	Endo S	2,500 units, lyophilized	104,088	> 95%	5.5-6.5 optimal	-20°C, up to 12 months
BA1001	10X Reaction Buffer 6	1 mL			5.5	4 to 25°C
GC0101 (optional)	Human Ig	0.1 mL (1 mg/mL)	150,000			-20°C, up to 12 months

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 Endo- $\beta$ -N-Acetylglucosaminidase F2 (Endo S; glycosyl hydrolase family 18, EC #3.2.1.96), cloned from *Streptococcus pyogenes* and expressed in *Escherichia coli* with an N-terminal 8xHis tag. The 8xHis tag may be removed by digestion with FasTEV<sup>TM</sup> (Cat #GE0501), a TEV protease with enhanced stability and catalytic activity.

Endo S catalyzes the cleavage of *N*-linked glycans from IgG heavy chain, as illustrated below.



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

**Unit definition:** One unit is defined as the amount of Endo S required to deglycosylate > 90% of 2.5  $\mu$ g of human IgG in 1 h at 37°C in 25  $\mu$ L 1X Reaction Buffer 6 (50 mM NaOAc, 5 mM CaCl<sub>2</sub>, pH 5.5).

**Product reconstitution:** Dissolve the lyophilized Endo S in 100  $\mu$ L molecular grade water to make a 25,000 units/mL (Cat #GE0801) solution in enzyme storage buffer (20 mM Tris-HCl, 50 mM NaCl, 1 mM EDTA, pH 7.5). Once reconstituted, store at 4°C for up to 7 days or -20°C for up to 3 months. Aliquoting is recommended to avoid repeated freeze-thaw cycles. *The optional control substrate (Cat #GC0101) is immunoglobulin (Ig) purified from human serum by Protein A affinity chromatography.* 

## **Activity assay:**

Endo S digestion:

1. Mix the following components in a microfuge tube:

Human Ig substrate (Cat #GC0101) 2.5  $\mu$ g 10X Reaction Buffer 6 (Cat #BA1001) 2.5  $\mu$ L Endo S (Cat #GE0801) 1.0-25 units Molecular grade water to 25  $\mu$ L final volume

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

Reference: Trastoy B, et al. Proc Natl Acad Sci U S A. 2014 May 6;111(18):6714-9. PMID: 24753590.