



Sialidase Sp

Alpha-(2-3) Nueraminidase, NANase

Source

recombinant from *Streptococcus pneumoniae*

Catalog Number

E-S007

Certification of Analysis Lot Number

401.1G

EC

3.2.1.18

Applications

- Structural analysis of oligosaccharides
- Determining sialic acid linkage
- Glycoprotein deglycosylation
- Removing heterogeneity from glycoproteins

Contents

- 1 vial: Sialidase Sp - 60 μ l (300 mU)
in 250 mM Sodium phosphate pH 7.5
- 1 vial: Reaction buffer – 400 μ l
250mM Sodium phosphate, pH 6.0

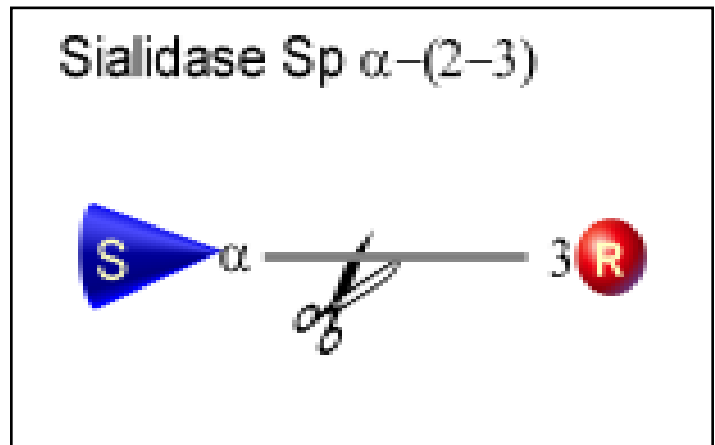
Specific Activity >75 U/mg

Activity >2.5 U/ml

Molecular Weight ~75,000 daltons

pH optimum 6.0, active over the range 4.5-7.

50 mM sodium phosphate (pH 6.0) provides the optimal buffer for enzyme activity with sialyllactose, a standard substrate. If glycosidase treatment is performed at suboptimal pH because of glycoprotein solubility or activity requirements, expect some diminution in enzyme activity.



Specific Activity

One unit of QA-Bio Sialidase Sp is defined as the amount of enzyme required to produce 1 μ mole of methylumbelliferone in 1 minute at 37°C, pH 5.0 from MU-NANA [2' .-(4-methyl-umbelliferyl)-*alpha*-D-N acetylneuraminic acid].

Specificity

All non-reducing terminal branched and unbranched α -(2-3) sialic acid.

Formulation

The enzyme is provided as a sterile-filtered solution in 250 mM Sodium phosphate pH 7.5.

Stability

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

Storage

Store enzyme at 4°C. Do not freeze.

Purity

QA-Bio Sialidase Sp 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.

E-S007 Sialadase Sp

Specifications - Protocol

Directions for use

1. Add up to 100 µg of glycoprotein or 1 nmol of oligosaccharide to tube.
2. Add de-ionized water to a total of 14 µl.
3. Add 4 µl 5x Reaction Buffer 6.0.
4. Add 2 µl Sialidase Sp.
5. Incubate at 37°C for 1 hour.

NOTE: longer incubation times are necessary if branched sialic acids are present.

Desialylation may be monitored by SDS-PAGE if the size differential between native and de-sialylated protein is sufficient for detection.

References:

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

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