

Product Information | Certification of Analysis

| Product Information

CAS: N/A Lot No.

rTrypsin-N, Mass Spec Grade

Part No.	Name	Size/pkg
HLS TRY001N	rTrypsin-N, Mass Spec Grade	20 µg

Description: rTrypsin-N, Mass Spec Grade, is an innovative protease that specifically hydrolyzes peptide bonds at the amino side of lysine and arginine residues. rTrypsin-N can be applied for post-translational modification (PTM) proteins research and is beat for protein sequencing. Peptides produced more b ions than y ions in the mass spectrometer.

Physical Appearance: Supplied by lyophilized powder with 0.47 mg HEPES; 0.05mg CaCl₂ per vial.

Molecular Weight: 29 kDa

Resuspension Buffer: 200 μ L distilled water, pH 7.5

Storage Conditions: Store the lyophilized powder at -20° C, and reconstituted enzyme at -80° C for up to 30 days. **Shelf life:** 24 months at -80 °C

Stability: rTrypsin-N is maximally active at pH 7.5.

In-Solution Protein Digestion Protocol:

- 1. For maximum activity, resuspend Mass Spec Grade rTrypsin-N in 200 µL distilled water, pH 7.5.
- 2. Use 20 50 mM HEPES solution, 0.1% BT Surfactant, or Tris-HCI (pH 7.5) for the protein mixture cleavage (recommended).
- 3. Add 0.5µg/µL rTrypsin-N to a final enzyme : substrate ratio of 1 : 30. Mix well and incubated at 37 °C for 4 h.

I Quality Control

Purity: > 99.5% rTrypsin-N peak area analyzed by HPLC at 280 nm.

Specificity: < 5% nonspecific cleavage with Human Serum Albumin (HSA) sample. Digested products were incubated at 37 °C for 4 h, and was nonspecific cleavage was analyzed by LC-MS/MS.

Activity: 152 units/mg

Unit Definition: One unit is the amount of Mass spectrum Grade rTrypsin-N will hydrolyze per minutes at pH 10 at 25 °C, OD366, Light path = 1 cm.

MALDI-TOF Analysis: rTrysin-N is analyzed by MALDI-TOF, impurity peak is not found.

LC-MS/MS Analysis: HSA samples were dissolved and denatured for at 37 °C for 1 h. The denatured HSA was diluted at pH 7.5 and incubated with rTrypsin-N for 4 h. The digest was analyzed by LC-MS/MS, and the peptides results matched the peptides generated in a theoretical digestion results of HSA by rTrypsin-N.





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