



Product Information & Manual

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IdeS Protease

Cat no. LDG0039RG

Product Overview

Description

IdeS Protease is a highly specific cysteine protease derived from *Streptococcus pyogenes* that cleaves human IgG antibodies at a single site in the lower hinge region. This enzyme is essential for therapeutic antibody and Fc fusion protein characterization, enabling the rapid generation of homogenous F(ab')₂ and Fc fragments for detailed analysis via mass spectrometry or chromatography.

Expression system

Escherichia coli

Unit definition

One unit is defined as the amount of enzyme required to cleave $\geq 95\%$ of 1 μ g recombinant human IgG in 30 minutes at 37°C.

Storage buffer

Lyophilized from 50 mM sodium phosphate, 150 mM NaCl, pH 6.6.

Reconstitution

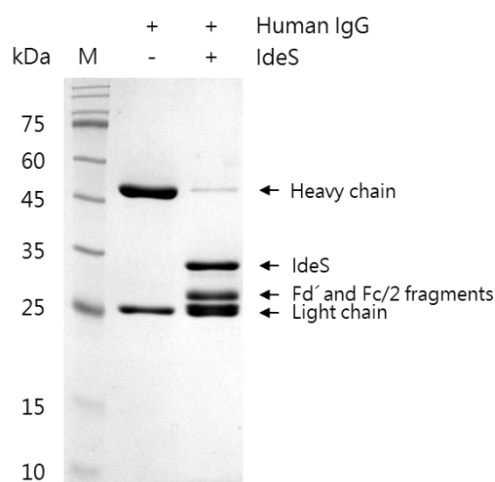
It is recommended to reconstitute the lyophilized protein by adding 36 μ L of sterile H₂O to the 2500 U vial, or 357 μ L of sterile H₂O to the 25 KU vial to a final concentration of 70 U/ μ L. Incubate the solution for at least 10 minutes to ensure sufficient re-dissolved.

Storage and Stability

This product is stable after storage at -20°C for 12 months in lyophilized state from date of receipt. Upon reconstitution, protein aliquots should be stored at 4°C.

Procedure

1. Prepare the antibody sample in a neutral reaction buffer (eg. 50 mM sodium phosphate, 150 mM NaCl, pH 6.6 or PBS, pH 7.4).
2. Add IdeS Protease to the sample at a ratio of 1 unit of enzyme per 1 μ g of IgG antibody.
3. Gently mix the reaction and incubate at 37°C for 30-60 minutes.
4. Determine F(ab')₂ and Fc fragments using SDS-PAGE analysis.



The standard assay was performed by incubating 2 units of IdeS Protease and 2 μ g of recombinant human IgG under the above condition and analyzed by SDS-PAGE

Disclaimer

For Research Use or Further Manufacturing Only.