

Sialidase 33B, *Clostridium perfringens*

CpNan33B (GH33)

| Catalogue number | Presentation |
|------------------|--------------|
| CZ02261 | 0.25 mg |
| CZ02262 | 3 x 0.25 mg |

Description

Sialidase 33B (CpNan33B), assigned the E.C. number 3.2.1.18, is a derivative of *Clostridium perfringens*. It is an exo- α -sialidase. The recombinant CpNan33B, purified from *Escherichia coli*, is a single-domain Glycoside Hydrolase family 33 (GH33) enzyme (see more details at www.cazy.org). The protein is supplied in a solution containing 35 mM NaHepes buffer (pH 7.5), 750 mM NaCl, 200 mM Imidazole, 3.5 mM CaCl₂, and 25% (v/v) glycerol, at a concentration of 0.25 mg/mL. Bulk quantities of this product can be made available upon request. To place an order, simply visit our website. We offer fast and secure shipping worldwide.

Electrophoretic Purity

The molecular integrity and purity of CpNan33B (GH33) were evaluated using sodium dodecyl sulfate-polyacrylamide gel electrophoresis (SDS-PAGE), followed by BlueSafe staining (MB15201) (Figure 1).

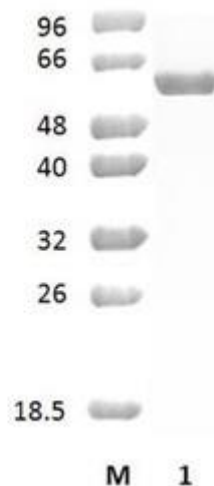


Figure 1. SDS-PAGE analysis of CpNan33B (GH33) was conducted in (Lane 1), employing a 14% polyacrylamide gel. The enzyme exhibits a band corresponding to a molecular weight of approximately 54,39 kDa. Lane M contains a Protein Marker for reference.

Storage temperature

The protein should be stored at -30°C to -15°C in a constant temperature freezer. The protein will remain stable till the expiry date if stored as specified.

Substrate specificity

CpNan33B (GH33) hydrolyses MUNeu5Ac and hydrolysis of glycosidic linkages of terminal sialic acid residues in oligosaccharides, glycoproteins, glycolipids, colominic acid and synthetic substrates.

Temperature and pH optima

The pH optimum for enzymatic activity is 6 while temperature optimum is 37 °C.

Enzyme activity

The substrate specificity and kinetic properties of CpNan33B (GH33) are detailed in the referenced publication provided below. To perform enzyme assays and determine specific activity values, adhere to the methodology outlined in the cited paper(s).

Reference

Boraston *et al.* (2007) *Biochemistry*. 46(40):11352-60.

Juge *et al.* (2016) *Biochem Soc Trans*. 44(1):166-75.

Customer Support

Our dedicated customer support team is always ready to assist you with any questions or technical issues you may have. Reach us via email at info@nzytech.com.

Quality control assay

Protein purity is determined to be $\geq 90\%$, as assessed by SDS-PAGE and subsequent BlueSafe staining (MB15201).

For life science research only. Not for use in diagnostic procedures.

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