Bacteroides Heparinase III



1-800-632-7799 info@neb.com www.neb.com

BioLabs

14 units	700 U/ml	Lot: 0031406
RECOMBINANT	Store at -80°C	Exp: 6/15

Description: Bacteroides Heparinase III cloned from Bacteroides eggerthii, also called Heparin Lyase III, is active on both heparin and heparan sulfate. The reaction yields oligosaccharide products containing unsaturated uronic acids which can be detected by UV spectroscopy at 232 nm.

Bacteroides Heparinase III BioLabs 1-800-632-7799 info@neb.com www.neb.com **P0737S** 700 U/ml Lot: 0031406 14 units

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Detailed Specificity:



All structural determinants for enzyme specificity are displayed in red.

Bacteroides Heparinase III specificity.

In stark contrast to the *Flavobacterium heparinum* Heparinase III that cleaves the glycosidic bond only between hexosamines and glucuronic acid residues, the Bacteroides Heparinase III can cleave the glycosidic bond between hexosamines and either iduronic acid or glucuronic acid residues. Flavobacterium heparinum Heparinase III is not active on an *N*-sulfoglucosamine with 6-sulfation, whereas the Bacteroides Heparinase III is active in the presence of 6-sulfation.

Source: Cloned from Bacteroides Eggerthii and expressed in pET21A.

Supplied in: 100 mM NaCl. 20 mM Tris-HCl (pH 7.5 @ 25°C), 1 mM Na EDTA and 5 mM CaCl.

Reagents Supplied with Enzyme:

10X Bacteroides Heparinase Reaction Buffer

Reaction Conditions:

- 1. Combine 10 µl of 1 mg/ml heparan sulfate. 10 µl Bacteroides Heparinase Reaction Buffer and H_oO in a total reaction volume of 100 µl.
- 2. Add 1 µl Bacteroides Heparinase III
- 3. Incubate reaction at 30°C for 1-24 hours (monitor absorbance at 232 nm for determination of partial or complete digestion).

Optimal incubation times and enzyme concentrations must be determined empirically for a particular substrate.

1X Bacteroides Heparinase Reaction Buffer:

100 mM NaCl (pH 7.0 @ 25°C) 20 mM Tris-HCI 1.5 mM CaCl

Unit Definition: One unit is defined as the amount of enzyme that will liberate 1.0 µmol unsaturated oligosaccharides from heparan sulfate per minute at 30°C and pH 7.0 in a total reaction volume of 100 ul.

Unit Definition Assay: Two fold dilutions of Bacteroides Heparinase III are incubated with 1 mg/ml porcine mucosal heparan sulfate in 1X *Bacteroides* Heparinase Reaction Buffer, in a 100 µl reaction. The reaction mix is incubated at 30°C. Liberation of unsaturated oligosaccharides is detected by real-time UV spectroscopy at 232 nm.

Molecular Weight: 75,000 daltons.

Quality Assurance: No contaminating exoglycosidase, sulfatase, uronidase or proteolytic activity could be detected (ND).

Quality Controls

Physical Purity: Purified to > 95% homogeneity as determined by SDS-PAGE analysis using Coomassie Blue detection and mass spectrometry analysis.

(see other side)

CERTIFICATE OF ANALYSIS

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(see other side)



Denotes either glucuronic acid or iduronic acid. All structural determinants for enzyme specificity are displayed in red.

Bacteroides Heparinase III specificity.

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Glycosidase and Sulfatase Assays: 1 unit of <i>Bacteroides</i> Heparinase III was incubated with 0.1 mM of AMC (7-amino-4-methyl-coumarin)		N,6- <i>O</i> -Sulfatase and Uronidase: \[\Delta\UA-(1-4)-GlcNS6S-AMC	ND
fluorescently-labeled oligosaccharides, in a 10 μ l reaction for 20 hours at 30°C. The reaction products were analyzed by TLC for digestion of substrate.		2-0-Sulfatase: <u>AUA2S-(1-4)-GICNS6S-AMC</u>	ND
No other glycosidase activities were detective with the following substrates:	ted (ND)	Protease Assay: After incubation of 5 uni Bacteroides Heparinase III with 0.2 nmol standard mixture of proteins for 20 hours	ts of of a at 37°C,
β -<i>N</i>-Acetylglucosaminidase: GlcNAcβ1-4GlcNAcβ1-4GlcNAc-AMC	ND	no proteolytic activity could be detected b PAGE.	y SDS-
β - <i>N</i> -Acetylgalactosaminidase:		Note: Avoid repeated freeze-thaw cycles.	
GalNAcβ1-4Galβ1-4Glc-AMC	ND	Heat Inactivation: 100°C for 1 minute.	
β -Galactosidase:	ND		
Galβ1-4GlcNAcβ1-3Galβ1-4Glc-AMC	ND		
β -Glucosidase: Glcβ1-4Glcβ1-4Glc-AMC	ND		

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Glycosidase and Sulfatase Assays: 1 unit of <i>Bacteroides</i> Heparinase III was incubated with 0.1 mM of AMC (7-amino-4-methyl-coumarin) fluorescently-labeled oligosaccharides, in a 10 µl reaction for 20 hours at 30°C. The reaction products were analyzed by TLC for digestion of substrate.				
No other glycosidase activities were detected (ND) with the following substrates:				
β -N-Acetylglucosaminidase: GlcNAcβ1-4GlcNAcβ1-4GlcNAc-AMC	ND			
β- N-Acetylgalactosaminidase: GalNAcβ1-4Galβ1-4Glc-AMC	ND			

β -Galactosidase:

β -Glucosidase: Glcβ1-4Glcβ1-4Glc-AMC	ND
Galβ1-4GlcNAcβ1-3Galβ1-4Glc-AMC	ND
Galβ1-3GlcNAcβ1-4Galβ1–4Glc-AMC	ND

Δ UA-(1-4)-GIcNS6S-AMC	ND		
2-<i>O</i>-Sulfatase: ΔUA2S-(1-4)-GIcNS6S-AMC	ND		
Protease Assay: After incubation of 5 units of <i>Bacteroides</i> Heparinase III with 0.2 nmol of a standard mixture of proteins for 20 hours at 37°C, no proteolytic activity could be detected by SDS-PAGE.			
Note: Avoid repeated freeze-thaw cycles.			
Heat Inactivation: 100°C for 1 minute.			

N,6-O-Sulfatase and Uronidase: