



P0730S

RX RX

100

BioLabs

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

www.neb.com

400 units 8.000 U/ml Lot: 0021210 RECOMBINANT Store at -20°C Exp: 10/14

Description: β 1-4 Galactosidase is a highly specific exoglycosidase that catalyzes the hydrolysis of β 1-4 linked D-galactopyranosyl residues from oligosaccharides.

Specificity:

Detailed Specificity: Specificity can vary depending on incubation time and branching structure.

β1-4	Enew encland
Galactosidase	BioLabsin.
P0730S 002121014101	1-800-632-7799 info@neb.com www.neb.com

P0730S



400 units 8.000 U/ml

RECOMBINANT Store at -20°C Exp: 10/14

Description: β1-4 Galactosidase is a highly specific exoglycosidase that catalyzes the hydrolysis of β 1-4 linked D-galactopyranosyl residues from oligosaccharides.

Specificity:



Detailed Specificity: Specificity can vary depending on incubation time and branching structure.

A) 0.1 nm/µl substrate, 1 hour incubation

 $Gal\beta(1-4)GlcNAc\beta(1-2)Man\alpha(1-6)$ ManB(1-4)GlcNAc $Gal\beta(1-4)GlcNAc\beta(1-2)Man\alpha(1-3)$

B) 0.1 nm/ul substrate. 1 hour incubation

 $Gal\beta(1-4)GlcNAc\beta(1-2)Man\alpha(1-6)$ Gal β (1-4)GlcNAc β (1-4)Man α (1-3)Man β (1-4)GlcNAc GalB(1-4)GlcNAcB(1-2)

C) 0.1 nm/µl substrate, 1 hour incubation



Figure 1: Detailed specificity of *β*1-4 Galactosidase. Reactions (A), (B) and (C) contained 2 units, 4 units and 8 units of β 1-4 Galactosidase, respectively, and either 1X G4 or 1X G6 Reaction Buffer in a total reaction volume of 10 µl. Reactions were incubated at 37°C.

A) 0.1 nm/ul substrate. 1 hour incubation



B) 0.1 nm/µl substrate, 1 hour incubation

 $Gal\beta(1-4)GlcNAc\beta(1-2)Man\alpha(1-6)$ GalB(1-4)GlcNAcB(1-4)Mana(1-3)ManB(1-4)GlcNAc GalB(1-4)GlcNAcB(1-2)

C) 0.1 nm/µl substrate, 1 hour incubation



Figure 1: Detailed specificity of B1-4 Galactosidase. Reactions (A). (B) and (C) contained 2 units. 4 units and 8 units of B1-4 Galactosidase, respectively, and either 1X G4 or 1X G6 Reaction Buffer in a total reaction volume of 10 µl. Reactions were incubated at 37°C.

Source: Cloned from Bacteroides fragilis and expressed in *E. coli* (1).

Supplied in: 50 mM NaCl, 20 mM Tris-HCl (pH 7.5 @ 25°C) and 1 mM Na₂EDTA.

Reagents Supplied with Enzyme: 10X G4 Reaction Buffer

Reaction Conditions:

1X G4 Reaction Buffer 50 mM Sodium Citrate (pH 6.0 @ 25°C) and 100 mM NaCl. Incubate at 37°C.

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

Unit Definition: One unit is defined as the amount of enzyme required to cleave > 95% of the terminal, β -D-galactose from 1 nmol Gal β 1-4GlcNAcB1-3GalB1-4Glc-7-amino-4-methylcoumarin (AMC), in 1 hour at 37°C in a total reaction volume of 10 µl.

Unit Definition Assay: Two fold serial dilutions of B1-4 Galactosidase are incubated with 1 nmol AMC-labeled substrate in 1X G4 Reaction Buffer, in

Source: Cloned from Bacteroides fragilis and expressed in E. coli (1).

Supplied in: 50 mM NaCl, 20 mM Tris-HCl (pH 7.5 @ 25°C) and 1 mM Na EDTA.

Reagents Supplied with Enzyme: 10X G4 Reaction Buffer

Reaction Conditions:

1X G4 Reaction Buffer 50 mM Sodium Citrate (pH 6.0 @ 25°C) and 100 mM NaCl. Incubate at 37°C.

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

Unit Definition: One unit is defined as the amount of enzyme required to cleave > 95% of the terminal, β -D-galactose from 1 nmol Gal β 1-4GlcNAcB1-3GalB1-4Glc-7-amino-4-methylcoumarin (AMC), in 1 hour at 37°C in a total reaction volume of 10 µl.

Unit Definition Assay: Two fold serial dilutions of B1-4 Galactosidase are incubated with 1 nmol AMC-labeled substrate in 1X G4 Reaction Buffer, in a 10 ul reaction. The reaction mix is incubated for 1 hour at 37°C. Separation of reaction products are visualized via thin layer chromatography (2).

Specific Activity: 50,000 units/mg

Molecular Weight: 94,000 daltons.

Quality Assurance: No contaminating exoglycosidase or proteolytic activity could be detected.

Quality Controls

Glycosidase Assays:

32 units of B1-4 Galactosidase were incubated with 0.1 mM of flourescently-labeled oligosaccharides and glycopeptides, in a 10 µl reaction for 20 hours at 37°C. The reaction products were analyzed by TLC for digestion of substrate.

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

(See other side)

CERTIFICATE OF ANALYSIS

a 10 µl reaction. The reaction mix is incubated for 1 hour at 37°C. Separation of reaction products are visualized via thin layer chromatography (2).

Specific Activity: 50,000 units/mg

Molecular Weight: 94,000 daltons.

Quality Assurance: No contaminating exoglycosidase or proteolytic activity could be detected.

Quality Controls

Glycosidase Assays:

32 units of B1-4 Galactosidase were incubated with 0.1 mM of flourescently-labeled oligosaccharides and glycopeptides, in a 10 µl reaction for 20 hours at 37°C. The reaction products were analyzed by TLC for digestion of substrate.

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

(See other side)

No other glycosidase activities were detect with the following substrates:	ed (ND)	Manα1-3Manβ1-4GlcNAc-AMC Manα1-6Manα1-6(Manα1-3)Man-AMC
β- N-Acetylglucosaminidase: GlcNAcβ1-4GlcNAcβ1-4GlcNAc-AMC	ND	β- Glucosidase: Glcβ1-4Glcβ1-4Glc-AMC
α -N-Acetylgalactosaminidase: GalNAc α 1-3(Fuc α 1-2)Gal β 1-4Glc-AMC	ND	α- Glucosidase: Glcα1-6Glcα1-4Glc-AMC
α -Fucosidase: Fuc α 1-2Gal β 1-4Glc-AMC	ND	β -Xylosidase: Xylβ1-4Xylβ1-4Xylβ1-4Xyl-AMC
Galβ1-4 (Fucα1-3)GicNAcβ1-3Galβ1- 4Gic-AMC	ND	β -Mannosidase: Manβ1-4Manβ1-4Man-AMC
α-Galactosidase:	ND	
	ND	Endo F ₁ , F ₂ , H: Dependented invertees high mennees
$Gal\alpha 1-6Gal\alpha 1-6Glc\alpha 1-2Fru-AMC$	ND	Dansylated invertase night maintose.
α -Neuraminidase: Neu5Ac α 2-3Gal β 1-3GlcNAc β 1-3Gal β 1-		Endo F ₂ , F ₃ : Dansylated fibrinogen biantennary.
4uic-Amic α- Mannosidase:	ND	PNGase F: Fluoresceinated fetuin triantennary.

Protease Assay: After incubation of 112 units of β 1-4 Galactosidase with 0.2 nmol of a standard mixture of proteins in a 20 μ l reaction, for 20 hours at 37°C, no proteolytic activity could be detected by SDS-PAGE.

Note: Recommended storage temperature is –20°C.

Heat Inactivation: 65°C for 10 minutes.

References:

ND

1. McLeod, E., New England Biolabs, Inc. unpublished results.

2. Wong-Madden, S.T. and Landry, D. (1995) *Glycobiology* 5, 19–28.

U.S. Patent No. 6,358,724

Page 2 (P0730)

ed (ND)	Manα1-3Manβ1-4GlcNAc-AMC Manα1-6Manα1-6(Manα1-3)Man-AMC
ND	β- Glucosidase: Glcβ1-4Glcβ1-4Glc-AMC
ND	α -Glucosidase: Glcα1-6Glcα1-4Glc-AMC
ND	β -Xylosidase: Xylβ1-4Xylβ1-4Xylβ1-4Xyl-AMC
ND	β -Mannosidase: Manβ1-4Manβ1-4Man-AMC
ND	
ND	Endo F_1, F_2, H :
ND	Dansylated invertase high mannose.
ND	Endo F₂, F₃: Dansylated fibrinogen biantennary.
α-Mannosidase:	
	ed (ND) ND ND ND ND ND ND

Protease Assay: After incubation of 112 units of β 1-4 Galactosidase with 0.2 nmol of a standard mixture of proteins in a 20 μ I reaction, for 20 hours at 37°C, no proteolytic activity could be detected by SDS-PAGE.

Note: Recommended storage temperature is -20°C.

Heat Inactivation: 65°C for 10 minutes.

References:

- 1. McLeod, E., New England Biolabs, Inc. unpublished results.
- 2. Wong-Madden, S.T. and Landry, D. (1995) *Glycobiology* 5, 19–28.

U.S. Patent No. 6,358,724