

PRODUCT INFORMATION **FITC Labeled Lectin**

Catalog Number:	F-3901-2	The following is a general Procedure and Trouble-Shooting Guide. The information is p your convenience. The success of your experiments are not guaranteed by EY Laboratories, In Tissue Sections			
Description:	Pure Maclura pomifera lectin (MPA) from Osage Orange, FITC conjugated.	 Wash and block tissue section. Do not use serum products, they contain glycoproteins to high levels of non specific background. After blocking, rinse briefly with Buffer (See Dilute Fluorescent Labeled Lectin to desired concentration 20-100 µg/ml using Buffer 	e reverse side).		
Lot Number:		 Incubate tissue section with Fluorescent Labeled Lectin for 30 minutes in a moist chamb Wash tissue section with Buffer three times. 	ber.		
Protein Concentration: (Based on OD 280)	2 mg purified MPA FITC /2 ml Buffer.	 Examine tissue section with Fluorescent microscope. Use appropriate filter. Ref. M. Immbar et. al., (1973). Intnl. Journal of Cancer, 12, 93-99 Cell Suspension 			
FITC / Protein Ratio: (OD 495/ OD 280)		 Wash cells with Buffer (See reverse side.) Collect cells by centrifugation. Dilute Fluorescent Labeled Lectin to 100 µg/ml using Buffer. 			
Purification Procedure	: Gel filtration performed after conjugation to remove free FITC.	 Bindo Thatecent Entret Entret Form of print using Burter. Incubate approximately 1x10⁶ cells with 1 ml diluted Fluorescent labeled Lectin for 15 temperature or in a 37°C water bath. Wash cells with Buffer three times using centrifugation. 	minutes at room		
Carbohydrate Specificity:	N-Acetylgalactosamine>Galactose.	 Examine cells, with or without fixation with Fluorescent microscope. Use appropriate fi Ref. K. Phiss. (1977). Experimental Pathology, 14, S15 			
Inhibitory Carbohydrate:	Melibiose [Gal α (1,6) Glc]> α -D-Galactose.	Fluorochromes must be protected from light. Perform incubation, when practical, in covered in foil.	a dark room or		
Activity:	Less than 5 μ g/ml will agglutinate type O human erythrocytes. Less than 0.1 μ g/ml will agglutinate neuraminidase treated cells.	Absorption and EmissionAbsorption/Excitation RateEmission Max.FITC492 nm517 nmTRITC554 nm570 nm			
Buffer:	0.02 M Sodium Bicarbonate, pH 9.0-9.5. Contains 0.05% sodium azide as a preservative.	Texas Red™ 596 nm 615 nm Carbohydrate Inhibition			
Chemical Used for Conjugation: Storage:	Fluorescein Isothiocyanate, FITC. Store liquid material frozen in aliquots in amber vials or covered with foil. Avoid	 carbohydrate for 30-60 minutes at room temperature. NOTE: Complete inhibition may 1 B. Preincubate diluted Fluorescent Labeled Lectin with inhibitory carbohydrate for 3 	A. Before incubating with Fluorescent Labeled Lectin , incubate section or cells with inhibitory carbohydrate for 30-60 minutes at room temperature. NOTE: Complete inhibition may NOT occur.		
Ota h Ultra	freeze thaw cycles. Clarify by centrifugation.	TROUBLE SHOOTING GUIDE			
Stability:	The liquid material is stable for at least 1 year when stored frozen in aliquots with 0.05% sodium azide added as a preservative.	Problem Cause Solution	n		
Caution:	Refer to the enclosed MSDS for information regarding Lectins. The aluminum seals have sharp edges and the vial itself may have cracks which can cause lacerations. Use caution when opening the vial.	Weak or no 1. Low concentration of specific Causes #1 - #3 Staining oligosaccharide on sample. a. Increase incubation time 2. Low concentration of lectin conjugate. b. Increase concentration of lectin conjugate. 3. Insufficient incubation time. a. Avoid exposure to light.	conjugate.		
Remarks: References:	 Bausch, N.J., et.al. (1977) Biochem. 16:5790. Jones, J.M., et.al. J.D. (1973) J. Immunol. 111:1765. Bird, G.W.G., et.al. (1973) Vox Sang, 24:48. 	1. Lectin conjugate is too concentrated. a. Decrease concentration 4. Decrease concentration b. Shorten incubation time 2. Insufficient washing. a. Perform multiple washing High b. Shorten incubation time Background 3. Autofluorescent sample. a. Decrease concentration b. Shorten incubation time a. Perform multiple washing time. a. Verform multiple washing time. a. Use fluorochrome with and emission spectrum. and emission spectrum.	of Lectin conjugate. es. ngs and prolong different excitation		
	5. Dird, G. W.O., Clail. (1775) VOX Bang, 27.76.	Unexpected Staining Pattern Multiple causes b. Use a different lectin co colloidal gold). a. Perform control reaction b. Use other cytochemical or disprove the findings.	ns. technique to prove		
107 North Amphle San Mateo, CA 94	ATORIES, INC. Tel: 650-342-3296 ett Blvd. Fax: 650-342-2648 4401 Orders: 1-800-821-0044 (Outside CA only)	EY LABORATORIES, INC. 107 North Amphlett Blvd. San Mateo, CA 94401 Tel: 650-1 Fax: 650-1 Orders: 1-800	342-3296 342-2648 0-821-0044 side CA only)		

General Procedure

Fluorescent Labeled Lectin



MATERIAL SAFETY DATA SHEET

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

PRODUCT IDENTIFICATION

Name:	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				
Catalog	FP-01, RP-01, TP-01, F-1102 to F-9000, R-1102 to R-9000, T-1102 to T-9000, FA-				
Number (s):	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.				
Synonyms:	Protein A, Avidin (egg white), Glycosylated Bovine Serum Albumin, Lectins,				
	Secondary and Monoclonal Antibodies labeled with FITC, TRITC, or Texas Red®				

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: SPECTUL FIRE FIGHTING CRECTULIONS:

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

Y LABORATORIES, INC.

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S: 1-800-821-0044 (Outside CA only)

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

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

NEACHWITT DATA					
STABILITY:		Stable. Decomposition products are not known to be hazardous.			
HAZARDOUS POLYMERIZATION:		Will NOT occur.			
INCOMPATIBILITY:		Alcohols, strong bases and acids, strong oxidizing agents, and heat. (Lead and copper may react with sodium azide).			
SPILL / LEAK PROCEDURES					
MATERIAL RELEASE /		ith powder or liquid. Clean up spill with a paper towel			
SPILL:	soaked in hour	sehold bleach. Do not allow solutions to dry on			
	environmental su	urfaces. Wash affected area with detergent after the area			
	has been treated	with bleach.			
WASTE DISPOSAL:	Incinerate, autoc	clave, or dispose of paper waste in accordance with all			
		d Federal regulations. Due to the small quantities of ed these products are generally not considered to be			
	environmental h	azards. 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

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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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