

Product Information

Lectin

from *Agaricus bisporus* (mushroom)

Catalog Number **L5640**

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

Product Description

Agaricus bisporus agglutinin (ABA) is a mixture of two phytohemagglutinins with similar specificities for carbohydrate receptors (PHA-A and PHA-B).¹ Both PHA-A and PHA-B are heterogeneous by SDS-PAGE. The mushroom PHA will have a molecular weight of 58.5 kDa as determined by gel filtration.

PHA-A and PHA-B agglutinate erythrocytes independent of their blood group type. Trypsin treatment of erythrocytes decreases lectin binding to the cell surface. The O-linked glycopeptide released by trypsin is a potent inhibitor of both the isolectins. Removal of the terminal sialic acid residue from the glycopeptide increases its inhibitory potency 8-fold. Periodate or β -galactosidase treatment of the trypsin released glycopeptide destroys all inhibitory activity. Simple sugars are very poor inhibitors of these isolectins. However, a galactose residue appears to play a major role in these lectin binding mechanisms. The sugar linkage is also important since a synthetic N-linked glycopeptide is not inhibitory, while the O-linked glycopeptide released by trypsinization is a potent inhibitor. The cell receptor for the isolectins has the structure Galactose-Gal-NAc-ser (or threonine).²

Lectins are proteins or glycoproteins of non-immune origin that agglutinate cells and/or precipitate complex carbohydrates. Lectins are capable of binding glycoproteins even in presence of various detergents.³ The agglutination activity of these highly specific carbohydrate-binding molecules is usually inhibited by a simple monosaccharide, but for some lectins, di, tri, and even polysaccharides are required.

Lectins are isolated from a wide variety of natural sources, including seeds, plant roots and bark, fungi, bacteria, seaweed and sponges, mollusks, fish eggs, body fluids of invertebrates and lower vertebrates, and from mammalian cell membranes.

The precise physiological role of lectins in nature is still unknown, but they have proved to be very valuable in a wide variety of applications *in vitro*, including:

- blood grouping and erythrocyte polyagglutination studies
- fractionation of cells and other particles
- histochemical studies of normal and pathological conditions.
- mitogenic stimulation of lymphocytes.
- lymphocyte subpopulation studies.

Sigma offers a range of lectins suitable for the above applications; most are highly purified by affinity chromatography, but some are offered as purified or partially purified lectins, suitable for specific applications.

Conjugation does not alter the specificity of the lectin. Many of the lectins are available conjugated to:

- fluorochromes (for detection by fluorimetry).
- enzymes (for enzyme-linked assays).
- insoluble matrices (for use as affinity media).

Please refer to the table for general information on the most common lectins.

Precautions and Disclaimer

This product is for R&D use only, not for drug, household, or other uses. Please consult the Material Safety Data Sheet for information regarding hazards and safe handling practices.

Reagent

Lyophilized powder containing ~10% protein (Biuret).

Preparation Instructions

Soluble in phosphate buffered saline, pH 6.8 (1 mg/ml), yielding a clear to hazy, faint yellow solution.

Storage/Stability

Aggregation is thought to occur in the presence of high concentrations of 2-mercaptoethanol.

Lectin	MW (kDa)	Subunits	Specificity		Mitogenic Activity
			Blood Group	Sugar	
<i>Abrus precatorius</i>			–		+
Agglutinin	134	4		gal	
Abrin A (toxin)	60	2		gal	
Abrin B (toxin)	63.8	2($\alpha\beta$)		gal	
<i>Agarius bisporus</i>	58.5	–	–	β -gal(1 \rightarrow 3)galNAc	
<i>Anguilla anguilla</i>	40	2	H	α -L-Fuc	
<i>Arachis hypogaea</i>	120	4	T	β -gal(1 \rightarrow 3)galNAc	
<i>Artocarpus integrifolia</i>	42	4	T	α -gal \rightarrow OME	+
<i>Bandeiraea simplicifolia</i>					
BS-I	114	4	A, B	α -gal, α -galNAc	
BS-I-A ₄	114	4	A	α -galNAc	
BS-I-B ₄	114	4	B	α -gal	
BS-II	113	4	acq, B, Tk, T	glcNAc	
<i>Bauhinia purpurea</i>	195	4	–	β -gal(1 \rightarrow 3)galNAc	+
<i>Caragana arborescens</i>	60; 120 ^a	2/4	–	galNAc	
<i>Cicer arietinum</i>	44	2	–	fetuin	
<i>Codium fragile</i>	60	4	–	galNAc	
<i>Concanavalin A</i>	102	4	–	α -man, α -glc	+
<i>Succinyl-Concanavalin A</i>	51	2	–	α -man, α -glc	+ ^b
<i>Cytisus scoparius</i>	–	–	–	galNAc, gal	
<i>Datura stramonium</i>	86	2($\alpha\beta$)	–	(glcNAc) ₂	
<i>Dolichos biflorus</i>	140	4	A ₁	α -galNAc	
<i>Erythrina corallodendron</i>	60	2	–	β -gal(1 \rightarrow 4)glcNAc	+
<i>Erythrina cristagalli</i>	56.8	2($\alpha\beta$)	–	β -gal(1 \rightarrow 4)glcNAc	
<i>Euonymus europaeus</i>	166	4($\alpha\beta$)	B, H	α -gal(1 \rightarrow 3)gal	+
<i>Galanthus nivalis</i>	52	4	(h)	non-reduc. α -man	
<i>Glycine max</i>	110	4	–	galNAc	+ ^c
<i>Helix aspersa</i>	79	–	A	galNAc	
<i>Helix pomatia</i>	79	6	A	galNAc	
<i>Lathyrus odoratus</i>	40-43	4($\alpha\beta$)	–	α -man	+
<i>Lens culinaris</i>	49	2	–	α -man	+
<i>Limulus polyphemus</i>	400	18	–	NeuNAc	
Bacterial agglutinin	–	–	–	galNAc, glcNAc	
<i>Lycopersicon esculentum</i>	71	–	–	(glcNAc) ₃	
<i>Maackia amurensis</i>	130	2($\alpha\beta$)	O	sialic acid	+
<i>Maclura pomifera</i>	40-43	2($\alpha\beta$)	–	α -gal, α -galNAc	
<i>Momordica charantia</i>	115-129	4($\alpha\beta$)	–	gal, galNAc	
<i>Naja mocambique mocambique</i>	–	–	–	–	
<i>Naja naja kaouthia</i>	–	–	–	–	
<i>Narcissus pseudonarcissus</i>	26	2	(h)	α -D-man	
<i>Perseu americana</i>	–	–	–	–	
<i>Phaseolus coccineus</i>	112	4	–	–	
<i>Phaseolus limensis</i>	247(II)	8	A	galNAc	+
	124(III)	4			
<i>Phaseolus vulgaris</i>					
PHA-E	128	4	–	oligosaccharide	+
PHA-L	128	4	–	oligosaccharide	+
PHA-P					
PHA-M					

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Lectin	MW (kDa)	Subunits	Specificity		Mitogenic Activity
			Blood Group	Sugar	
<i>Phytolacca americana</i>	32	–	–	(glcNAc) ₃	+
<i>Pisum sativum</i>	49	4(αβ)	–	α-man	+
<i>Pseudomonas aeruginosa PA-I</i>	13-13.7	–	–	gal	+ ^c
<i>Psophocarpus tetragonolobus</i>	35	1	–	galNAc, gal	
<i>Ptilota plumosa</i>	65; 170	–	B	α-gal	
<i>Ricinus communis</i>					
Toxin, RCA ₆₀	60	2	–	galNAc, β-gal	
Toxin, RCA ₁₂₀	120	4	–	β-gal	
<i>Sambucus nigra</i>	140	4(αβ)	–	αNeuNAC(2→6)gal galNAc	+ ^c
<i>Solanum tuberosum</i>	50; 100 ^a	1, 2	–	(glcNAc) ₃	
<i>Sophora japonica</i>	133	4	A, B	β-galNAc	
<i>Tetragonolobus purpureas</i>	120(A)	4	H	α-L-fuc	
	58(BA)	2	H	α-L-fuc	
	117(C)	4	H	α-L-fuc	
<i>Triticum vulgare</i>	36	2	–	(glcNAc) ₂ , NeuNAc	+
<i>Ulex europaeus</i>					
UEA I	68	–	H	α-L-fuc	
UEA II	68	–	–	(glcNAc) ₂	
<i>Vicia faba</i>	50	4(αβ)	–	man, glc	+
<i>Vicia sativa</i>	40	4(αβ)	–	glc, man	+
<i>Vicia villosa</i>	139	4	A ₁ +T _n	galNAc	
A ₄	134	4	A ₁	galNAc	
B ₄	143	4	T _n	galNAc	
<i>Vigna radiata</i>	160	4	–	α-gal	
<i>Viscum album</i>	115	4(αβ)	–	β-gal	
<i>Wisteria floribunda</i>	68	2	–	galNAc	

^a Concentration-dependent molecular weight

^b Non-agglutinating and mitogenic

^c Mitogenic for neuraminidase-treated lymphocytes

References

1. Presant, C.A. and Kornfeld, S., *J. Biol. Chem.*, **247**, 6937 (1972).
2. Young, N. M., et al., *J. Biol. Chem.*, **246**, 1596 (1971).
3. Rueben, L., et al., Activities of lectins and their immobilized derivatives in detergent solutions. Implications on the use of lectin affinity chromatography for the purification of membrane glycoproteins. *Biochemistry*, **16**, 1787-1794 (1977).

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