

# BIOFILES

FOR LIFE SCIENCE RESEARCH

2007  
Volume 2  
Number 3



ENZYMES, KITS AND  
REAGENTS FOR ANALYSIS  
OF:

AGAROSE

ALGINIC ACID

CELLULOSE, LICHENEN  
AND GLUCANS

HEMICELLULOSE AND  
XYLAN

CHITIN AND CHITOSAN

CHONDROITINS

DEXTRAN

HEPARANS

HYALURONIC ACID

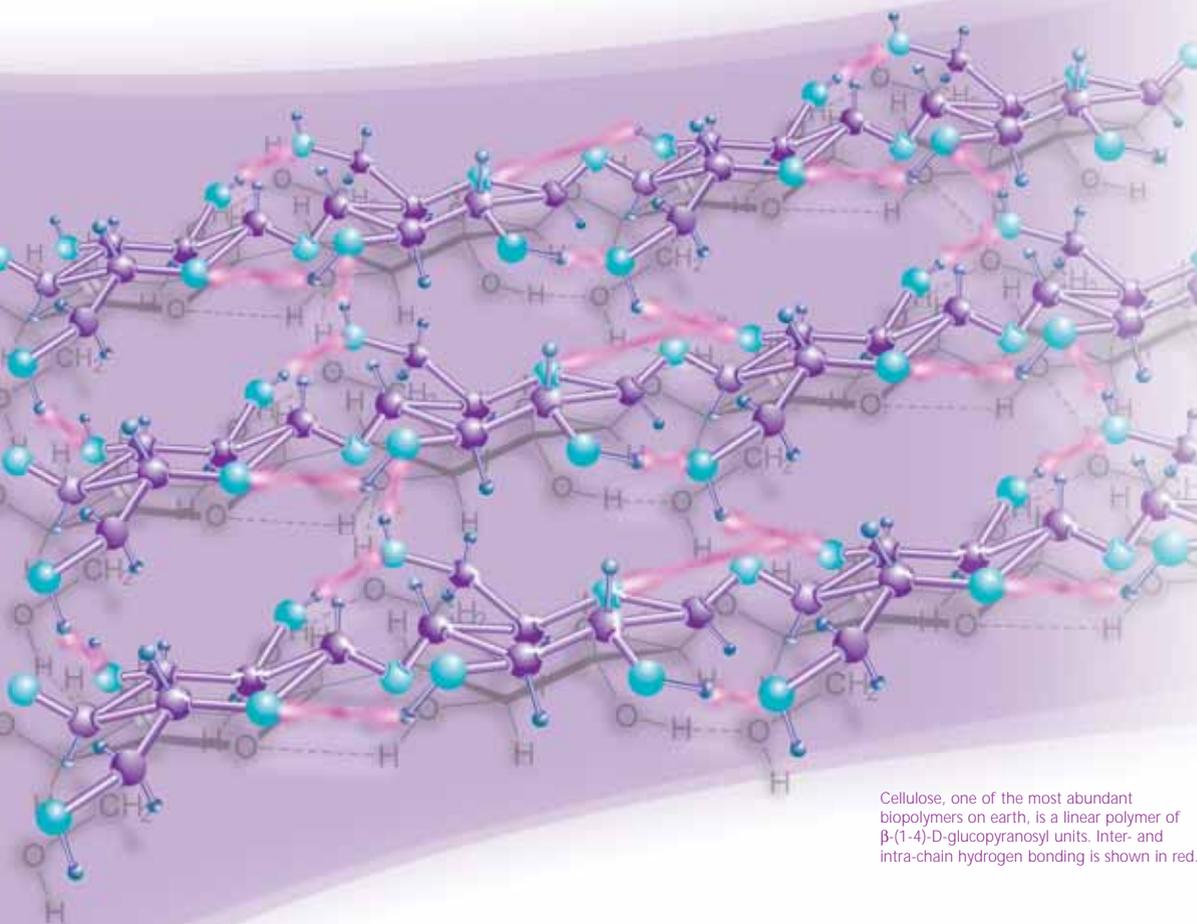
INULIN

PEPTIDOGLYCAN

PECTIN

PULLULAN

STARCH AND GLYCOGEN



Cellulose, one of the most abundant biopolymers on earth, is a linear polymer of  $\beta$ -(1-4)-D-glucopyranosyl units. Inter- and intra-chain hydrogen bonding is shown in red.

## Complex Carbohydrate Analysis: Enzymes, Kits and Reagents

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- **New!** Search for Plants Associated with Physiological Activity
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## Introduction

Complex carbohydrates compose the most abundant class of biopolymers on earth. Because of their structural and functional diversity, they have found applications in biomedical, nutritional, textile, cosmetic and countless other industries.

The complex carbohydrates of the extracellular matrices such as hyaluronic acid and chondroitin sulfate are finding utility in anti-inflammatory and cell proliferation applications. *In vivo*, hyaluronic acid forms a coating around chondrocytes in articular cartilage and together with the proteoglycan, aggrecan, is responsible for the uptake and retention of water. The two major glycan components of aggrecan are chondroitin sulfate and keratan sulfate. Hyaluronic acid may also interact with cell surface receptors, such as CD44, involved in lymphocyte activation. The degradation products of hyaluronic acid may also interact with Toll-like receptors in macrophages. Heparan sulfate is commonly found as a component of cell surface proteoglycans. It is also found in the extracellular matrix. Heparan sulfate appears to have a broad range of biological functions including regulation of thrombosis, growth factor signaling, cell proliferation, adhesion and mobility. Depending on its morphology, location and ligands, heparan sulfate may inhibit or promote metastasis. Heparan sulfate is known to bind several protein ligands. Most notably, its binding affinity with antithrombin has been extensively utilized in the form of the anticoagulant, heparin.

Dextrans also help to decrease vascular thrombosis. By binding to the endothelium, platelets and red blood cells, dextrans impart an electronegative environment in the blood vessel resulting in a reduction of red blood cell aggregation and platelet adhesion to the vascular endothelium. *In vivo*, dextran solutions have also been used for blood volume expansion. Conversely, chitosan has the ability to induce clot formation. It is used in wound healing, particularly as a coating for bandages. Chitin may also aid in wound healing by accelerating collagen production. Chitosan is also used to enhance plant growth and may help plants resist fungal infection.

Chitin and starch are used as binders in the paper, dye, textiles and adhesives industries. Chitin and chitosan are also used as filtration aids, particularly in the waste water treatment industry. Chitosan aids in particulate aggregation as well as removal of phosphorus, metals, and grease from waste water. Modified agarose, chitin, starch and dextrans have been manipulated to produce media with controlled pore size for chemical separations. Beaded forms of cross-linked agarose and dextrans are the components of size exclusion, ion exchange and affinity chromatography media. Agarose and soluble starch are commonly used as electrophoresis media.

In the food industry, starches, agglutinates, agarose, chitins, chitosans and pectins are used as gelling, thickening and encapsulating agents. Pectins and inulins are common components of dietary fiber supplements and may help to increase nutrient uptake. Pullulan is a common component of edible films.

### The Enzyme Explorer

Your Comprehensive Source for Products and Technical Resources for Glycobiology [sigma-aldrich.com/enzymeexplorer](http://sigma-aldrich.com/enzymeexplorer)

#### Carbohydrate Analysis

- Complex Carbohydrate and Polysaccharide Analysis
- Proteoglycan and Glycoprotein Analysis

#### Carbohydrate Metabolism

- Carbohydrate Metabolite Library
- Enzymes Involved in Carbohydrate Metabolism
- Metabolic Pathway Charts and Animations

#### Enzymatic-Based Kits for the Quantitation of Carbohydrates

- Total Dietary Fiber
- Starch
- Glucose
- Fructose
- Sucrose

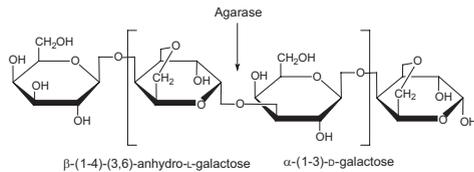
For additional technical information including literature citations pertaining to the content in this publication, visit the Enzyme Explorer's "Enzymatic Carbohydrate Analysis Resource"

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The Enzyme Research Resource

## Agarase

### Agarase Specificity



#### Agarose

Agarose is the principal neutral gelling component of agar extracted from algae. Agarose is a complex range of polysaccharide chains composed of alternating  $\alpha$ -(1-3)-D-galactosyl- $\beta$ -(1-4)-anhydro-L-galactosyl units.

### Agarase

Agarase catalyzes the hydrolysis of 1,3- $\beta$ -D-galactosidic linkages in agarose, giving the tetramer as the predominant product.

#### Agarase from *Pseudomonas atlantica*

Agarose 3-glycanohydrolase  
[37288-57-6] E.C. 3.2.1.81

#### lyophilized powder, activity: 1,000-3,000 units/mg solid

Contains phosphate buffer salts. May contain bovine serum albumin to standardize protein content

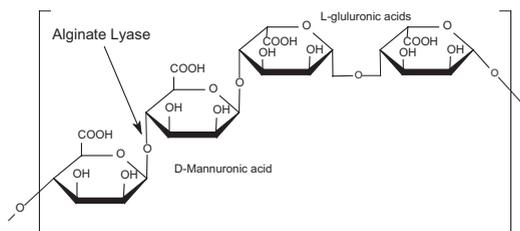
One unit will produce 1.0  $\mu$ g of reducing sugar (measured as D-galactose) from agar per min at pH 6.0 at 40 °C.

2-8°C

A6306-1KU	1,000 units
A6306-5KU	5,000 units

## Alginate Lyase

### Alginate Lyase Specificity



#### Alginic Acid

Composed of blocks of  $\beta$ -D-(1-4) mannuronic acid homopolymeric regions (MMMM...),  $\alpha$ -L-(1-4)-guluronic acid (GGGG...) homopolymeric regions, and alternating copolymer regions of  $\beta$ -D-(1-4) mannuronic acid -  $\alpha$ -L-(1-4)-guluronic acid (GMGMGM...). Bacterial alginic acid can be acetylated at the 2 or 3 positions on mannuronic acid.

Alginate lyase cleaves at the  $\beta$ -(1-4)-D-mannuronic bonds residues to yield oligosaccharides with 4-deoxy- $\alpha$ -L-erythro-hex-4-enopyranuronosyl groups at their non-reducing terminus.

#### Alginate Lyase from *Flavobacterium* sp.

E.C. 4.2.2.3

#### powder

#### activity: $\geq 10,000$ units/g solid

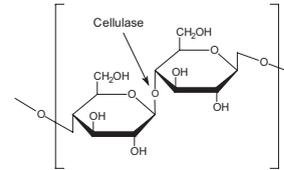
Add 0.15 mL of enzyme solution (1 un/mL) to 4.5 mL of 0.1% sodium alginate (pH 6.3). Incubate at 37 °C for 30 minutes. Terminate reaction by addition of 4.65 mL of 0.1 N NaOH.

One unit will produce an increase the A235 nm of 1.0 per minute per mL of sodium alginate solution at pH 6.3 at 37 °C

2-8°C

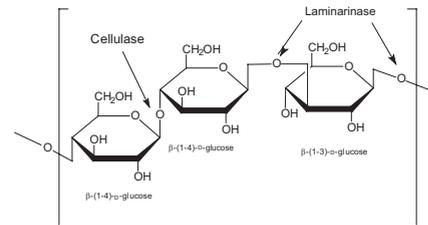
A1603-100MG	100 mg
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## Cellulose, Lichenan and Glucan Degrading Enzymes



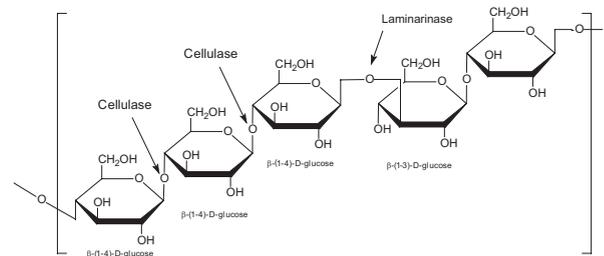
#### Cellulose

Polymer of  $\beta$ -(1-4)-D-glucopyranosyl units



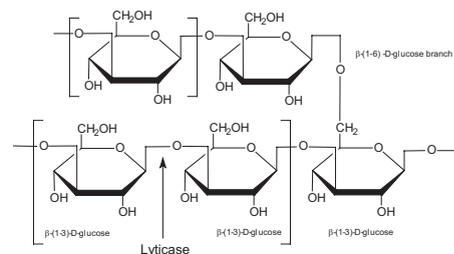
#### Lichenan

Repeating linear polymer of two  $\beta$ -(1-4)-D-glucopyranosyl and one  $\beta$ -(1-3)-D-glucopyranosyl unit.



#### Cerial $\beta$ -Glucan

Polymer of  $\beta$ -(1-4)-D-glucopyranosyl units occurring as predominantly as cellotriose and cellotetraose separated by single  $\beta$ -(1-3)-D-glucopyranosyl units. Cross-linking can occur within the consecutive cellotriose regions.



#### Yeast $\beta$ -Glucan

Polymer of  $\beta$ -(1-3)-D-glucopyranosyl units with branching at  $\beta$ -(1-6)-D-glucopyranosyl units.

## Cellulase

Cellulase catalyzes the endohydrolysis of 1,4- $\beta$ -D-glucosidic linkages in cellulose, lichenin and cereal  $\beta$ -D-glucans

### Cellulase from *Aspergillus* sp.

#### Carezyme® 1000L

[9012-54-8] E.C. 3.2.1.4

**activity:  $\geq 1000$  U/g**

Produced by submerged fermentation of a genetically modified *Aspergillus* microorganism

A product of Novozyme Corp.

2-8°C

C2605-50ML	50 mL
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C2605-250ML	250 mL
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### Cellulase from *Aspergillus niger*

1,4-(1,3:1,4)- $\beta$ -D-Glucan 4-glucano-hydrolase

[9012-54-8] E.C. 3.2.1.4 EC No. 2327344

**powder, activity:  $\geq 0.3$  units/mg solid**

One unit will liberate 1.0  $\mu$ mole of glucose from cellulose in 1 hr at pH 5.0 at 37 °C (2 hr incubation time).

2-8°C

C1184-5KU	5,000 units
-----------	-------------

C1184-25KU	25,000 units
------------	--------------

C1184-100KU	100,000 units
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### Cellulase from *Trichoderma reesei* ATCC 26921

1,4-(1,3:1,4)- $\beta$ -D-Glucan 4-glucano-hydrolase

[9012-54-8] E.C. 3.2.1.4

#### Celluclast® 1.5L

**▶ aqueous solution, activity:  $\geq 700$  U/g**

Produced by submerged fermentation of a selected strain of the fungus *Trichoderma reesei* and catalyzes the breakdown of cellulose into glucose, cellobiose, and higher glucose polymers. density.....1.2 g/mL, 25 °C

A product of Novozyme Corp.

2-8°C

C2730-50ML	50 mL
------------	-------

**▶ lyophilized powder, activity:  $\geq 1$  unit/mg solid**

One unit will liberate 1.0  $\mu$ mole of glucose from cellulose in 1 hr at pH 5.0 at 37 °C (2 hr incubation time).

2-8°C

C8546-2.5KU	2,500 units
-------------	-------------

C8546-5KU	5,000 units
-----------	-------------

C8546-10KU	10,000 units
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### Cellulase from *Trichoderma viride*

1,4-(1,3:1,4)- $\beta$ -D-Glucan 4-glucano-hydrolase

[9012-54-8] E.C. 3.2.1.4 EC No. 2327344

**▶ plant cell culture tested, activity: 3-10 units/mg solid**

**Composition: protein ~50% (biuret); contains lactose and glucose**

One unit will liberate 1.0  $\mu$ mole of glucose from cellulose in 1 hr at pH 5.0 at 37 °C (2 hr incubation time).

2-8°C

C1794-5KU	5,000 units
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C1794-10KU	10,000 units
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**▶ crude powder, activity: 3-10 units/mg solid**

**Composition: protein ~50% (biuret)**

One unit will liberate 1.0  $\mu$ mole of glucose from cellulose in 1 hr at pH 5.0 at 37 °C (2 hr incubation time).

2-8°C

C9422-5KU	5,000 units
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C9422-10KU	10,000 units
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**▶ Onozuka RS, powder, activity:  $\geq 5,000$  units/g solid**

Manufactured by Yakult

One unit will liberate 1.0  $\mu$ mole of glucose from cellulose in 1 hr at pH 5.0 at 37 °C (2 hr incubation time).

2-8°C

C0615-1G	1 g
----------	-----

## Driselase

### Driselase from *Basidiomycetes* sp.

[85186-71-6]

**powder, Protein: ~15%**

Crude powder containing laminarinase, xylanase and cellulase.

-20°C

D9515-1G	1 g
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D9515-5G	5 g
----------	-----

D9515-25G	25 g
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## Cellulose, Lichenan and Glucan Degradins Enzymes

### $\beta$ -Glucanase/Laminarinase

Laminarinase catalyzes the endohydrolysis of 1,3- or 1,4-linkages in  $\beta$ -D-glucans when the glucose residue whose reducing group is involved in the linkage to be hydrolyzed is itself substituted at C-3.

#### $\beta$ -Glucanase from *Aspergillus niger*

[9074-98-0] E.C. 3.2.1.6 EC No. 2329802

**BioChemika, powder, dark-brown, activity: ~1 unit/mg**

One Unit corresponds to the amount of enzyme which will release 1  $\mu$ mol of reducing sugar equivalents (expressed as glucose) per minute at pH 5.0 and 55 °C, using  $\beta$ -D-glucan (Catalog No. 49102) as substrate

**2-8°C**

**49101-100MG 100 mg**

**49101-500MG 500 mg**

#### $\beta$ -1,3-D-Glucanase from *Helix pomatia*

[9044-93-3] E.C. 3.2.1.39 EC No. 2329273

**BioChemika, powder, light beige, activity: 0.5-1.5 units/mg**

One Unit corresponds to the amount of enzyme which liberates 1  $\mu$ mol glucose from laminarin (Catalog No. 61340) per minute at pH 5.0 and 37 °C.

Improved filterability of wines by enzymic decomposition of carbohydrate-containing colloids<sup>1</sup>; Induction of hydrolases as a defense reaction against pathogens, review<sup>2</sup>

**Lit. cited:** 1. Wucherpfennig, K., and Dietrich, H., Weinwirtschaft **118**, 598 (1982)

2. Boller, T., UCLA Symp. Mol. Cell Biol., New Ser. **22**, 247 (1985)

**-20°C** WET ICE

**49103-10MG 10 mg**

**49103-50MG 50 mg**

#### Laminarinase from *Penicillium sp.*

endo-1,3(4)- $\beta$ -glucanase; 1,3-(1,3;1,4)- $\beta$ -D-Glucan 3(4)-glucanohydrolase [62213-14-3] E.C. 3.2.1.6

**lyophilized powder, activity: 5-10 units/mg protein**

Lyophilized powder containing acetate buffer salts

#### composition

Protein ~70% (biuret)

One unit will liberate 1.0 mg of reducing sugar (measured as glucose) from laminarin per min at pH 5.0 at 37 °C.

Contains cellulase and  $\alpha$ -amylase

**2-8°C**

**L9259-25UN 25 units**

#### Laminarinase from *Trichoderma sp.*

endo-1,3(4)- $\beta$ -glucanase; 1,3-[1,3;1,4]- $\beta$ -D-Glucan 3(4)-glucanohydrolase [62213-14-3] E.C. 3.2.1.6

**powder, activity: 100-400 units/g solid**

Contains chitinase activity.

One unit will liberate 1.0 mg of reducing sugar (measured as glucose) from laminarin per min at pH 5.0 at 37 °C.

Contains cellulase and  $\alpha$ -amylase

**2-8°C**

**L5272-5UN 5 units**

**L5272-25UN 25 units**

### Lyticase

Lyticase hydrolyzes poly- $\beta$ -(1-3)-glucose such as yeast cell wall  $\beta$ -glucan.

#### Lyticase from *Arthrobacter luteus*

[37340-57-1]

Yeast cells are difficult to disrupt because the cell walls may form capsules or resistant spores. DNA can be extracted from yeast by using lysing enzymes such as lyticase, chitinase, zymolase, and gluculase to induce partial spheroplast formation; spheroplasts are subsequently lysed to release DNA. Lyticase is preferred to digest cell walls of yeast and generate spheroplasts from fungi for transformation. Reported to be useful for lysis of *Ashbya*, *Candida*, *Debaryomyces*, *Eremothecium*, *Endomyces*, *Hansenula*, *Hanseniaspora*, *Kloeckera*, *Kluyveromyces*, *Lipomyces*, *Metschikowia*, *Pichia*, *Pullularia*, *Torulopsis*, *Saccharomyces*, *Saccharomycopsis*, *Saccharomyces*, and *Schwanniomyces* species.

One unit will produce a  $\Delta$ A800 of 0.001 per min at pH 7.5 at 25 °C, using a suspension of yeast as substrate in a 3 mL reaction mixture.

**▶ Lyophilized powder, activity:  $\geq$ 2,000 units/mg protein, Protein:  $\geq$ 20%**

Partially purified, lyophilized powder containing potassium phosphate buffer salts and stabilizers

**-20°C**

**L2524-10KU 10,000 units**

**L2524-25KU 25,000 units**

**L2524-50KU 50,000 units**

**L2524-200KU 200,000 units**

**▶ Lyophilized powder, activity:  $\geq$ 200 units/mg solid**

**-20°C**

**L4025-25KU 25,000 units**

**L4025-50KU 50,000 units**

**L4025-100KU 100,000 units**

**L4025-250KU 250,000 units**

**L4025-1MU 1,000,000 units**

**▶ partially purified powder, activity:  $\geq$ 2,000 units/mg protein**

Partially purified powder containing ammonium sulfate and stabilizer

#### composition

Protein ~20% (biuret)

**2-8°C**

**L5263-25KU 25,000 units**

**L5263-50KU 50,000 units**

**L5263-200KU 200,000 units**

### Lyticase from *Oerskovia xanthineolytica*

[37340-57-1]

**recombinant, expressed in *Escherichia coli*, lyophilized powder**

Purified recombinant  $\beta$ -(1,3)-glucanase preparation that is protease-free. Vial of  $\geq 500$  units.

One unit will produce a  $\Delta A_{800}$  of 0.001 per min at pH 7.5 at 25 °C, using a suspension of yeast as substrate in a 3 mL reaction mixture.

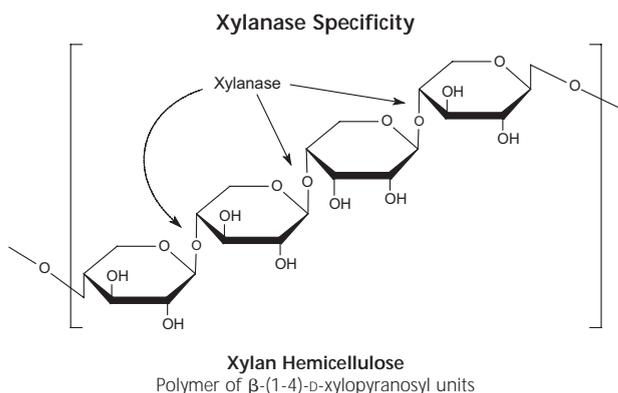
An exceptionally stable enzyme preparation with very low levels of nucleic acid and nuclease contamination.

-20°C

L4276-1VL

1 vial

## Hemicellulose and Xylan Degrading Enzymes



Xylan hemicelluloses are a group of plant-derived heteropolysaccharides associated with cellulose and lignin. The most common hemicelluloses are: xylan, glucuronoxylan, arabinoxylan, glucomannan and xyloglucan. In angiosperms, the principal hemicellulose component, xylan, is a polymer of  $\beta$ -(1-4)-D-xylopyranose. In arabinoxylan, branching occurs at the C2 & C3 positions with  $\alpha$ -L-arabinofuranose. Glucuronoxylan, also found in angiosperms, has the xylan backbone with 4-O methylglucuronic acid branching. In addition, arabinose branching as well as acetylation may be present. Gymnosperms contain glucomannans comprised primarily of D-mannosyl and D-glucosyl residues.

### Hemicellulase

#### Hemicellulase from *Aspergillus niger*

[9025-56-3]

**powder, activity: 0.3-3.0 units/mg solid (using a  $\beta$ -galactose dehydrogenase system and locust bean gum as substrate)**

An undefined mixture of glycolytic enzymes usually containing xylanase, mananase and other activities.

Contains lactose as standardization of activity

One unit will produce a relative fluidity change of 1 per 5 minutes using locust bean gum as substrate at pH 4.5 at 40 °C

-20°C

H2125-150KU

150,000 units

### Xylanase

Xylanase catalyzes the endohydrolysis of  $\beta$ -(1-4)-D-xylosidic linkages in xylans yielding various  $\beta$ -(1-4)-D-xylooligosaccharides.

#### Xylanase from *Thermomyces lanuginosus*

**Pentopan Mono BG®**

[37278-89-0]

**powder, activity:  $\geq 2500$  units/g, recombinant, expressed in *Aspergillus oryzae***

Purified endo (1,4)- $\beta$ -xylanase from *Thermomyces lanuginosus*. Produced by submerged fermentation of a genetically modified *Aspergillus oryzae* microorganism.

A product of Novozyme Corp.

2-8°C

X2753-10G

10 g

X2753-50G

50 g

#### Xylanase from *Trichoderma viride*

endo-1,4- $\beta$ -Xylanase; 1,4- $\beta$ -D-Xylanxylanohydrolase  
[9025-57-4] E. C. 3.2.1.8 EC No. 2534397

**lyophilized powder, activity: 100-300 units/mg protein**

Contains sorbitol and sodium acetate buffer salts

**composition**

Protein ~50% (biuret)

One unit will liberate 1  $\mu$ mole of reducing sugar measured as xylose equivalents from xylan (X0627) per min at pH 4.5 at 30 °C.

cellulase .....	<0.2%	$\beta$ -xylosidase .....	<0.002%
$\beta$ -glucosidase .....	<0.01%		

2-8°C

X3876-250UN

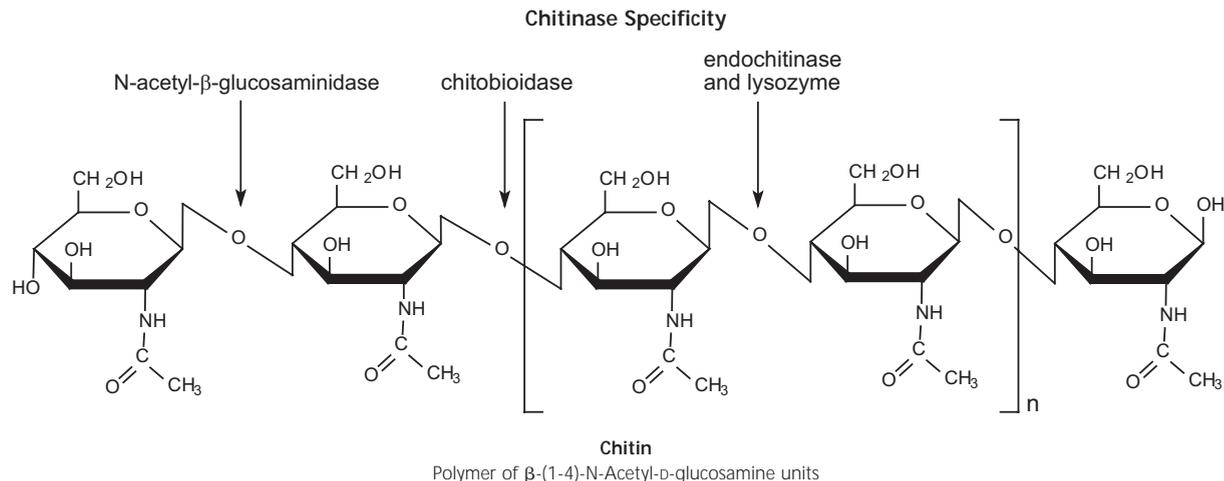
250 units

X3876-1KU

1,000 units

## Chitin and Chitosan Degrading Enzymes

### Chitinase



Chitinases have been detected in many organisms, including bacteria, fungi, plants, invertebrates and vertebrates. Chitinases are broadly classified as endo- and exochitinases. The endochitinase activity is defined as the random cleavage at internal points in the chitin chain. The exochitinase activity is defined as the progressive action starting at the non reducing end of chitin with the release of chitobiose or N-acetylglucosamine units. Chitobiosidase and N-acetyl- $\beta$ -glucosaminidase are considered exochitinases. The combination of endo- and exochitinases results in a synergistic increase in the chitinolytic activity.

#### Chitinase from *Serratia marcescens*

Chitodextrinase; Poly(1,4- $\beta$ -[2-acetamido-2-deoxy-D-glucoside])  
glycanohydrolase  
[9001-06-3] E.C. 3.2.1.14

##### lyophilized powder, activity: 400-1,200 units/g solid

Lyophilized powder containing phosphate buffer salts

##### composition

Protein 20-40% (biuret)

One unit will liberate 1.0 mg of N-acetyl-D-glucosamine from chitin per hour at pH 6.0 at 25 °C in a 2 hour assay.

$-20^{\circ}\text{C}$

C7809-1UN	1 unit
C7809-5UN	5 units
C7809-10UN	10 units

#### Chitinase from *Streptomyces griseus*

Chitodextrinase; Poly(1,4- $\beta$ -[2-acetamido-2-deoxy-D-glucoside])  
glycanohydrolase  
[9001-06-3] E.C. 3.2.1.14 EC No. 2325787

##### lyophilized powder (Essentially salt free), activity: 200-800 units/g solid

One unit will liberate 1.0 mg of N-acetyl-D-glucosamine from chitin per hour at pH 6.0 at 25 °C in a 2 hour assay.

$-20^{\circ}\text{C}$

C6137-5UN	5 units
C6137-25UN	25 units
C6137-50UN	50 units

#### Chitinase from *Trichoderma viride*

N-acetyl- $\beta$ -glucosaminidase and chitodextrinase  
E.C. 3.2.1.14 and 3.2.1.52

##### lyophilized powder, activity: $\geq 600$ units/g solid

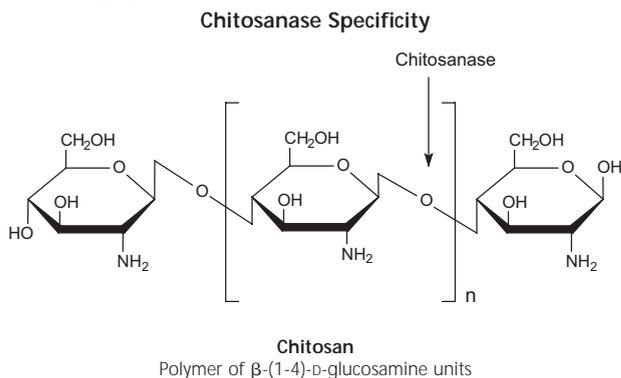
The chitinolytic enzymes from *T. viride* are a mixture of extracellular chitinolytic enzymes, which exhibit exo- and endochitinase activities including N-acetyl- $\beta$ -glucosaminidase and chitobiosidase.

One unit will liberate 1.0 mg of N-acetyl-D-glucosamine from chitin per hour at pH 6.0 at 25 °C in a 2 hour assay.

$-20^{\circ}\text{C}$  WET ICE

C8241-25UN	25 units
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## Chitosanase



Chitosanase catalyzes the endohydrolysis of  $\beta$ -(1-4)-linkages between D-glucosamine (GlcN-GlcN) residues in chitosan. The enzyme from *Streptomyces* has been reported to also hydrolyze the GlcNAc-GlcN linkage in partially acetylated chitosan.

### Chitosanase from *Streptomyces* sp.

Chitosan N-acetylglucosaminohydrolase  
[51570-20-8] E.C. 3.2.1.132

**buffered aqueous glycerol solution, activity:**  
 **$\geq 15$  units/mg protein**

Solution in 50% glycerol containing 100 mM sodium acetate,  
pH 5.0

One unit will liberate 1.0  $\mu$ mole of reducing sugar (measured as  
D-glucosamine equivalents) from chitosan per minute at pH 5.5 at  
37 °C.

**-20°C**

**C0794-10UN**

**10 units**

### Chitosanase from *Streptomyces griseus*

Chitosan N-acetylglucosaminohydrolase  
[51570-20-8] E.C. 3.2.1.132

**lyophilized powder, activity: >50 units/mg protein**  
**(Bradford)**

Lyophilized powder containing potassium phosphate buffer salts.  
Purified by chromatography

One unit will release 1  $\mu$ mole of glucosamine from chitosan per  
min at pH 5.0 at 37 °C

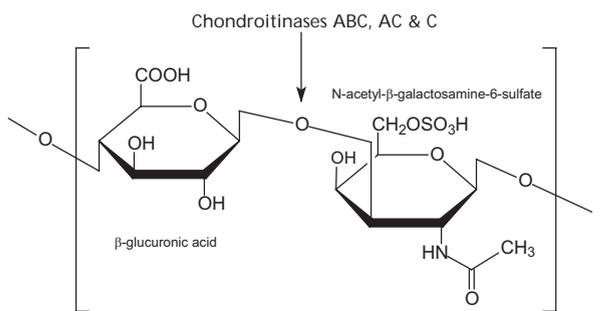
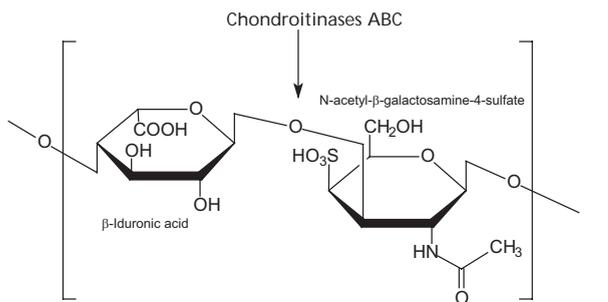
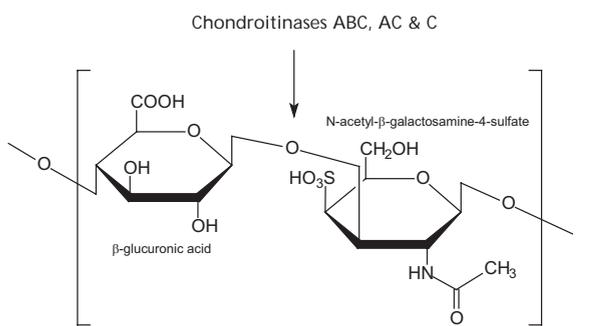
chitinase ..... <1.0 unit/mg protein

**-20°C**

**C9830-10UN**

**10 units**

## Chondroitinases



## Chondroitinases

### Chondroitinase ABC

Chondroitinase ABC catalyzes the eliminative degradation of polysaccharides containing  $\beta$ -(1-4)-D-hexosaminy and  $\beta$ -(1-3)-D-glucuronosyl or  $\alpha$ -(1-3)-L-iduronosyl linkages to disaccharides containing 4-deoxy- $\beta$ -D-gluc-4-enuronosyl groups. It acts on chondroitin 4-sulfate, chondroitin 6-sulfate, dermatan sulfate, and acts slowly on hyaluronate. Initial rates of degradation of chondroitin sulfate B, chondroitin, and hyaluronic acid were 40%, 20%, and 2%, respectively, that of chondroitin sulfate A and chondroitin sulfate C.

#### Chondroitinase ABC from *Proteus vulgaris*

Chondroitin ABC Lyase  
[9024-13-9] E.C. 4.2.2.4 EC No. 2327779

#### lyophilized powder, activity: 50-250 units/mg protein (using chondroitin sulfate C as substrate)

Contains potassium phosphate buffer salts and stabilizer; BSA-free.

#### composition, protein ~10% (Lowry)

Affinity purified

Packages based on chondroitinase C

One unit will liberate 1.0  $\mu$ mole of 2-acetamido-2-deoxy-3-O-( $\beta$ -D-gluc-4-ene-pyranosyluronic acid)-4-O-sulfo-D-galactose from chondroitin sulfate A or 1.0  $\mu$ mole of 2-acetamido-2-deoxy-3-O-( $\beta$ -D-gluc-4-ene-pyranosyluronic acid)-6-O-sulfo-D-galactose from chondroitin sulfate C per min at pH 8.0 at 37 °C.

protease ..... essentially free

-20°C

C3667-5UN 5 units

C3667-10UN 10 units

### Chondroitinase AC

Chondroitinase AC is an eliminase that degrades chondroitin sulfates A and C, but not chondroitin sulfate B. The enzyme cleaves, via an elimination mechanism, sulfated and non-sulfated polysaccharide chains containing  $\beta$ -(1-4) and  $\beta$ -(1-3) linkages between hexosamines and glucuronic acid residues. The reaction yields oligosaccharide products, mainly disaccharides, containing unsaturated uronic acids that can be detected by UV spectroscopy at 232 nm. The enzyme shows approximately equal activity with chondroitin sulfates A and C, while the activity observed with chondroitin sulfate B is approximately 7% of this value. This activity is most likely due to the presence of chondroitin sulfates A and C (10%) in the chondroitin sulfate B.

#### Chondroitinase AC from *Flavobacterium heparinum*

Chondroitin AC lyase  
[9047-57-8] E.C. 4.2.2.5

#### lyophilized powder, activity: 0.5-1.5 units/mg solid (using chondroitin sulfate A as substrate, also cleaves chondroitin sulfate C)

Contains potassium phosphate buffer salts and BSA as stabilizer.

#### composition, protein ~15% (Lowry)

One unit will cause a  $\Delta A_{232}$  of 1.0 per minute due to the release of unsaturated disaccharide from chondroitin sulfate A at pH 7.3 at 37 °C. Reaction volume: 3.1 mL (light path 1 cm).

Glycosaminoglycan (GAG) degradation enzymes ..... may contain trace amount

-20°C

C2780-5UN 5 units

### Chondroitinase C

Chondroitinase C cleaves chondroitin sulfate C producing tetrasaccharide plus an unsaturated 6-sulfated disaccharide (delta Di-6S). It also cleaves hyaluronic acid producing unsaturated nonsulfated disaccharide ( $\Delta$  Di-OS). Chondroitin sulfate A is also degraded producing oligosaccharides and delta Di-6S, but not delta Di-4S. Chondroitinase C cleaves the GalNAc bond of the pentasaccharides or hexasaccharides derived from the linkage region of chondroitin sulfate chains and tolerates sulfation of the C-4 or C-6 of the GalNAc residue and C-6 of the Gal residues, as well as 2-O-phosphorylation of the Xyl residue. In contrast, it does not act on a GalNAc-GlcA linkage when attached to a 4-O-sulfated Gal residue.

#### Chondroitinase C from *Flavobacterium heparinum*

Chondroitin C lyase  
[60184-91-0] E.C. 4.2.2.—

#### lyophilized powder, activity: $\geq 200$ units/mg solid

One unit will form 0.1  $\mu$ mole of unsaturated uronic acid per hr at pH 8.0 at 25 °C using chondroitin sulfate C as substrate.

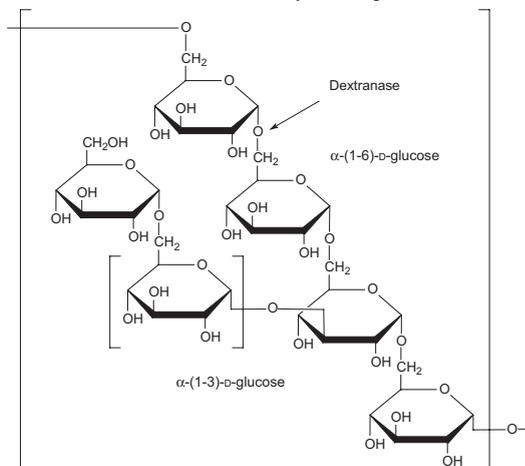
-20°C

C0954-75UN

75 units

## Dextranases

### Dextranase Specificity



Dextran is composed of approximately 95%  $\alpha$ -(1-6)-D-linkages. The remaining  $\alpha$ -(1-3) linkages account for the branching of dextran. Conflicting data on the branch lengths implies that the average branch length is less than three glucose units. However, other methods indicate branches of greater than 50 glucose units exist. Native dextran has been found to have a molecular weight (MW) in the range of 9 million to 500 million Da. Lower MW dextrans will exhibit slightly less branching and have a more narrow range of MW distribution. Dextrans with MW greater than 10,000 Da behave as if they are highly branched. As the MW increases, dextran molecules attain greater symmetry. Dextrans with MW of 2,000 to 10,000 Da exhibit the properties of an expandable coil. At MWs below 2,000 Da dextran is more rod-like.

## Dextranase

Dextranase catalyzes the endohydrolysis of  $\alpha$ -(1-6)-D-glucosidic linkages in dextran.

### Dextranase from *Chaetomium erraticum*

1,6- $\alpha$ -D-Glucan 6-glucohydrolase

E.C. 3.2.1.11

A product of Novozymes Corp.

#### solution

A fungal dextranase produced by submerged fermentation of *Chaetomium erraticum*.

Stable in the pH range of 3-7 and at temperatures up to approx. 70 °C. For most applications, the preferred conditions are pH 5-6 and a temperature of 50-60 °C.

[-8°C]

D0443-50ML	50 mL
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D0443-250ML	250 mL
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### Dextranase from *Penicillium sp.*

1,6- $\alpha$ -D-Glucan 6-glucohydrolase

E.C. 3.2.1.11

One unit will liberate 1.0  $\mu$ mole of isomaltose (measured as maltose) per min at pH 6.0 at 37 °C, using dextran as substrate.

► **Lyophilized powder, activity: 400-800 units/mg protein composition, protein 35% (Lowry)**

[-8°C]

D8144-500UN	500 units
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D8144-1KU	1,000 units
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► **Lyophilized powder, activity: 100-250 units/mg protein**

Partially purified, lyophilized powder

**composition, protein ~25% (Lowry)**

[-8°C]

D4668-500UN	500 units
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D4668-1KU	1,000 units
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► **Lyophilized powder, activity: 10-25 units/mg solid**

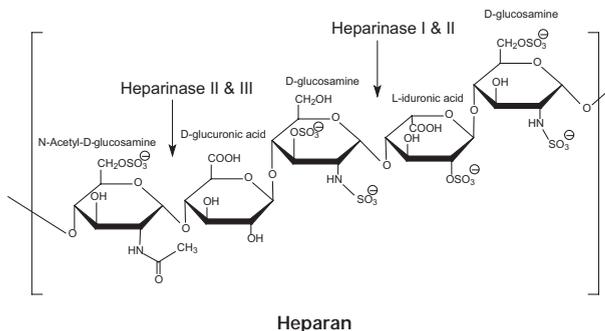
Crude

[-8°C]

D5884-5KU	5,000 units
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## Heparinases

### Heparinase Specificities



Heparan and heparin glycosaminoglycans (GAGs) are complex heterogeneous mixtures of repeating disaccharide units consisting of a uronic acid (D-glucuronic or L-iduronic acid) and D-glucosamine or N-acetyl-D-glucosamine. Various degrees of sulfation occur (at O and/or N) on each monosaccharide unit, ranging from zero to tri-sulfation. In general, heparan is less sulfated than heparin.

Heparinase selectively cleaves sulfated glycans containing  $\alpha$ -(1-4)-glycosidic linkages between the glucosamine and uronic acid residues in the heparin polymer. The cleavage proceeds via an elimination reaction, resulting in the formation of oligosaccharides containing unsaturated uronic acid residues (double bond between C4 and C5). These cleavage products can be detected by UV spectroscopy (232 nm). The three forms of heparinase (I, II, and III) have varying substrate specificities.

### Heparinase I

Heparinase I cleaves heparin and heparan sulfate (relative activity about 3:1) at the linkages between hexosamines and O-sulfated iduronic acids, yielding mainly disaccharides. The enzyme also cleaves the antithrombin III binding pentasaccharide domain in the heparin molecule.

#### Heparinase I from *Flavobacterium heparinum*

Heparinase; Heparin lyase I

[9025-39-2] E.C. 4.2.2.7

**Lyophilized powder stabilized with approx. 25% bovine serum albumin, activity: 200-600 units/mg solid**

mol wt 42.8 kDa

One unit will form 0.1  $\mu$ mole of unsaturated uronic acid per hr at pH 7.5 at 25 °C. One International Unit (I.U.) is equivalent to approx. 600 Sigma units.

[-20°C]

H2519-50UN	50 units
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H2519-100UN	100 units
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H2519-250UN	250 units
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## Heparinases

### Heparinase II

Heparinase II cleaves heparan sulfate, and to a lesser extent heparin (relative activity about 2:1), at the  $\alpha$ -(1-4) linkages between hexosamines and uronic acid residues (both glucuronic and iduronic), yielding mainly disaccharides.

#### Heparinase II from *Flavobacterium heparinum*

Heparin lyase II  
[149371-12-0]

Lyophilized powder stabilized with approx. 25% bovine serum albumin, lyophilized powder, activity: 100-300 units/mg solid mol wt 84.1 kDa

One unit will form 0.1  $\mu$ mol of unsaturated uronic acid per hr at pH 7.0 at 25 °C. One International Unit (I.U.) is equivalent to approx. 600 Sigma units.

-20°C

<b>H6512-10UN</b>	<b>10 units</b>
<b>H6512-25UN</b>	<b>25 units</b>
<b>H6512-100UN</b>	<b>100 units</b>

### Heparinase III

Heparinase III cleaves at the  $\alpha$ -(1-4) linkages between hexosamine and glucuronic acid residues in heparan sulfate, yielding mainly disaccharides. The enzyme is not active towards heparin. Sulfation at the 6-position of glucosamine inhibits cleavage by heparinase III,

#### Heparinase III from *Flavobacterium heparinum*

Heparin Lyase III; Heparitinase from *Flavobacterium heparinum*; Heparitinase I [37290-86-1] E.C. 4.2.2.8

**Lyophilized powder stabilized with approx. 25% bovine serum albumin, activity: 200-600 unit/mg solid**

Heparin-degrading lyase that recognizes heparin sulfate proteoglycan as its primary substrate.

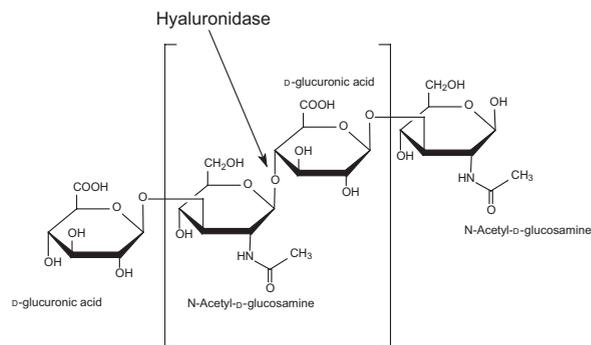
mol wt 70.8 kDa

One unit will form 0.1  $\mu$ mol of unsaturated uronic acid per hr at pH 7.5 at 25 °C. One International Unit (I.U.) is equivalent to approx. 600 Sigma units.

-20°C

<b>H8891-5UN</b>	<b>5 units</b>
<b>H8891-10UN</b>	<b>10 units</b>
<b>H8891-50UN</b>	<b>50 units</b>

## Hyaluronidases



**Hyaluronic Acid**

Composed of alternating residues of  $\beta$ -D-(1-3) glucuronic acid and  $\beta$ -D-(1-4)-N-acetylglucosamine

The mammalian hyaluronidases (EC 3.2.1.35) cleave hyaluronic acid and similar glycosaminoglycans by hydrolysis. The enzyme from *Streptomyces* (EC 4.2.2.1) is a lyase that catalyzes cleavage by an elimination reaction yielding a 4-deoxy-4,5-unsaturated oligosaccharides. Its specificity towards chondroitins and other glycosaminoglycans is unclear.

### Mammalian Hyaluronidase

The mammalian glycolytic hyaluronidase (EC 3.2.1.35) catalyzes the random hydrolysis of the 1-4 bond between N-acetyl-D-glucosamine and D-glucuronic acid in hyaluronic acid. It also hydrolyzes  $\beta$ -(1-4)-D-glycosidic linkages between N-acetyl-galactosamine or N-acetylgalactosamine sulfate and glucuronic acid in chondroitin sulfates A and C, and dermatan.

#### Hyaluronidase from bovine testes

Hyaluronate 4-glycanohydrolase; Hyaluronoglucosaminidase [37326-33-3] E.C. 3.2.1.35 EC No. 2534643

mol wt ~55 kDa (four subunits of 14 kDa each)

One unit is based on the change in absorbance at 600 nm (change in turbidity) of a USP reference standard hyaluronidase which is assayed concurrently with each lot.

► **Type I-S, lyophilized powder, activity: 400-1000 units/mg solid**

-20°C

<b>H3506-100MG</b>	<b>100 mg</b>
<b>H3506-500MG</b>	<b>500 mg</b>
<b>H3506-1G</b>	<b>1 g</b>
<b>H3506-5G</b>	<b>5 g</b>

► **Type IV-S, lyophilized powder (essentially salt-free), activity: 750-1500 units/mg solid**

-20°C

<b>H3884-50MG</b>	<b>50 mg</b>
<b>H3884-100MG</b>	<b>100 mg</b>
<b>H3884-500MG</b>	<b>500 mg</b>
<b>H3884-1G</b>	<b>1 g</b>

▶ **Type VIII, lyophilized powder, activity: ~300 units/mg**

Prepared from sterile filtered solution of Type I-S.

$-20^{\circ}\text{C}$

H3757-100MG	100 mg
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▶ **Type VI-S, lyophilized powder, activity: 3,000-15,000 units/mg solid**

Chromatographically purified

$-20^{\circ}\text{C}$

H3631-3KU	3,000 units
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H3631-15KU	15,000 units
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H3631-30KU	30,000 units
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### Hyaluronidase from sheep testes

Hyaluronate 4-glycanohydrolase; Hyaluronoglucosaminidase [37326-33-3]  
E.C. 3.2.1.35 EC No. 2534643

mol wt 55 kDa

One unit is based on the change in absorbance at 600 nm (change in turbidity) of a USP reference standard hyaluronidase which is assayed concurrently with each lot.

▶ **Type V, lyophilized powder, activity:  $\geq 1,500$  units/mg solid**

$-20^{\circ}\text{C}$

H6254-500MG	500 mg
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H6254-1G	1 g
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▶ **Type II, lyophilized powder, activity:  $\geq 300$  units/mg**

Lyophilized powder containing lactose

$-20^{\circ}\text{C}$

H2126-100MG	100 mg
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H2126-500MG	500 mg
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H2126-1G	1 g
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H2126-5G	5 g
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▶ **Type III, lyophilized powder, activity:  $\geq 500$  units/mg**

Lyophilized powder containing 20-50% lactose

$-20^{\circ}\text{C}$

H2251-100MG	100 mg
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H2251-500MG	500 mg
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H2251-1G	1 g
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H2251-5G	5 g
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## Hyaluronate Lyase (*Streptomyces Hyaluronidase*)

### Hyaluronidase from *Streptomyces hyalurolyticus*

Hyaluronate Lyase from *Streptomyces hyalurolyticus*  
[9001-54-1] E.C. 4.2.2.1 EC No. 2326141

#### lyophilized powder

Hyaluronate lyase cleaves hyaluronic acid at the  $\beta$ -D-GalNAc-(1-4)- $\beta$ -D-GlcA bond, yielding 3-(4-deoxy- $\beta$ -D-gluc-4-enuronosyl)-N-acetyl-D-glucosamine tetra- and hexasaccharides. Unlike other hyaluronidases, this enzyme is specific for hyaluronic acid and is inactive with chondroitin and chondroitin sulfate.<sup>1</sup>

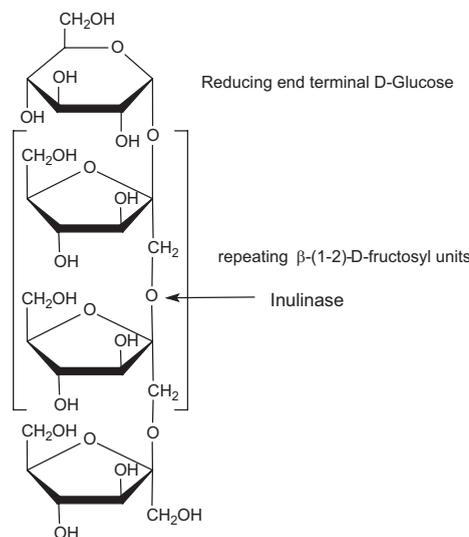
**Lit. cited:** 1. Ohya, T., and Kaneko, Y., *Biochim. Biophys. Acta* **198**, 607 (1970)

$-20^{\circ}\text{C}$

H1136-1AMP	1 amp
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## Inulinase

### Inulinase Specificity



### Inulin

Inulins are fructan oligosaccharides composed  $\alpha$ -D-glucopyranosyl- $[\beta$ -(2-1)-D-fructofuranosyl-D-fructofuranosides. Inulins can generally contain 2 to 140 fructose units.

### Inulinase from *Aspergillus niger*

Fructozyme L<sup>TM</sup>; Inulase  
E.C. 3.2.1.7

Inulinase catalyzes endohydrolysis of  $\beta$ -(2-1)-D-fructosidic linkages in inulin.

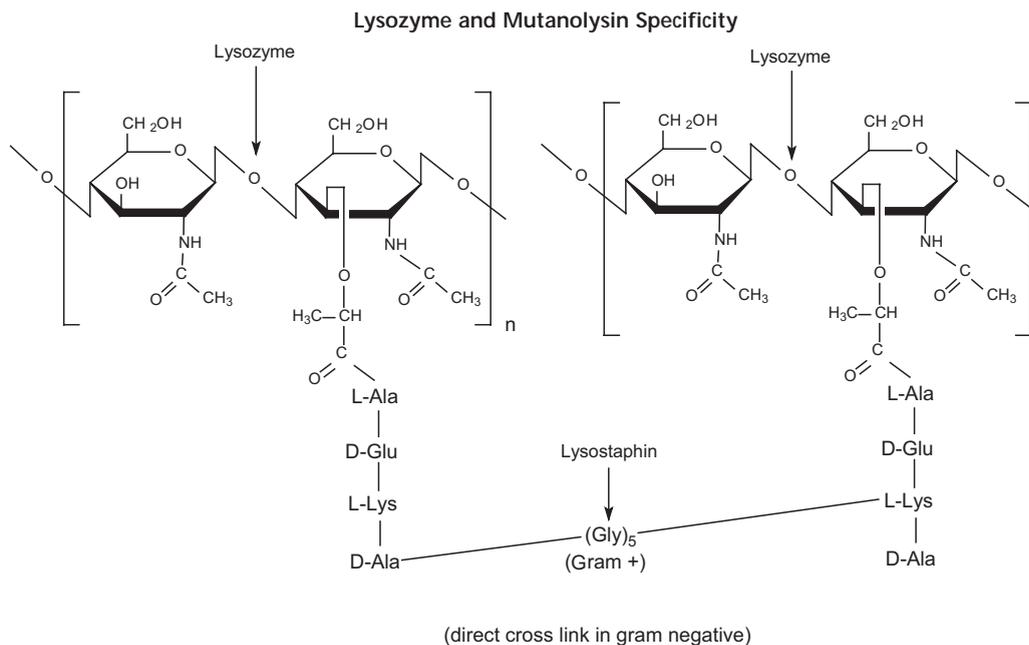
A product of Novozyme Corp.

$2-8^{\circ}\text{C}$

I2017-50ML	50 mL
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I2017-250ML	250 mL
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## Peptidoglycan Degrading Enzymes



### Peptidoglycan

Polymer of  $\beta$ -(1-4)-N-Acetyl-D-glucosamine units. Alternating residues are modified to form N-acetylmuramic acid with the addition of lactate to form branching links to the tetrapeptide.

## Lysozyme

### Lysozyme from chicken egg white

Mucopeptide N-acetylmuramoylhydrolase; Muramidase  
[12650-88-3] E.C. 3.2.1.17

Lysozyme hydrolyzes  $\beta$ -(1-4) linkages between N-acetylmuramic acid and N-acetyl-D-glucosamine residues in peptidoglycan and between N-acetyl-D-glucosamine residues in chitodextrin. Gram-positive cells are quite susceptible to this hydrolysis as their cell walls have a high proportion of peptidoglycan. Gram-negative bacteria are less susceptible due to the presence of an outer membrane and a lower proportion of peptidoglycan. However, these cells may be hydrolyzed in the presence of EDTA that chelates metal ions in the outer bacterial membrane.

The enzyme is active over a broad pH range (6.0 to 9.0). At pH 6.2, maximal activity is observed over a wider range of ionic strengths (0.02 to 0.100 M) than at pH 9.2 (0.01 to 0.06 M).

Used to prepare spheroplasts.

Single-chain mol wt 14.7 kDa

One unit will produce a  $\Delta A_{450}$  of 0.001 per min at pH 6.24 at 25 °C, using a suspension of *Micrococcus lysodeikticus* as substrate, in a 2.6 mL reaction mixture (1 cm light path).

#### ► lyophilized powder, Protein: ~95%, activity: ~50,000 units/mg protein

Dialyzed and lyophilized, containing buffer salts as sodium acetate and sodium chloride

3× crystallized

-20°C

L6876-1G	1 g
L6876-5G	5 g
L6876-10G	10 g
L6876-25G	25 g
L6876-100G	100 g

#### ► lyophilized powder, Protein: ~95%, activity: ~50,000 units/mg protein (E1%282)

##### Features and Benefits

- Highly purified by repeated crystallization and dialysis
- Each lot is use-tested for isolation of plasmid DNA from *E. coli* essentially salt-free

3× crystallized

-20°C

L7651-1G	1 g
L7651-5G	5 g
L7651-10G	10 g
L7651-25G	25 g
L7651-100G	100 g

#### ► aseptically filled, Lyophilized powder

Prepared from L6876

-20°C

L7773-50MG	50 mg
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**Lysozyme chloride form from chicken egg white**

Mucopolysaccharide N-acetylmuramoylhydrolase; Muramidase  
[9066-59-5] E.C. 3.2.1.17

**Grade VI, activity: ~60,000 units/mg protein**

Enzyme which breaks down the cell walls of bacteria; used to prepare spheroplasts.

Lyophilized powder containing sodium chloride and sodium acetate

mol wt ~14.3 kDa

**Composition protein ~90%**

3× Crystallized

One unit will produce a  $\Delta A_{450}$  of 0.001 per min at pH 6.24 at 25 °C, using a suspension of *Micrococcus lysodeikticus* as substrate, in a 2.6 mL reaction mixture (1 cm light path).

**-20°C**

<b>L2879-1G</b>	<b>1 g</b>
<b>L2879-5G</b>	<b>5 g</b>
<b>L2879-25G</b>	<b>25 g</b>

**Lysozyme from human milk**

Mucopolysaccharide N-acetylmuramoylhydrolase; Muramidase  
[12671-19-1] E.C. 3.2.1.17

**lyophilized powder, activity: ≥100,000 units/mg protein**

Lyophilized powder containing sodium phosphate and sodium chloride

**composition protein ~10%**

One unit will produce a  $\Delta A_{450}$  of 0.001 per min at pH 6.24 at 25 °C, using a suspension of *Micrococcus lysodeikticus* as substrate, in a 2.6 mL reaction mixture (1 cm light path).

**-20°C**

<b>L6394-25KU</b>	<b>25,000 units</b>
<b>L6394-100KU</b>	<b>100,000 units</b>

**Lysozyme from human neutrophils**

Mucopolysaccharide N-acetylmuramoylhydrolase; Muramidase  
[9001-63-2] E.C. 3.2.1.17

**≥95% (SDS-PAGE), lyophilized powder, activity: ≥100,000 units/mg protein**

Lyophilized from 50 mM sodium acetate, pH 6.0, with 100 mM NaCl

One unit will produce a  $\Delta A_{450}$  of 0.001 per min at pH 6.24 at 25 °C, using a suspension of *Micrococcus lysodeikticus* as substrate, in a 2.6 mL reaction mixture (1 cm light path).

**-20°C**

<b>L8402-.1MG</b>	<b>0.1 mg</b>
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**Mutanolysin****Mutanolysin from *Streptomyces globisporus* ATCC 21553**

[55466-22-3]

Mutanolysin is an N-acetylmuramidase. Like lysozyme, it is a muralytic enzyme that cleaves the  $\beta$ -N-acetylmuramyl-(1-4)-N-acetylglucosamine linkage of the bacterial cell wall polymer peptidoglycanopolysaccharide. Its carboxy terminal moieties are involved in the recognition and binding of unique cell wall polymers. Mutanolysin lyses *Listeria* and other Gram-positive bacteria such as *Lactobacillus* and *Lactococcus*.

Provides gentle cell lysis for the isolation of easily degradable biomolecules and RNA from bacteria. It has been used in the formation of spheroplasts for isolation of DNA.

mol wt 23 kDa

One unit will produce a  $\Delta A_{600}$  of 0.01 per minute at pH 6.0 at 37 °C in a 1 mL volume using a suspension of *Streptococcus faecalis* cell wall as substrate.

**▶ lyophilized powder, activity: ≥4000 units/mg protein (biuret),**

Chromatographically purified

Lyophilized powder containing Ficoll® and sodium succinate buffer salts

**-20°C**

<b>M9901-1KU</b>	<b>1,000 units</b>
<b>M9901-5KU</b>	<b>5,000 units</b>
<b>M9901-10KU</b>	<b>10,000 units</b>
<b>M9901-50KU</b>	<b>50,000 units</b>

**▶ aseptically filled, lyophilized powder, activity: ≥4000 units/mg protein (biuret)**

Lyophilized powder containing Ficoll® and sodium succinate buffer salts

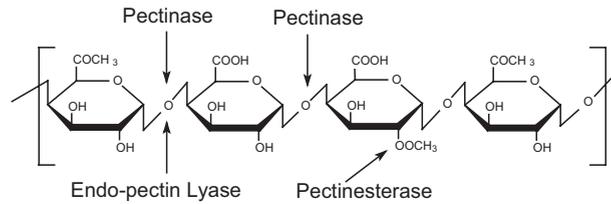
Prepared from M9901

**-20°C**

<b>M4782-5KU</b>	<b>5,000 units</b>
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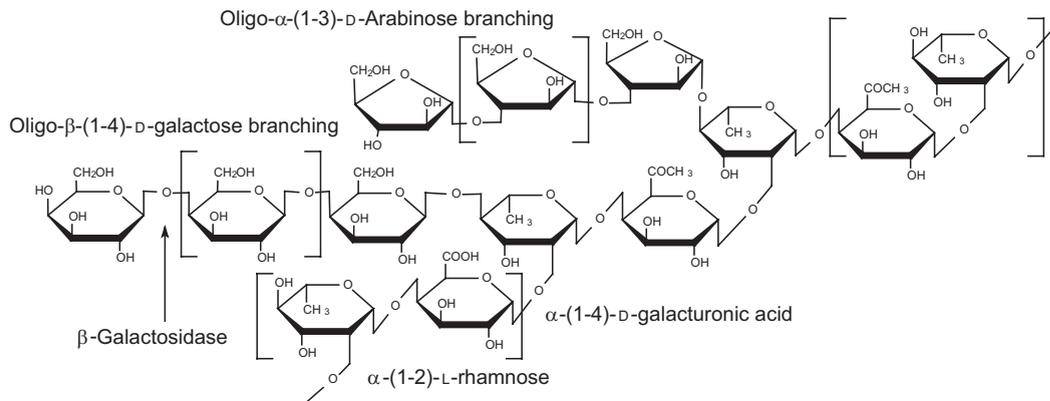
# Pectin Degrading Enzymes

## Pectinase and Pectinesterase Specificities



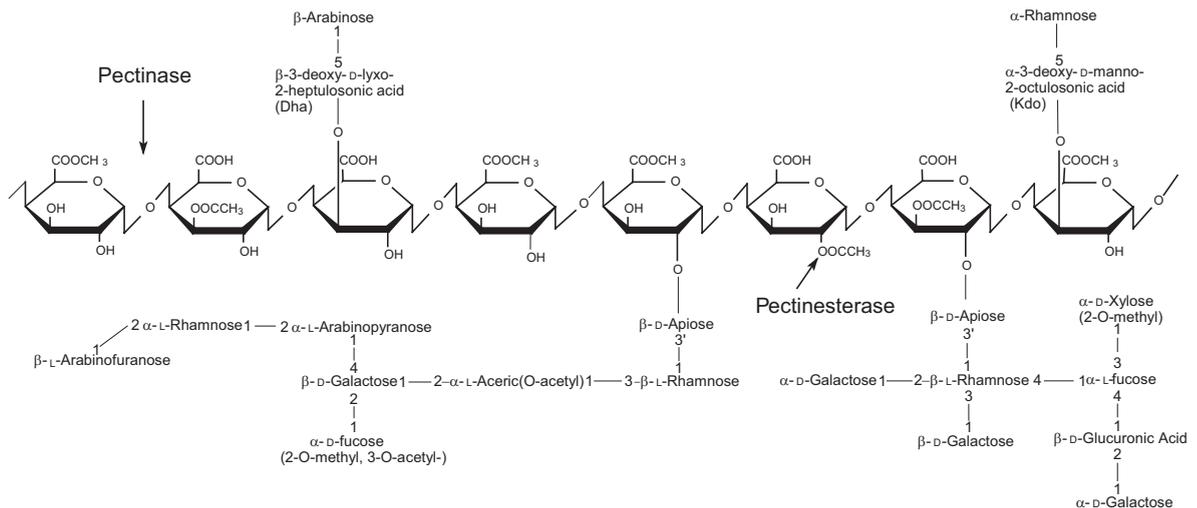
### Homogalacturonan

Poly- $\alpha$ -(1-4)-D-galacturonic acid backbone with random-partial methylation and acetylation



### Rhamnogalacturonan I

Alternating  $\alpha$ -(1-2)-L-rhamnosyl-  $\alpha$ -(1-4)-D-galacturonosyl backbone with two types of branching composed of arabinofuranose or galactose oligomers



### Rhamnogalacturonan II

Poly- $\alpha$ -(1-4)-D-galacturonic acid backbone with random-partial methylation, acetylation and four different types of branching

Pectins are complex branched heteropolysaccharides primarily containing an  $\alpha$ -(1-4) polygalacturonic acid backbone which can be randomly acetylated and methylated. Three different pectins have been isolated from plant cell walls. Homogalacturonans are composed of the simple  $\alpha$ -(1-4) polygalacturonic acid backbone. Substituted homogalacturonans are modifications of this backbone with  $\beta$ -D-xylose branching at C3, or apiofuranose substitutions in the backbone with  $\beta$ -D-Apiofuranosyl-(1,3')- $\beta$ -D-Apiofuranose branching. Rhamnogalacturonan I contains alternating  $\alpha$ -(1-4) galacturonosyl and  $\alpha$ -(1-2) rhamnosyl residues, with primarily oligo  $\alpha$ -(1-3) arabinose and oligo  $\beta$ -(1-4) galactose branching. Rhamnogalacturonan II is composed of the simple  $\alpha$ -(1-4) polygalacturonic acid backbone with complex branching composed of up to 11 different monosaccharide types.

## Pectinase

Pectinase catalyzes the random hydrolysis of  $\alpha$ -(1-4)-D-galactosiduronic linkages in pectin and other galacturonans.

### Pectinase from *Aspergillus aculeatus*

**Pectinex Ultra SPL®**

**aqueous solution, activity:  $\geq 26,000$  units/mL**

Highly active pectolytic enzyme preparation produced by a selected strain of *Aspergillus aculeatus*

A product of Novozyme Corp.

2-8°C

P2611-50ML	50 mL
P2611-250ML	250 mL

### Pectinase from *Aspergillus niger*

**Pectinex® 3XL**

**aqueous solution**

Pectolytic enzyme preparation produced from a selected strain of *Aspergillus niger*: contains mainly pectintranseliminase, polygalacturonase, and pectinesterase and small amounts of hemicellulases and cellulases.

A product of Novozyme Corp.

2-8°C

P2736-50ML	50 mL
P2736-250ML	250 mL

### Pectinase solution from *Aspergillus niger*

Polygalacturonase solution from *Aspergillus niger*:

Poly-(1,4- $\alpha$ -D-galacturonide) glycanohydrolase

[9032-75-1] E.C. 3.2.1.15

Used in plant protoplast preparation to digest cell wall prior to organelle isolation.

Solution in 40% glycerol

One unit will liberate 1.0  $\mu$ mole of galacturonic acid from polygalacturonic acid per min at pH 4.0 at 25 °C.

▶ **aqueous glycerol solution, activity:  $\geq 5$  units/mg protein (Lowry)**

2-8°C

P4716-5KU	5,000 units
P4716-10KU	10,000 units
P4716-25KU	25,000 units
P4716-100KU	100,000 units

▶ **plant cell culture tested, aqueous glycerol solution, activity:  $\geq 5$  units/mg protein (Lowry)**

2-8°C

P0690-10KU	10,000 units
P0690-25KU	25,000 units

### Pectinase from *Rhizopus* sp.

Macerozyme R-10; Polygalacturonase; Poly-(1,4- $\alpha$ -D-galacturonide) glycanohydrolase

[9032-75-1] E.C. 3.2.1.15 EC No. 2328856

Used in plant protoplast preparation to digest cell wall prior to organelle isolation.

One unit will liberate 1.0  $\mu$ mole of galacturonic acid from polygalacturonic acid per min at pH 4.0 at 25 °C.

▶ **powder, activity: 400-800 units/g solid**

Crude source of pectinase activity, also containing cellulase and hemicellulase activities.

-20°C

P2401-500UN	500 units
P2401-1KU	1,000 units
P2401-5KU	5,000 units

▶ **plant cell culture tested, crude powder activity: 400-800 units/g solid**

-20°C

P4300-1KU	1,000 units
P4300-5KU	5,000 units

Visit the  
**Enzyme Explorer  
Assay Library**

Features over 600 detailed procedures for measuring enzyme activity and related metabolites. The Library is the result of over ten years of in-house process development by Sigma-Aldrich scientists.

[sigma-aldrich.com/enzymeexplorer](http://sigma-aldrich.com/enzymeexplorer)

## Pectin Degrading Enzymes

### Pectinesterase

Pectinesterase catalyzes the hydrolysis of the methyl esters of pectin to yield pectate and methanol.

#### Pectinesterase from orange peel

Pectin methylsterase; Pectin pectylhydrolase  
[9025-98-3] E.C. 3.1.1.11 EC No. 2328070

**lyophilized powder, activity:**  $\geq 150$  units/mg protein

Contains  $(\text{NH}_4)_2\text{SO}_4$  and sodium chloride

**Composition protein 20-50% (biuret)**

One unit will release 1.0 microequivalent of acid from pectin per min at pH 7.5 at 30 °C.

Protein determined by biuret.

2-8°C

P5400-1KU	1,000 units
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### Pectolyase

Pectolyase catalyzes the eliminative cleavage of  $\alpha$ -(1-4)-D-galacturonan methyl ester to give oligosaccharides with 4-deoxy-6-O-methyl- $\alpha$ -D-galact-4-enuronosyl groups at their non-reducing ends.

#### Pectolyase from *Aspergillus japonicus*

E.C. 3.2.1.15

Reported to contain two types of pectinase, endopolygalacturonase (EC 3.2.1.15), endo-pectin lyase (EC 4.2.2.10) and a maceration stimulating factor.

Used in plant protoplast preparation to digest cell wall prior to organelle isolation.

Lyophilized powder containing lactose

One unit will liberate 1.0  $\mu$ mole of galacturonic acid from polygalacturonic acid per min at pH 5.5 at 25 °C.

► **lyophilized powder, activity:**  $\geq 0.3$  units/mg solid

2-8°C

P3026-100MG	100 mg
P3026-250MG	250 mg
P3026-1G	1 g

► **plant cell culture tested, lyophilized powder activity:**  $\geq 0.3$  unit/mg solid

**Composition protein ~60% (Lowry)**

2-8°C

P5936-100MG	100 mg
P5936-250MG	250 mg
P5936-1G	1 g

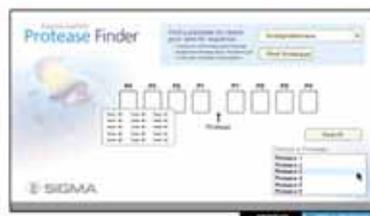
## The Enzyme Explorer The Protease Finder

Cleave proteins exactly where you want to with Sigma's new Protease Finder. The Protease Finder will identify the protease needed to cleave a specific peptide sequence at your desired location.

#### Simple to use:

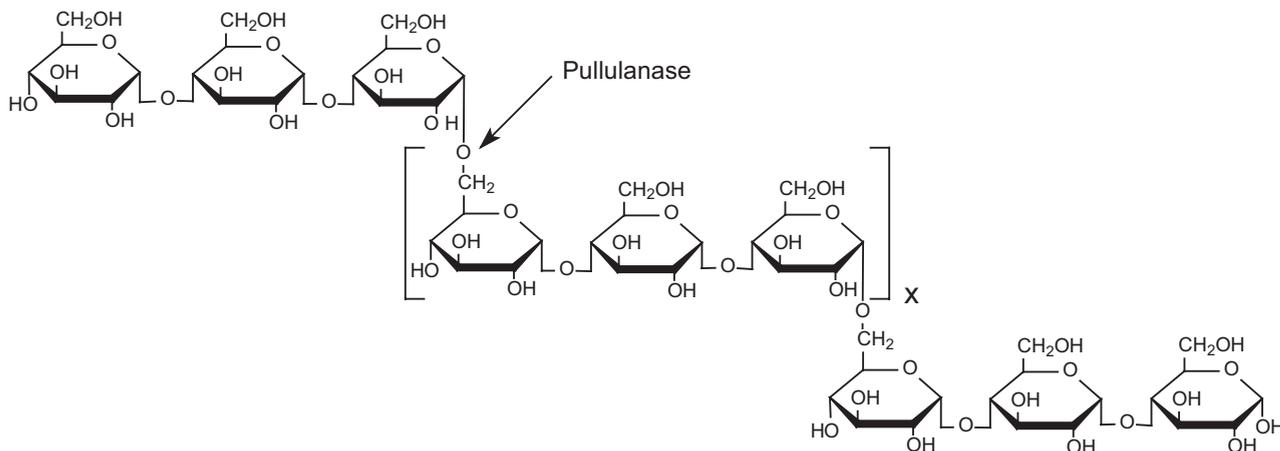
- Select either Endo- or Exoproteolytic cleavage.
- Enter your protein sequence into the positional boxes
- Submit the request to instantly receive the protease(s) capable of the cleavage.

[sigma-aldrich.com/proteasefinder](http://sigma-aldrich.com/proteasefinder)



## Pullulanase

### Pullulanase Specificity



### Pullulan

Linear polymer of  $\alpha$ -(1-6)-linked maltotriose units

Pullulanase catalyzes the hydrolysis of  $\alpha$ -(1-6)-D-glucosidic linkages in pullulan (a linear polymer of  $\alpha$ -(1-6)-linked maltotriose units), and, similar to isoamylase, in amylopectin and glycogen, and the  $\alpha$ - and  $\beta$ - limit dextrans of amylopectin and glycogen.

### Pullulanase from *Bacillus acidopullulyticus*

Pullulan 6-glucano-hydrolase  
[9075-68-7] E.C. 3.2.1.41

#### Promozyme® 400 L

aqueous solution,  $\geq 400$  units/mL

Heat-stable debranching enzyme obtained from a selected strain of *Bacillus acidopullulyticus*, and belongs to the group of debranching enzymes known as pullulanases.

One unit is defined as the amount of enzyme which hydrolyzes pullulan, liberating reducing carbohydrate with a reducing power equivalent to 1.0  $\mu$ mole glucose per minute at pH 5.0 and 40 °C.  
density ..... 1.25 g/mL, 25 °C  
A product of Novozymes Corp.

2-8°C

P2986-50ML	50 mL
P2986-250ML	250 mL

### Pullulanase from *Klebsiella pneumoniae*

Amylopectin 6-gluconohydrolase; Limit dextrinase  
[9075-68-7] E.C. 3.2.1.41

One unit will liberate 1.0  $\mu$ mole of maltotriose (measured as glucose) from pullulan per min at pH 5.0 at 25 °C.

#### ▶ lyophilized powder, activity: 10-30 units/mg protein

Lyophilized powder containing potassium phosphate buffer salts and stabilizer

#### Composition protein ~10% (Lowry)

-20°C

P1067-100UN	100 units
P1067-250UN	250 units

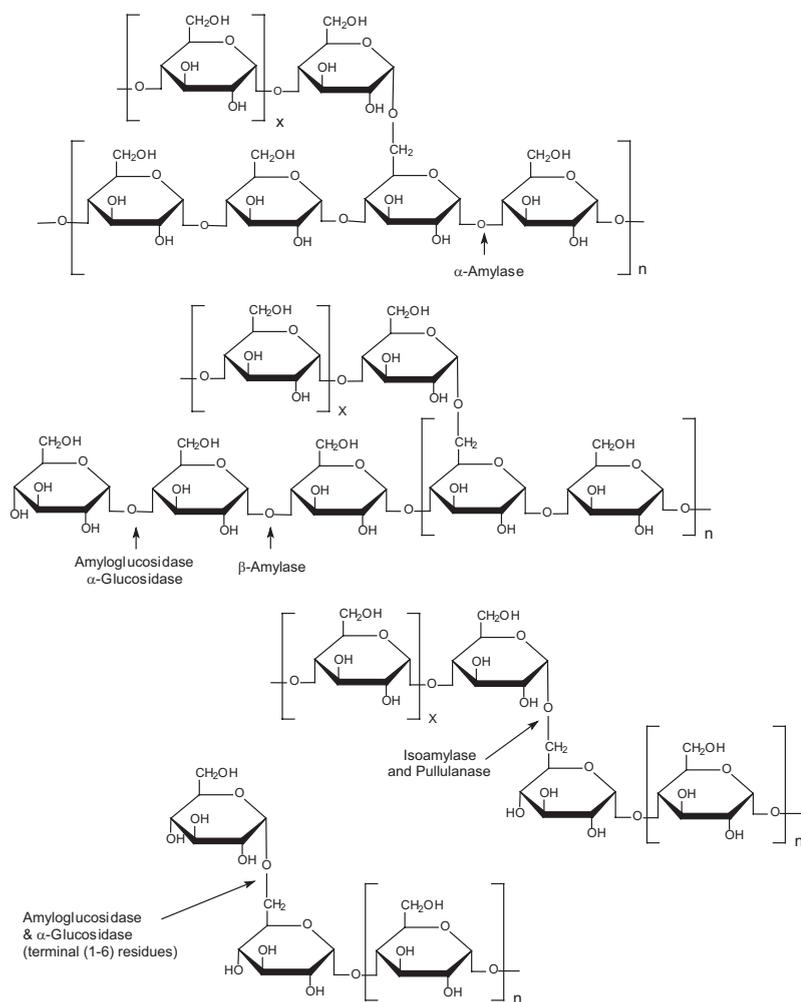
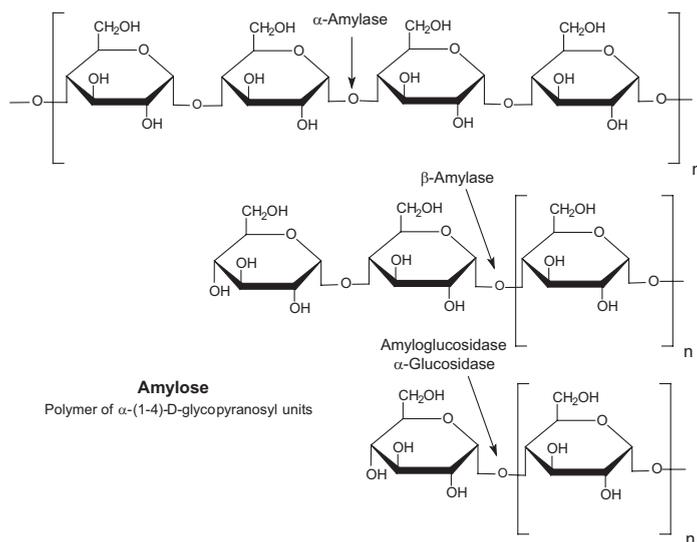
#### ▶ ammonium sulfate suspension, activity: $\geq 5$ units/mg protein (biuret)

Suspension in 3.2 M  $(\text{NH}_4)_2\text{SO}_4$  solution, pH 6.2  
Highly purified by a modification of ion exchange chromatography.

2-8°C

P5420-100UN	100 units
P5420-250UN	250 units

## Starch and Glycogen Degrading Enzymes



**$\alpha$ -Amylase**

$\alpha$ -Amylase catalyzes the endohydrolysis of  $\alpha$ -(1-4)-D-glucosidic linkages in polysaccharides containing three or more  $\alpha$ -(1-4)-linked D-glucose units.

 **$\alpha$ -Amylase from *Bacillus licheniformis***

1,4- $\alpha$ -D-Glucan-glucanohydrolase  
[9000-85-5] E.C. 3.2.1.1

**lyophilized powder, activity: 500-1,500 units/mg protein, ~95% (SDS-PAGE)**

Lyophilized powder containing potassium

**Composition** protein ~70% (BCA)

One unit will liberate 1.0 mg of maltose from starch in 3 min at pH 6.9 at 20 °C.

heat ..... (Stable)

**2-8°C**

A4551-100MG	100 mg
A4551-1G	1 g

 **$\alpha$ -Amylase from *Aspergillus oryzae***

1,4- $\alpha$ -D-Glucan-glucanohydrolase  
[9001-19-8] E.C. 3.2.1.1 EC No. 2325881

**▶ lyophilized powder, activity: 150-250 units/mg protein (biuret) Crude**

One unit will liberate 1.0 mg of maltose from starch in 3 min at pH 6.9 at 20 °C.

**-20°C**

A6211-250KU	250,000 units
A6211-1MU	1,000,000 units
A6211-5MU	5,000,000 units

**▶ Fungamyl® 800L aqueous solution, activity:  $\geq$ 0.8 units/g**

A product of Novozyme Corp.

**2-8°C**

A8220-50ML	50 mL
A8220-250ML	250 mL

**Taka-Diastase from *Aspergillus oryzae***

$\alpha$ -Amylase; 1,4- $\alpha$ -D-Glucan-glucanohydrolase; Taka-Amylase A  
[9001-19-8] E.C. 3.2.1.1 EC No. 2325881

**BioChemika, powder, activity: ~100 units/mg**

One unit corresponds to the amount of enzyme which liberates 1  $\mu$ mol maltose per minute at pH 6.0 and 25 °C (starch according to Zulkowsky, Catalog No. 85642, as substrate).

**2-8°C**

86247-25G	25 g
86247-100G	100 g

 **$\alpha$ -Amylase from *Bacillus sp.***

1,4- $\alpha$ -D-Glucan-glucanohydrolase from *Bacillus sp.*  
[9000-90-2] E.C. 3.2.1.1

**▶ powder, activity:  $\geq$ 400 units/mg protein (Lowry)**

Contains starch as an extender.

One unit will liberate 1.0 mg of maltose from starch in 3 min at pH 6.9 at 20 °C.

**-20°C**

A6814-1MU	1,000,000 units
A6814-5MU	5,000,000 units
A6814-25MU	25,000,000 units

**▶ Duramyl®, liquid, activity:  $\geq$ 300 units/g**

Protein-engineered  $\alpha$ -amylase produced by submerged fermentation of a genetically modified species of *Bacillus*.

A product of Novozyme Corp.

**2-8°C**

A7720-50ML	50 mL
A7720-250ML	250 mL

**▶ Type II-A, lyophilized powder, activity: 1,500-3,000 units/mg protein (biuret)**

mol wt 50-55 kDa by SDS-PAGE

4 $\times$  crystallized

One unit will liberate 1.0 mg of maltose from starch in 3 min at pH 6.9 at 20 °C.

**-20°C**

A6380-25MG	25 mg
A6380-100MG	100 mg
A6380-250MG	250 mg
A6380-500MG	500 mg
A6380-1G	1 g

 **$\alpha$ -Amylase from *Bacillus amyloliquefaciens***

BAN™ 240L; 1,4- $\alpha$ -D-Glucan glucanohydrolase  
[9000-85-5] E.C. 3.2.1.1. EC No. 2325656

**liquid, activity:  $\geq$ 250 units/g**

This enzyme is active at high temperatures (70-90 °C).

mol wt 55 kDa

One unit is the amount of enzyme which dextrinizes 5.26 g dry starch per hour under standard conditions.

A product of Novozyme Corp.

**2-8°C**

A7595-50ML	50 mL
A7595-250ML	250 mL

## Starch and Glycogen Degrading Enzymes

### $\alpha$ -Amylase from *Bacillus licheniformis*

1,4- $\alpha$ -D-Glucan-glucohydrolase  
[9000-85-5] E.C. 3.2.1.1  
Termamyl® 120

► **Type XII-A, saline solution, activity: 500-1,000 units/mg protein (biuret)**

Aqueous solution containing approx. 15% sodium chloride and 25% sucrose.

One unit will liberate 1.0 mg of maltose from starch in 3 min at pH 6.9 at 20 °C.

Reported to be heat stable at temperatures as high as ~90 °C.

A product of Novozyme Corp.

2-8°C

A3403-500KU	500,000 units
A3403-1MU	1,000,000 units
A3403-5MU	5,000,000 units

► **suitable for determination of starch (Kit STA-20)**

2-8°C

A4582-.5ML	0.5 mL
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### $\alpha$ -Amylase, heat-stable

$\alpha$ -Amylase: 1,4- $\alpha$ -D-Glucan-glucohydrolase  
[9000-85-5] E.C. 3.2.1.1

**solution, For use in Total Dietary Fiber Assay, TDF-100A**

2-8°C

A3306-10ML	10 mL
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### $\alpha$ -Amylase from barley malt

1,4- $\alpha$ -D-Glucan-glucohydrolase  
[9000-90-2] E.C. 3.2.1.1

**Type VIII-A, powder**

$\alpha$ -amylase activity:  $\geq 1$  unit/mg solid

$\beta$ -amylase activity:  $\geq 1$  unit/mg solid

contains lactose as standardization of activity

Package size based on  $\alpha$ -amylase activity

One unit will liberate 1.0 mg of maltose from starch in 3 min at pH 6.9 at 20 °C.

2-8°C

A2771-10KU	10 KU
A2771-50KU	50 KU

### $\alpha$ -Amylase from human pancreas

[9000-90-2] E.C. 3.2.1.1 EC No. 2325656

**lyophilized powder, activity:  $\geq 100$  units/mg protein**

Lyophilized from Tris buffer containing NaCl and CaCl<sub>2</sub>.

purified by 3 $\times$  crystallization

Prepared by modified method of Levitzki et al.

One unit will liberate 1.0 mg of maltose from starch in 3 min at pH 6.9 at 20 °C.

-20°C

A9972-100UG	100 $\mu$ g
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### $\alpha$ -Amylase from human saliva

1,4- $\alpha$ -D-Glucan-glucohydrolase  
[9000-90-2] E.C. 3.2.1.1 EC No. 2325656

**composition**

Protein ~10% (biuret)

One unit will liberate 1.0 mg of maltose from starch in 3 min at pH 6.9 at 20 °C.

► **Type XIII-A, lyophilized powder, activity: 300-1,000 units/mg protein**

Lyophilized powder containing (NH<sub>4</sub>)<sub>2</sub>SO<sub>4</sub> and sodium citrate.

-20°C

A1031-1KU	1,000 units
A1031-5KU	5,000 units

► **Type IX-A, lyophilized powder, activity: 1,000-3,000 units/mg protein**

Lyophilized powder containing (NH<sub>4</sub>)<sub>2</sub>SO<sub>4</sub> and sodium citrate  
Chromatographically purified

-20°C

A0521-100UN	100 units
A0521-500UN	500 units
A0521-2.5KU	2,500 units

### $\alpha$ -Amylase from porcine pancreas

E.C. 3.2.1.1

Molecular Weight: 51-54 kDa.

$\alpha$ -Amylase isolated from porcine pancreas is a glycoprotein. It is a single polypeptide chain of approximately 475 residues containing two SH groups and four disulfide bridges and a tightly bound Ca<sup>2+</sup> necessary for stability. Chloride ions are necessary for activity and stability. The pH range for activity is 5.5 to 8.0, with the pH optimum at 7.

One unit will liberate 1.0 mg of maltose from starch in 3 min at pH 6.9 at 20 °C.

► **Type VI-B, activity: 10-30 units/mg solid**

Contains lactose

Package size based on  $\alpha$ -amylase activity

2-8°C

A3176-500KU	500,000 units
A3176-1MU	1,000,000 units
A3176-2.5MU	2,500,000 units
A3176-5MU	5,000,000 units
A3176-10MU	10,000,000 units

► **Type I-A, DFP Treated, saline suspension, activity: 700-1400 units/mg protein**

Suspension in 2.9 M NaCl solution containing 3 mM CaCl<sub>2</sub>

DFP treated. 2 $\times$  crystallized

2-8°C

A6255-10MG	10 mg
A6255-25MG	25 mg
A6255-100MG	100 mg

▶ **Type VII-A, DFP treated, ammonium sulfate suspension,**

**activity:  $\geq 500$  units/mg protein**

Suspension in 3.2 M  $(\text{NH}_4)_2\text{SO}_4$ , pH 6.1

**2-8°C**

**A2643-10MG 10 mg**

**A2643-50MG 50 mg**

▶ **Type I-A, PMSF treated, saline suspension,**

**activity: 700-1400 units/mg protein**

Suspension in 2.9 M NaCl solution containing 3 mM  $\text{CaCl}_2$ .

2× crystallized

**2-8°C**

**A4268-25MG 25 mg**

**A4268-100MG 100 mg**

### $\beta$ -Amylase

$\beta$ -Amylase catalyzes the exo-hydrolysis of  $\alpha$ -(1-4)-D-glucosidic linkages in polysaccharides resulting in the successive liberation of maltose units from the non-reducing ends of the chains.

#### $\beta$ -Amylase from barley

1,4- $\alpha$ -D-Glucan maltohydrolase  
[9000-91-3] E.C. 3.2.1.2 EC No. 2325661

**Type II-B, activity: 20-80 units/mg protein (biuret)**

Crude

One unit will liberate 1.0 mg of maltose from starch in 3 min at pH 4.8 at 20 °C.

**2-8°C**

**A7130-10KU 10,000 units**

**A7130-50KU 50,000 units**

**A7130-250KU 250,000 units**

#### $\beta$ -Amylase from sweet potato

1,4- $\alpha$ -D-Glucan maltohydrolase  
[9000-91-3] E.C. 3.2.1.2 EC No. 2325661

**Type I-B, ammonium sulfate suspension, activity:  
750-1,000 units/mg protein**

Crystalline suspension in 2.3 M  $(\text{NH}_4)_2\text{SO}_4$

One unit will liberate 1.0 mg of maltose from starch in 3 min at pH 4.8 at 20 °C.

**2-8°C**

**A7005-10KU 10,000 units**

**A7005-25KU 25,000 units**

**A7005-50KU 50,000 units**

**A7005-100KU 100,000 units**

### Amyloglucosidase

Amyloglucosidase catalyzes the hydrolysis of terminal  $\alpha$ -(1-4)-linked D-glucose residues successively from the non-reducing ends of maltooligo- and polysaccharides with release of  $\beta$ -D-glucose. Most forms of the enzyme can rapidly hydrolyze  $\alpha$ -(1-6)-D-glucosidic bonds when the next bond in the sequence is 1,4- and some preparations of this enzyme hydrolyze 1,6- and  $\alpha$ -(1-3)-D-glucosidic bonds in other polysaccharides.

#### Amyloglucosidase from *Aspergillus niger*

Exo-1,4- $\alpha$ -glucosidase; 1,4- $\alpha$ -D-Glucan glucohydrolase; Glucoamylase  
[9032-08-0] E.C. 3.2.1.3 EC No. 2328772

▶ **lyophilized powder, activity:  $\geq 80$  units/mg protein (biuret)**

Lyophilized powder containing less than 0.02% glucose

One unit will liberate 1.0 mg of glucose from starch in 3 min at pH 4.5 at 55 °C.

**-20°C**

**A7420-5MG 5 mg**

**A7420-25MG 25 mg**

**A7420-100MG 100 mg**

▶ **AMG™ 300L,  $\geq 300$  units/mL**

Stabilized with glucose

aqueous solution

density ..... ~1.2 g/mL, 25 °C

A product of Novozymes Corp.

**2-8°C**

**A7095-50ML 50 mL**

▶ **aqueous glucose solution activity:  $\geq 5000$  units/mL**

Solution in 1 M glucose containing 0.5% sodium benzoate as preservative

One unit will liberate 1.0 mg of glucose from starch in 3 min at pH 4.5 at 55 °C.

**2-8°C**

**A3042-50ML 50 mL**

▶ **ammonium sulfate suspension, activity:**

**$\geq 40$  units/mg protein** Suspension in 3.2 M  $(\text{NH}_4)_2\text{SO}_4$  solution, pH approx. 6.0

One unit will liberate 1.0 mg of glucose from starch in 3 min at pH 4.5 at 55 °C.

**2-8°C**

**A1602-25MG 25 mg**

**A1602-100MG 100 mg**

## Starch and Glycogen Degrading Enzymes

### Amyloglucosidase from *Candida tsukubaensis*

Exo-1,4- $\alpha$ -glucosidase; 1,4- $\alpha$ -D-Glucan glucohydrolase; Glucoamylase [9032-08-0] E.C. 3.2.1.3

**ammonium sulfate suspension, activity: 50-150 units/mg protein**

An acid-stable amyloglucosidase, maintaining high activity at pH values down to 2.5.

Suspension in 3.2 M  $(\text{NH}_4)_2\text{SO}_4$ , pH approx. 5.5.

One unit will liberate 1.0 mg of glucose from starch in 3 min at pH 4.5 at 55 °C.

2-8°C

**A2330-25UN** 25 units

**A2330-100UN** 100 units

### Amyloglucosidase from *Rhizopus* sp.

Exo-1,4- $\alpha$ -glucosidase; 1,4- $\alpha$ -D-Glucan glucohydrolase; Glucoamylase [9032-08-0] E.C. 3.2.1.3 EC No. 2328772

**activity:  $\geq 40,000$  units/g solid**

Lyophilized salt-free powder

**Composition protein ~35% (biuret)**

One unit will liberate 1.0 mg of glucose from starch in 3 min at pH 4.5 at 55 °C.

2-8°C

**A9228-1G** 1 g

### $\alpha$ -Glucosidase

$\alpha$ -Glucosidase catalyzes the hydrolysis of terminal  $\alpha$ -(1-4)-linked D-glucose residues successively from the non-reducing ends of maltooligo- and to a lesser extent polysaccharides with release of  $\beta$ -D-glucose. Most forms of the enzyme can slowly hydrolyze  $\alpha$ -(1-6)-D-glucosidic bonds.

### $\alpha$ -Glucosidase from *Bacillus stearothermophilus*

$\alpha$ -D-Glucoside glucohydrolase; Maltase [9001-42-7] E.C. 3.2.1.20

**lyophilized powder, activity:  $\geq 50$  units/mg protein**

Lyophilized powder containing potassium phosphate buffer salt

One unit will liberate 1.0  $\mu$ mole of D-glucose from p-nitrophenyl  $\alpha$ -D-glucoside per min at pH 6.8 at 37 °C.

Protein determined by biuret.

$\beta$ -Glucosidase,  $\alpha$ -galactosidase and  $\beta$ -galactosidase .....  $\leq 0.1\%$

2-8°C

**G3651-250UN** 250 units

### $\alpha$ -Glucosidase from rice

$\alpha$ -D-Glucoside glucohydrolase; Maltase [9001-42-7] E.C. 3.2.1.20

**Type V, ammonium sulfate suspension, activity: 40-80 units/mg protein**

Suspension in 2.8 M  $(\text{NH}_4)_2\text{SO}_4$  solution

One unit will convert 1.0  $\mu$ mole of maltose to 2.0  $\mu$ moles of D-glucose per min at pH 4.0 at 37 °C.

Protein determined by biuret.

2-8°C

**G9259-100UN** 100 units

### $\alpha$ -Glucosidase from *Saccharomyces cerevisiae*

$\alpha$ -D-Glucosidase;  $\alpha$ -D-Glucoside glucohydrolase; Maltase from yeast [9001-42-7] E.C. 3.2.1.20 EC No. 2326047

For the determination of  $\alpha$ -amylase and the synthesis of various 1'-osucrose and 1-O-fructose esters

Protein determined by biuret.

**▶ recombinant, expressed in unspecified host, lyophilized powder, activity:  $\geq 125$  units/mg protein**

Lyophilized powder containing potassium phosphate buffer salt pH 7.15 and approx. 70% lactose

One unit will liberate 1.0  $\mu$ mole of D-glucose from p-nitrophenyl  $\alpha$ -D-glucoside

per min at pH 6.8 at 37 °C.

2-8°C

**G0660-750UN** 750 units

**▶ Type I, lyophilized powder, activity:  $\geq 10$  units/mg protein (using p-nitrophenyl  $\alpha$ -D-glucoside as substrate.)**

contains phosphate buffer salts and EDTA as balance composition

Protein ~50%

Sold on basis of p-nitrophenyl  $\alpha$ -D-glucoside units.

One unit will liberate 1.0  $\mu$ mole of D-glucose from p-nitrophenyl  $\alpha$ -D-glucoside per min at pH 6.8 at 37 °C.

$\beta$ -Glucosidase,  $\alpha$ -galactosidase and  $\beta$ -galactosidase .....  $\leq 0.1\%$

-20°C

**G5003-100UN** 100 units

**G5003-1KU** 1,000 units

## Kits for Carbohydrate Analysis

Sigma manufactures several unique enzymatic-based kits for the quantitation of important carbohydrates. These kits utilize spectrophotometric, and gravimetric detection making them easy-to-use, yielding high sensitivity, and consistent results.

### Total Dietary Fiber Assay Kit, Cat. No. TDF100A-1KT

sufficient for ~100 assays

For the determination of total dietary fiber. Uses a combination of enzymatic and gravimetric methods to analyze samples of dried, defatted foods to determine soluble fiber, protein, and ash content. This procedure is based on the method published by AOAC.<sup>1</sup>

#### Reference:

<sup>1</sup>Official Methods of Analysis, 16th ed., AOAC, Arlington, VA, Vol. II, Sec. 45.4.07, Method 985.29, 1105 (1997).

### Total Dietary Fiber Assay Procedure

Heat stable  $\alpha$ -Amylase, incubation at pH 6.0, 15 min., 95 °C



Protease incubation at pH 7.5, 30 min., 60 °C



Amyloglucosidase incubation at pH 4.5, 30 min., 60 °C



Ethanol precipitation of Soluble Dietary Fiber



Alcohol and acetone washes



Drying



Kjeldahl Protein  
Determination

Ash Determination  
5 hours, 525 °C



Calculation of Total Dietary Fiber

### Dietary Fiber, Total, Assay Control Kit, Cat. No. TDFC10-1KT

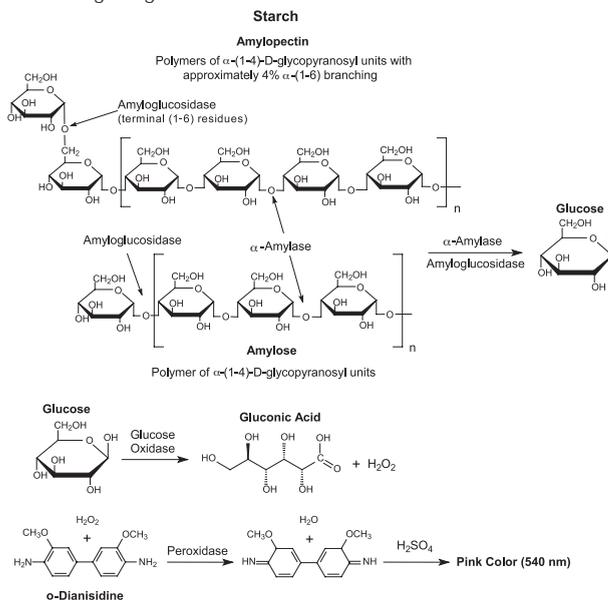
sufficient for ~10 assays

Set of 6 standards for use as internal controls in conjunction with the Total Dietary Fiber Assay Kit (TDF100A)

### Starch (GO/P) Assay Kit, Cat. No. STA20-1KT

sufficient for 20 assays

For the quantitative, enzymatic determination of starch in food and other materials. The hydrolysis of starch to glucose is catalyzed by  $\alpha$ -amylase and amyloglucosidase. Glucose is oxidized to gluconic acid and hydrogen peroxide by glucose oxidase. Hydrogen peroxide reacts with *o*-dianisidine in the presence of peroxidase to form a colored product. Oxidized *o*-dianisidine reacts with sulfuric acid to form a more stable colored product. The intensity of the pink color measured at 540 nm is proportional to the original glucose concentration.

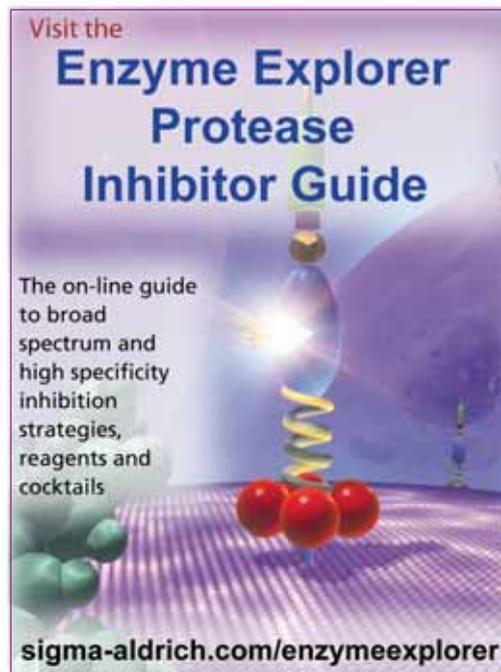


Visit the

## Enzyme Explorer Protease Inhibitor Guide

The on-line guide to broad spectrum and high specificity inhibition strategies, reagents and cocktails

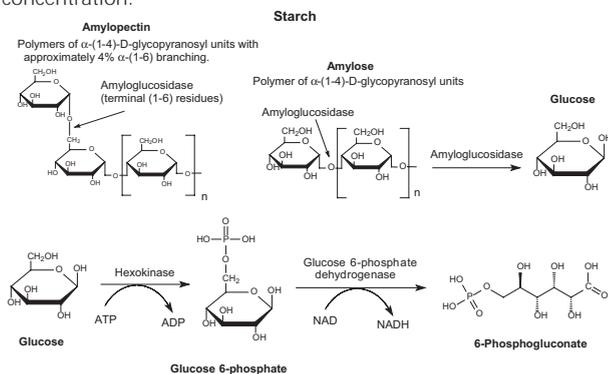
[sigma-aldrich.com/enzymeexplorer](http://sigma-aldrich.com/enzymeexplorer)



**Starch (HK) Assay Kit, Cat. No. SA20-1KT**

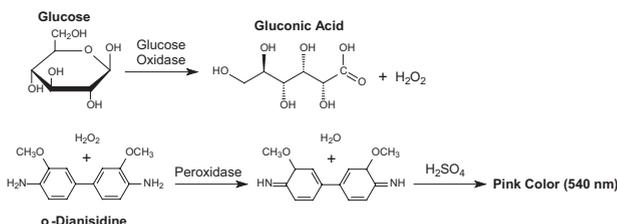
sufficient for 20 assays

For the quantitative, enzymatic determination of native starch in food and other materials. The hydrolysis of starch to glucose is catalyzed by amyloglucosidase. Glucose is phosphorylated by hexokinase. Glucose-6-phosphate is then oxidized to 6-phosphogluconate in the presence of NAD in a reaction catalyzed by glucose 6-phosphate dehydrogenase. The increase in absorbance at 340 nm is directly proportional to the glucose concentration.

**Glucose (GO) Assay Kit, Cat. No. GAGO20-1KT**

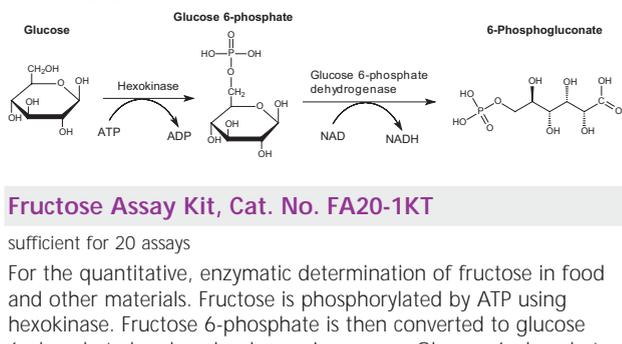
sufficient for 20 assays

For the quantitative, enzymatic determination of glucose in food and other materials. Glucose is oxidized to gluconic acid and hydrogen peroxide by glucose oxidase. Hydrogen peroxide reacts with *o*-dianisidine in the presence of peroxidase to form a colored product. Oxidized *o*-dianisidine reacts with sulfuric acid to form a more stable colored product. The intensity of the pink color measured at 540 nm is proportional to the original glucose concentration.

**Glucose (HK) Assay Kit, Cat. No. GAHK20-1KT**

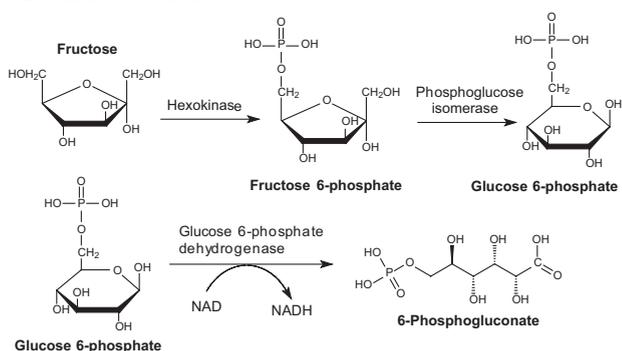
sufficient for 20 assays

For the quantitative, enzymatic determination of glucose in food and other materials. Glucose is phosphorylated by hexokinase to form glucose 6-phosphate. Glucose 6-phosphate is then oxidized to 6-phosphogluconate in the presence of NAD by glucose 6-phosphate dehydrogenase. During this oxidation, an equimolar amount of NAD is reduced to NADH. The consequent increase in absorbance at 340 nm is directly proportional to glucose concentration.

**Fructose Assay Kit, Cat. No. FA20-1KT**

sufficient for 20 assays

For the quantitative, enzymatic determination of fructose in food and other materials. Fructose is phosphorylated by ATP using hexokinase. Fructose 6-phosphate is then converted to glucose 6-phosphate by phosphoglucose isomerase. Glucose 6-phosphate is then oxidized to 6-phosphogluconate in the presence of NAD by glucose 6-phosphate dehydrogenase. During this oxidation, an equimolar amount of NAD is reduced to NADH. The consequent increase in absorbance at 340 nm is directly proportional to fructose concentration.



## Complex Carbohydrates

The following table contains selected polysaccharides, glycosaminoglycans, and related reagents for use as standards, glycan detection and measurement of enzymatic activity.

Product Name	Description	Cat. No.
Alginate ammonium calcium salt <i>Macrocystis pyrifera</i> (kelp)	-	A7253-100G
Alginate from brown algae	-	A7003-100G
		A7003-250G
		A7003-500G
		A7003-1KG
Alginate sodium from salt brown algae	for immobilization of microorganisms	71238-50G
		71238-250G
		71238-1KG
Alginate sodium from salt brown algae	Low viscosity	A2158-100G
		A2158-250G
		A2158-500G
		A2158-1KG
Alginate sodium from salt brown algae	Medium viscosity	A2033-100G
		A2033-250G
		A2033-500G
		A2033-1KG
Alginate sodium from salt brown algae	Low viscosity, plant cell culture tested, powder	A0682-100G
Amylopectin potato starch	-	A8515-25G
		A8515-100G
Amylopectin Azure	Amylase Substrate	A4640-1G
		A4640-5G
		A4640-25G
		A4640-50G
Amylose from potato	Essentially Free of Amylopectin	A0512-250MG
		A0512-1G
		A0512-5G
		A0512-25G
Amylose-Remazol Brilliant Blue R	Amylase Substrates	A3508-1G
		A3508-5G
Chitin from crab shells	suitable for analysis of chitinase, purified powder	C9752-250MG
		C9752-1G
		C9752-5G
Chitin from crab shells	practical grade, powder	C7170-100G
		C7170-1KG
Chitin from crab shells	practical grade, coarse flakes	C9213-500G
Chitin azure	Chitinase Substrate	C9213-1KG
		C3020-100MG
Chitosan from crab shells	≥75% deacetylated	C3020-1G
		C3646-10G
		C3646-25G
		C3646-100G
Chondroitin disaccharide Δdi-0S sodium salt	-	C3646-500G
		C3920-5MG
Chondroitin disaccharide Δdi-4S sodium salt	-	C3920-10MG
		C4045-5MG
Chondroitin disaccharide Δdi-6S sodium salt	-	C4045-10MG
		C4170-5MG
Chondroitin disaccharide Δdi-UA-2S sodium salt	-	C4170-25MG
		C5820-1MG
Chondroitin 6-sulfate sodium salt from shark cartilage	~90%, balance is chondroitin sulfate A	C4384-250MG
		C4384-1G
		C4384-5G
		C4384-25G
Chondroitin sulfate B sodium salt	from porcine intestinal mucosa, ≥90%, lyophilized powder	C3788-25MG
		C3788-100MG
Monoclonal Anti-Chondroitin Sulfate antibody produced in mouse	clone CS-56, ascites fluid	C8035-.2ML
		C8035-.5ML
Chondroitin sulfate A sodium salt from bovine trachea	cell culture tested	C9819-5G
		C9819-25G

## Complex Carbohydrates

Product Name	Description	Cat. No.
Curdlan from <i>Alcaligenes faecalis</i>	-	C7821-5G
Dammar Resin	-	30424-250G 30424-1KG
Dextran from <i>Leuconostoc mesenteroides</i>	average mol wt 9,000-11,000	D9260-10G D9260-50G D9260-100G D9260-500G
Dextran from <i>Leuconostoc mesenteroides</i>	average mol wt 64,000-76,000	D4751-10G D4751-50G D4751-100G D4751-500G D4751-1KG
Dextran from <i>Leuconostoc mesenteroides</i>	average mol wt 35,000-45,000	D1662-10G D1662-50G D1662-100G D1662-500G
Dextran from <i>Leuconostoc mesenteroides</i>	average mol wt 425,000-575,000	D1037-50G D1037-100G D1037-500G
Dextran <i>Leuconostoc mesenteroides</i>	average mol wt 100,000-200,000	D4876-50G D4876-100G D4876-500G D4876-1KG
Dextran from <i>Leuconostoc mesenteroides</i>	industrial grade, average mol wt 5,000,000-40,000,000	D5501-100G D5501-500G D5501-1KG
Dextran from <i>Leuconostoc mesenteroides</i>	average mol wt ~2,000,000	D5376-100G D5376-500G
Dextran solution from <i>Leuconostoc mesenteroides</i>	20 % (w/w) (Autoclaved)	D8802-25ML D8802-50ML
$\beta$ -D-Glucan from barley	powder	G6513-50MG G6513-100MG G6513-500MG G6513-1G G6513-5G
Glucan from baker's yeast ( <i>S. cerevisiae</i> )	-	G5011-25MG G5011-100MG
Glycogen from bovine liver	-	G0885-1G G0885-5G G0885-10G G0885-25G
Glycogen <i>Crepidula fornicata</i> (slipper limpet)	-	G1633-5G
Glycogen from <i>Mytilus edulis</i> (Blue mussel)	-	G1508-5G G1508-25G
Glycogen from oyster	-	G1765-5MG G1765-10MG G1765-25MG
Glycogen from oyster	-	G8751-5G G8751-25G G8751-100G
Glycogen from rabbit liver	-	G8876-500MG G8876-1G G8876-5G G8876-10G
Glycogen azure	from rabbit liver, suitable for substrate for $\alpha$ -amylase	G5510-1G
Glycol chitosan	$\geq 60\%$ (colloidal titration), crystalline	G7753-500MG G7753-1G G7753-5G

## Complex Carbohydrates

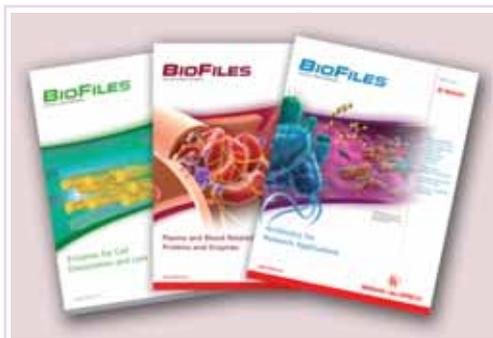
Product Name	Description	Cat. No.
Guar	-	G4129-250G
		G4129-500G
		G4129-1KG
Gum accroides	-	G9627-500G
Gum arabic from acacia tree	reagent grade	G9752-500G
		G9752-1KG
Heparan sulfate sodium salt from bovine kidney	-	H7640-1MG
		H7640-5MG
		H7640-10MG
Heparan sulfate fast-moving fraction sodium salt from porcine intestinal mucosa	≥90% (electrophoresis)	H9902-1MG
		H9902-5MG
Heparan sulfate proteoglycan	≥400 µg/mL glycosaminoglycan, sterile-filtered	H4777-.1MG
Heparin sodium salt from porcine intestinal mucosa	mol wt 4,000-6,000 Da	H8537-500MG
		H8537-100MG
		H8537-250MG
		H8537-1G
Heparin sodium salt from porcine intestinal mucosa	average mol wt ~3,000	H3400-50MG
		H3400-100MG
		H3400-250MG
		H3400-1G
Heparin sodium salt from porcine intestinal mucosa	Grade I-A, activity: ~170 USP units/mg	H3393-10KU
		H3393-25KU
		H3393-50KU
		H3393-100KU
		H3393-250KU
		H3393-500KU
Heparin-benzalkonium	activity: ~60 USP units/mg	H7280-1G
Heparin, deaminated sodium salt from porcine intestinal mucosa	Low molecular weight mono-aldehyde, heparin activity: >75 USP units/mg	H7405-250MG
		H7405-1G
Heparin disaccharide I-A sodium salt	$\alpha$ - $\delta$ UA-2S-[1→4]-GlcNAc-6S	H9517-1MG
Heparin disaccharide I-H sodium salt	$\alpha$ - $\delta$ UA-2S-[1→4]-Glc-6S	H8892-1MG
Heparin disaccharide I-S sodium salt	( $\alpha$ - $\Delta$ UA-2S-[1→4]-GlcNS-6S)	H9267-1MG
Heparin disaccharide II-H sodium salt	( $\alpha$ - $\delta$ UA-[1→4]-GlcN-6S)	H9017-1MG
Heparin disaccharide III-H sodium salt	( $\alpha$ - $\delta$ UA-S2-[1→4]-GlcN)	H9142-1MG
Heparin disaccharide III-S sodium salt	( $\alpha$ - $\Delta$ UA-2S-[1→4]-GlcNS)	H9392-1MG
Heparin disaccharide IV-A sodium salt	( $\alpha$ - $\Delta$ UA-[1→4]-GlcNAc)	H0895-.5MG
Heparin disaccharide IV-H ≥95%	$\alpha$ - $\Delta$ UA-[1→4]-GlcN	H9276-1MG
Hyaluronan biotin sodium salt	>97%, soluble powder	B1557-5MG
Hyaluronic acid potassium salt from human umbilical cord	suitable as substrate for hyaluronidase	H1504-50MG
		H1504-100MG
		H1504-500MG
		H1504-1G
Hyaluronic acid potassium salt from human umbilical cord	Highly polymerized	H1751-500MG
Hyaluronic acid sodium salt from bovine vitreous humor	-	H7630-10MG
		H7630-50MG
Hyaluronic acid sodium salt from rooster comb	-	H5388-100MG
		H5388-250MG
		H5388-1G
Hyaluronic acid sodium salt from <i>Streptococcus equi</i>	-	53747-1G
		53747-10G
Hyaluronic acid sodium salt from <i>Streptococcus zooepidemicus</i>	-	H9390-1G
Hyaluronic acid disaccharide $\delta$ DIHA sodium salt	≥95%	H9649-1MG
Inulin from chicory	-	I2255-10G
		I2255-25G
		I2255-100G
		I2255-1KG

## Complex Carbohydrates

Product Name	Description	Cat. No.
Inulin from dahlia tubers	Mr <sup>2</sup> 5000	I3754-25G
		I3754-100G
		I3754-1KG
Inulin-FITC	from dahlia tuber	F3272-1G
Lichenan from <i>Cetraria islandica</i>	practical grade, powder	L6133-250MG L6133-1G
Pectin from apple	meets USP testing specifications	P8471-100G
		P8471-500G
Pectin from citrus peel	Galacturonic acid: $\geq 74.0\%$	P9135-100G
		P9135-500G
		P9135-1KG
Pectin, esterified from citrus fruit	extent of labeling: ~90% esterified	P9561-5G
		P9561-25G
Pectin, esterified potassium salt from citrus fruit	extent of labeling: ~60% esterified	P9436-5G
		P9436-25G
		P9436-50G
Pectin, esterified potassium salt from citrus fruit	extent of labeling: ~30% esterified	P9311-5G
		P9311-25G
Peptidoglycan from <i>Bacillus subtilis</i>	-	69554-10MG-F
Peptidoglycan from <i>Micrococcus luteus</i>	-	53243-10MG-F
Peptidoglycan from <i>Saccharomyces cerevisiae</i>	-	72789-10MG-F
Peptidoglycan from <i>Staphylococcus aureus</i>	-	77140-10MG
		77140-25MG
Peptidoglycan from <i>Streptomyces</i> sp.	-	79682-10MG-F
Pullulan from <i>Aureobasidium pullulans</i>	suitable for substrate for pullulanase	P4516-1G
		P4516-5G
		P4516-25G
Stachyose hydrate from <i>Stachys tubrifera</i>	$\geq 98\%$	S4001-10MG
		S4001-100MG
		S4001-500MG
		S4001-1G
		S4001-5G
Starch	from potatoes, for electrophoresis	85645-100G
		85645-1KG
Starch	from potatoes	85650-1KG
Starch from corn	Standard for Starch Assay Kits SA-20 and STA-20	S5296-5G
Starch from corn	practical grade	S4180-100G
		S4180-500G
		S4180-1KG
Starch from corn	Unmodified regular corn starch containing approx. 73% amylopectin and 27% amylose	S4126-2KG
		S4126-5KG
Starch from potato	for electrophoresis	S5651-100G
		S5651-500G
		S5651-1KG
		S5651-2KG
		S5651-5KG
Starch from potato	Soluble	S2630-100G
		S2630-500G
		S2630-1KG
Starch from potato	Soluble	S2004-500G
		S2004-1KG
Starch from potato	Powder	S4251-2KG
		S4251-5KG
Starch from rice	-	S7260-100G
		S7260-500G
		S7260-1KG
Starch from wheat	Unmodified S	5127-100G
		S5127-500G S5127-5KG

## Complex Carbohydrates

Product Name	Description	Cat. No.
Starch from wheat	Purified	S2760-500G
Starch from Azure	Potato starch covalently linked with Remazol Brilliant Blue	S7629-1G S7629-5G S7629-25G
Starch from Azure	Insoluble corn starch covalently linked with Remazol Brilliant Blue	S7776-1G S7776-5G
Starch, from soluble	ACS reagent	S9765-100G S9765-250G S9765-500G S9765-1KG
Xylan from beechwood	-	X4252-10G X4252-25G X4252-100G
Xylan from birch wood	Xylose residues: $\geq 90\%$	X0502-10G X0502-25G X0502-100G
Xylan from birch wood	-	95588-10G 95588-25G 95588-100G
Xylan from oat spelts	Xylose: $\geq 70\%$	X0627-10G X0627-25G X0627-100G
Xylan from oat spelts	-	95590-10G 95590-50G



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