PRODUCT INFORMATION **TRITC Labeled Lectin**

Catalog Number:	R-3901-1		
Description:	Pure Maclura pomifera lectin (MPA) fro	om Osage Ora	nge, TRITC conjugated.
Lot Number:			
Protein Concentration: (Based on OD 280)	1 mg purified MPA TRITC / 1 ml Buffer	r.	
TRITC / Protein Ratio: (OD 550 / OD 280)			
Purification Procedure:	Gel filtration performed after conjugatio	on to remove fr	ree TRITC.
Carbohydrate Specificity:	N-Acetylgalactosamine>Galactose.		
Inhibitory Carbohydrate:	Melibiose [Gal α(1,6) Glc]>α-D-Galacto	ose.	
Activity:	Less than 5 μ g/ml will agglutinate ty 0.1 μ g/ml will agglutinate neuraminidase		erythrocytes. Less than
Buffer:	0.02M Sodium Bicarbonate, pH 9.0-9. preservative.	.5. Contains	0.05% sodium azide as a
Chemical Used for Conjugation:	Tetramethylrhodamine Isothiocyanate, TR	RITC.	
Storage:	Store liquid material frozen in aliquots in freeze thaw cycles. Clarify by centrifugati		r covered with foil. Avoid
Stability:	The liquid material is stable for at least with 0.05% sodium azide added as a pres		stored frozen in aliquots
Caution:	Refer to the enclosed MSDS for inform seals have sharp edges and the vial its lacerations. Use caution when opening t	self may have	
Remarks:	Fluorescent Conjugates are extremely lig	ght sensitive.	
References:	 Bausch, N.J., et.al. (1977) Biochem Jones, J.M., et.al. J.D. (1973) J. Imr Bird, G.W.G., et.al. (1973) Vox San 	nunol. 111 :17	55.
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	DRIES, INC.	Tel:	650-342-3296
107 North Amphlett E San Mateo, CA 9440	3lvd. 1	Fax: Orders:	650-342-2648 1-800-821-0044

General Procedure Fluorescent Labeled Lectin

The following is a general Procedure and Trouble-Shooting Guide. The information is provided only for your convenience. The success of your experiments are not guaranteed by EY Laboratories, Inc.

Tissue Sections

1.					
	Wash and block tissue section. Do not use serum products, they contain glycoproteins which may lead to high levels of non specific background. After blocking, rinse briefly with Buffer (See reverse side).				
2.	Dilute Fluorescent Labeled Lectin to desired concentration 20-100 µg/ml using Buffer.				
3.	Incubate tissue section with Fluorescent Labeled Lectin for 30 minutes in a moist chamber.				
4.	Wash tissue section with Buffer three times.				
5.	Examine tissue section with Fluorescent microscope. Use appropriate filter.				
	Ref. M. Imm	bar et. al., (1973). Intnl. Journal of Cancer, 12	, 93-99		
		Cell Suspension	n		
1.	Wash cells with Buffer (See reverse side.)				
2.	Collect cells by centrifugation.				
3.	Dilute Fluorescent Labeled Lectin to 100 µg/ml using Buffer.				
4.					
5.	Wash cells with Buffer three times using centrifugation.				
6.	Examine cells	s, with or without fixation with Fluorescent mi	croscope. Use appropriate filter.		
	Ref. K. Phiss	. (1977). Experimental Pathology, 14, S15			
	rochromes mu red in foil.	ust be protected from light. Perform incub	ation, when practical, in a dark room or		
		Absorption and Emi	ission		
		Absorption/Excitation Ra	te Emission Max.		
	FIT		517 nm		
		TC 554 nm	570 nm		
	Tex	as Red [™] 596 nm	615 nm		
		Carbohydrate Inhib	bition		
In hall					
innibi	ition of lectin	binding may be accomplished by using one of	two procedures:		
A. B.	Before incut carbohydrate Preincubate	pating with Fluorescent Labeled Lectin , for 30-60 minutes at room temperature. NOT	incubate section or cells with inhibitory		
A.	Before incut carbohydrate Preincubate	bating with Fluorescent Labeled Lectin , for 30-60 minutes at room temperature. NOT diluted Fluorescent Labeled Lectin with i ature before applying to section or cells.	incubate section or cells with inhibitory rE: Complete inhibition may NOT occur. nhibitory carbohydrate for 30-60 minutes at		
А. В.	Before incut carbohydrate Preincubate o room tempera	bating with Fluorescent Labeled Lectin, for 30-60 minutes at room temperature. NOT diluted Fluorescent Labeled Lectin with i ature before applying to section or cells. TROUBLE SHOOTING	incubate section or cells with inhibitory TE: Complete inhibition may NOT occur. nhibitory carbohydrate for 30-60 minutes at GUIDE		
А. В.	Before incut carbohydrate Preincubate	bating with Fluorescent Labeled Lectin, for 30-60 minutes at room temperature. NOI diluted Fluorescent Labeled Lectin with i ature before applying to section or cells. TROUBLE SHOOTING Cause	incubate section or cells with inhibitory TE: Complete inhibition may NOT occur. nhibitory carbohydrate for 30-60 minutes at GUIDE		
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А. В. Р W	Before incub carbohydrate Preincubate o room tempera	bating with Fluorescent Labeled Lectin, for 30-60 minutes at room temperature. NOT diluted Fluorescent Labeled Lectin with i ature before applying to section or cells. TROUBLE SHOOTING Cause 1. Low concentration of specific	incubate section or cells with inhibitory TE: Complete inhibition may NOT occur. nhibitory carbohydrate for 30-60 minutes at GUIDE Solution Causes #1 - #3		
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А. В. Р W	Before incub carbohydrate Preincubate o room tempera Problem	hating with Fluorescent Labeled Lectin, for 30-60 minutes at room temperature. NOT diluted Fluorescent Labeled Lectin with i ature before applying to section or cells. TROUBLE SHOOTING Cause 1. Low concentration of specific oligosaccharide on sample. 2. Low concentration of lectin conjugate. 3. Insufficient incubation time.	incubate section or cells with inhibitory TE: Complete inhibition may NOT occur. nhibitory carbohydrate for 30-60 minutes at GUIDE Causes #1 - #3 a. Increase incubation time. b. Increase concentration conjugate. a. Avoid exposure to light. a. Decrease concentration of Lectin conjugate.		
А. В. Р W	Before incub carbohydrate Preincubate o room tempera Problem	bating with Fluorescent Labeled Lectin, for 30-60 minutes at room temperature. NOT diluted Fluorescent Labeled Lectin with in ature before applying to section or cells. TROUBLE SHOOTING Cause 1. Low concentration of specific oligosaccharide on sample. 2. Low concentration of lectin conjugate. 3. Insufficient incubation time. 4. Photobleaching 1. Lectin conjugate is too concentrated.	incubate section or cells with inhibitory TE: Complete inhibition may NOT occur. nhibitory carbohydrate for 30-60 minutes at GUIDE Causes #1 - #3 a. Increase incubation time. b. Increase concentration conjugate. a. Avoid exposure to light. a. Decrease concentration of Lectin conjugate. b. Shorten incubation times.		
А. В. Р W	Before incut carbohydrate Preincubate e room tempera roblem Veak or no Staining	bating with Fluorescent Labeled Lectin, for 30-60 minutes at room temperature. NOT diluted Fluorescent Labeled Lectin with i ature before applying to section or cells. TROUBLE SHOOTING Cause 1. Low concentration of specific oligosaccharide on sample. 2. Low concentration of lectin conjugate. 3. Insufficient incubation time. 4. Photobleaching	incubate section or cells with inhibitory E: Complete inhibition may NOT occur. nhibitory carbohydrate for 30-60 minutes at GUIDE Solution Causes #1 - #3 a. Increase incubation time. b. Increase concentration conjugate. a. Avoid exposure to light. a. Decrease concentration of Lectin conjugate. b. Shorten incubation times. a. Perform multiple washings and prolong		
A. B. W	Before incul carbohydrate Preincubate e room tempera Problem Veak or no Staining High	bating with Fluorescent Labeled Lectin, for 30-60 minutes at room temperature. NOT diluted Fluorescent Labeled Lectin with i ature before applying to section or cells. TROUBLE SHOOTING • Cause 1. Low concentration of specific oligosaccharide on sample. 2. Low concentration of lectin conjugate. 3. Insufficient incubation time. 4. Photobleaching 1. Lectin conjugate is too concentrated. 2. Insufficient washing.	incubate section or cells with inhibitory TE: Complete inhibition may NOT occur. nhibitory carbohydrate for 30-60 minutes at GUIDE Solution Causes #1 - #3 a. Increase incubation time. b. Increase concentration conjugate. a. Avoid exposure to light. a. Decrease concentration of Lectin conjugate. b. Shorten incubation times. a. Perform multiple washings and prolong washing time.		
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A. B. W	Before incul carbohydrate Preincubate e room tempera Problem Veak or no Staining High	bating with Fluorescent Labeled Lectin, for 30-60 minutes at room temperature. NOT diluted Fluorescent Labeled Lectin with i ature before applying to section or cells. TROUBLE SHOOTING • Cause 1. Low concentration of specific oligosaccharide on sample. 2. Low concentration of lectin conjugate. 3. Insufficient incubation time. 4. Photobleaching 1. Lectin conjugate is too concentrated. 2. Insufficient washing.	incubate section or cells with inhibitory TE: Complete inhibition may NOT occur. nhibitory carbohydrate for 30-60 minutes at GUIDE Solution Causes #1 - #3 a. Increase incubation time. b. Increase concentration conjugate. a. Avoid exposure to light. a. Decrease concentration of Lectin conjugate. b. Shorten incubation times. a. Perform multiple washings and prolong washing time. a. Use fluorochrome with different excitation and emission spectrum. b. Use a different lectin conjugate (enzyme or colloidal gold).		
A. B. W	Before incul carbohydrate Preincubate e room tempera roblem /eak or no Staining High ackground	 bating with Fluorescent Labeled Lectin, for 30-60 minutes at room temperature. NOI diluted Fluorescent Labeled Lectin with i ature before applying to section or cells. TROUBLE SHOOTING Cause Low concentration of specific oligosaccharide on sample. Low concentration of specific oligosaccharide on sample. Low concentration of lectin conjugate. Insufficient incubation time. Photobleaching Lectin conjugate is too concentrated. Insufficient washing. Autofluorescent sample. 	incubate section or cells with inhibitory TE: Complete inhibition may NOT occur. nhibitory carbohydrate for 30-60 minutes at GUIDE Solution Causes #1 - #3 a. Increase incubation time. b. Increase concentration conjugate. a. Avoid exposure to light. a. Decrease concentration of Lectin conjugate. b. Shorten incubation times. a. Perform multiple washings and prolong washing time. a. Use fluorochrome with different excitation and emission spectrum. b. Use a different lectin conjugate (enzyme <u>or</u> colloidal gold). a. Perform control reactions.		
A. B. W	Before incul carbohydrate Preincubate e room tempera Problem Veak or no Staining High	bating with Fluorescent Labeled Lectin, for 30-60 minutes at room temperature. NOT diluted Fluorescent Labeled Lectin with i ature before applying to section or cells. TROUBLE SHOOTING • Cause 1. Low concentration of specific oligosaccharide on sample. 2. Low concentration of lectin conjugate. 3. Insufficient incubation time. 4. Photobleaching 1. Lectin conjugate is too concentrated. 2. Insufficient washing.	incubate section or cells with inhibitory TE: Complete inhibition may NOT occur. nhibitory carbohydrate for 30-60 minutes at GUIDE Solution Causes #1 - #3 a. Increase incubation time. b. Increase concentration conjugate. a. Avoid exposure to light. a. Decrease concentration of Lectin conjugate. b. Shorten incubation times. a. Perform multiple washings and prolong washing time. a. Use fluorochrome with different excitation and emission spectrum. b. Use a different lectin conjugate (enzyme or colloidal gold).		



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MSDS for Fluorescent labeled Purified Proteins Continued - page 2 of 2.

MATERIAL SAFETY DATA SHEET

Effective Date: March 31, 2006 Revision 4 Page 1 of 2

PRODUCT IDENTIFICATION

Purified proteins labeled with fluorescein isothiocyanate (FITC),
tetramethylrhodamine isothiocyanate (TRITC), or Texas Red a trademark of
Molecular Probes for the sulfonyl chloride derivative of sulforhodamine 101
FP-01, RP-01, TP-01, F-1102 to F-9000, R-1102 to R-9000, T-1102 to T-9000, FA-
2100 to FA-2701, RA-2100 to RA-2701, TA-2100 to TA-2701, FAF-001 to FAF-
2354, RAF-001 to RAF-2354, TAF-001 to TAF-2354, FAL-1104 to FAL-4701,
RAL-1104 to RAL-4701, TAL-1104 to TAL-4701, FA-01 to FA-013, TA-01 to
TA-013, DM1011F to DM1064F, FNP-01 to FNP-05, BA-101, BA-102, BA-612.
Protein A, Avidin (egg white), Glycosylated Bovine Serum Albumin, Lectins,
Secondary and Monoclonal Antibodies labeled with FITC, TRITC, or Texas $\operatorname{Red}^{\otimes}$

EMERGENCY INFORMATION

EY Laboratories, Inc. 107 North Amphlett Blvd. San Mateo, CA 94401 EMERGENCY PHONE: 650-342-3296

HAZARDOUS COMPONENTS

Specific protein(s) as listed on the vial label. Solutions are at a concentration generally greater than 0.5mg protein/ml. Biological activity of these labeled proteins will vary. FITC, TRITC, and Texas Red \circledast are possible carcinogens in their pure form. Compounds with similar chemical structures are known to be reactive with proteins and other biomolecules. The complete properties of the dyes after labeling have not been evaluated. These compounds should be treated as potentially hazardous. All solutions contain less than 0.05% sodium azide as a preservative.

HEALTH HAZARD INFORMATION

EXPOSURE LIMITS:	None established. The toxicological properties of these products have not
	been thoroughly investigated. Care should be taken when handling any of
	these materials.
EFFECTS OF	Causes localized eye, skin, or mucous membrane irritation. Some sensitive
OVEREXPOSURE:	individuals may develop a chronic allergic reaction with exposure. The
	known effects are due to the protein. No specific effects of the bound dye are known at this time.
ROUTES OF EXPOSURE:	Inhalation of powders and skin contact with liquids are the primary routes of exposure. Care should be taken to avoid the formation of aerosols when
	handling any of the solutions.

PHYSICAL CHARACTERISTICS

APPEARANCE: SOLUBILITY:

Powders are a light orange. Solutions will be yellow to dark purple. Powders are completely soluble in many biological buffers and water. I liquids are completely miscible in water and biological buffers.

FIRE AND EXPLOSION HAZARDS

Not considered to be a vire hazard. At high concentrations the chemicals may emit toxic fumes. Such high concentrations are not normally found in a research laboratory.

EXTINGUISHING MEDIA: SPECIAL FIRE FIGHTING CRECAUTIONS:

Dry chemical powder or CO₂. Wear self-contained breathing apparatus and protective clothing to prevent contact with skin and eyes.

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NOTE: Most solutions contain less than 0.05% sodium azide as a preservative. Azide may react with lead and copper plumbing to form explosive metal azides. Flush with copious amounts of water when disposing material in the sink.

REACTIVITY DATA

STABILITY: HAZARDOUS POLYMER INCOMPATIBILITY:	IZATION:	Stable. Decomposition products are not known to be hazardous. Will NOT occur. Alcohols, strong bases and acids, strong oxidizing agents, and heat. (Lead and copper may react with sodium azide).
SPILL / LEAK PROCEDU MATERIAL RELEASE / SPILL:	URES Avoid contact with powder or liquid. Clean up spill with a paper towel soaked in household bleach. Do not allow solutions to dry on environmental surfaces. Wash affected area with detergent after the area	
WASTE DISPOSAL:	Incinerate, au	ed with bleach. toclave, or dispose of paper waste in accordance with all and Eederal regulations. Due to the small quantities of

Local, State, and Federal regulations. Due to the small quantities of material involved these products are generally not considered to be environmental hazards. All of these proteins are fully biodegradable.

EMERGENCY FIRST AID PROCEDURES

May be harmful if swallowed, inhaled, or allowed to absorb through the skin. Wash contacted area with water for 15 minutes. If inhaled remove to fresh air. Report exposure to the appropriate safety official. Consult a physician if irritation occurs or if there is any indication of an allergic response, such as watering eyes, sneezing, or difficulty breathing.

SPECIAL HANDLING PRECAUTIONS

VENTILATION:	No special ventilation is required but it is recommended to handle these reagents in a fume hood when possible.
EYE PROTECTION:	Required. Goggles or safety glasses with a side shield are recommended.
RESPIRATORY	Recommended as a safety precaution, specifically when working with
PROTECTION:	powders. An approved respirator may be required for those individuals
	already known to be sensitive to these materials.
PROTECTIVE GLOVES:	Required when handling any of these materials.

SPECIAL PRECAUTIONS

This material is for research and experimental application only. It is not intended for food, drug, household, agricultural, or cosmetic use. All materials should be handled only by technically qualified individuals experienced with working with potentially hazardous chemicals. The above information is correct to the best of our knowledge. The user should make independent decisions regarding completeness of the information, based on all sources available. EY Laboratories, Inc. shall not be held liable for any damage resulting from handling or contact with the above product.



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