PRODUCT INFORMATION Texas Red[®] Labeled Lectin

| | Texas Red Labeled Lectin |
|---|---|
| Catalog Number: | T-1301-2 |
| Description: | Pure Glycine max lectin (SBA) from soybean, Texas Red [®] conjugated. |
| Lot Number: | |
| Protein Concentration: (Based on OD 280) | 2 mg purified SBA Texas $\text{Red}^{\otimes}/2$ ml Buffer. |
| Texas Red [®] / Protein Ratio:(OD 595 / OD 280) | |
| Purification Procedure: | Gel filtration performed after conjugation to remove free Texas Red [®] . |
| Carbohydrate Specificity: | α and $\beta\text{-}$ N-Acetylgalactosamine $>\alpha$ and $\beta\text{-}Galactose.$ |
| Inhibitory Carbohydrate: | Terminal α - and β -N-Acetylgalactosamine > Galactose. |
| Activity: | Less than 4 $\mu g/ml$ will agglutinate fresh A_1 cells. Older B cells can react stronger than A_2 cells. |
| Buffer: | 0.01M Phosphate - 0.15M NaCl, pH 7.2 - 7.4. Contains 0.05% sodium azide as a preservative. |
| Chemical Used for Conjugation: | Texas Red [®] . |
| Storage: | Store liquid material frozen in aliquots in amber vials or covered with foil. Avoid freeze thaw cycles. Clarify by centrifugation. |
| 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. |
| 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. |
| Remarks: | Fluorescent Conjugates are <u>extremely</u> light sensitive. |
| References: | Wada, S., et.al. (1958) J. Biol.Chem. 233 : 395. Lis, H., et.al. (1973) Ann.Rev. of Biochem., Vol.42 : 541 - 574. Lis, H., et.al. (1970) Biochem. Biophys. Acta. 211 : 582. Hammerstrom, et. al. (1977) Biochemistry. Summer, J.B., et. al. (1926) J. Bacteriol. 132 : 227. Den, H., et. al. (1975) J. Cell. Biol. 67 : 826-834. |
| texastred is a registered tr | ademark of Molecular Probes, Inc. |
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107 North Amphlett Blvd. San Mateo, CA 94401 Tel: 650-342-3296 Fax: 650-342-2648 Orders: 1-800-821-0044 (Outside CA only)

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

| | | Tissue Se | ctions | | |
|-------|--|---|--|--|--|
| 1. | Wash and block tissue section. Do not use serum products, they contain glycoproteins which may l 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 tiss | Wash tissue section with Buffer three times. | | | |
| 5. | Examine | tissue section with Fluorescent microscope. U | Jse appropriate filter. | | |
| | Ref. M. Ir | nmbar et. al., (1973). Intnl. Journal of Cancer | r, 12 , 93-99 | | |
| | | Cell Suspen | sion | | |
| 1. | Wash cell | s with Buffer (See reverse side.) | | | |
| 2. | Collect ce | lls by centrifugation. | | | |
| 3. | Dilute Flu | orescent Labeled Lectin to 100 Hg/ml using | Buffer. | | |
| 4. | Incubate approximately 1×10^6 cells with 1 ml diluted Fluorescent labeled Lectin for 15 minutes at room temperature or in a 37°C water bath. | | | | |
| 5. | Wash cell | s with Buffer three times using centrifugation | L. | | |
| 6. | Examine | cells, with or without fixation with Fluorescer | t microscope. Use appropriate filter. | | |
| | Ref. K. Pl | niss. (1977). Experimental Pathology, 14, S15 | 5 | | |
| | | must be protected from light. Perform | incubation, when practical, in a dark room or | | |
| cove | red in foil. | | | | |
| | | Absorption and E | Emission | | |
| | | Absorption/Excitation | Rate Emission Max. | | |
| | | FITC 492 nm | 517 nm | | |
| | | TRITC 554 nm | 570 nm | | |
| | | Texas Red [™] 596 nm | 615 nm | | |
| | | Carbohydrate In | hibition | | |
| Inhib | ition of lec | tin binding may be accomplished by using on | e of two procedures: | | |
| A. | Before ir | cubating with Fluorescent Labeled Leo | tin, incubate section or cells with inhibitory | | |
| B. | carbohydrate for 30-60 minutes at room temperature. NOTE: Complete inhibition may NOT occur. | | | | |
| | room tem | perature before applying to section or cells. | | | |
| | | TROUBLE SHOOT | ING GUIDE | | |
| Pr | oblem | Cause | Solution | | |
| | | 1. Low concentration of specific | Causes #1 -#3 | | |
| | | aliaceacharide an commle | Increase incubation time. | | |
| We | eak or no | oligosaccharide on sample. | | | |
| | eak or no taining | 2. Low concentration of lectin conjugate. | b. Increase concentration conjugate. | | |
| | | Low concentration of lectin conjugate. Insufficient incubation time. | b. Increase concentration conjugate. | | |
| | | Low concentration of lectin conjugate. Insufficient incubation time. Photobleaching | b. Increase concentration conjugate.a. Avoid exposure to light. | | |
| | | Low concentration of lectin conjugate. Insufficient incubation time. | b. Increase concentration conjugate. | | |



Multiple causes

High

Background

Unexpected

Staining

Pattern

2. Insufficient washing.

3. Autofluorescent sample.

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a. Perform multiple washings and prolong

a. Use fluorochrome with different excitation

b. Use a different lectin conjugate (enzyme or

b. Use other cytochemical technique to prove

washing time.

colloidal gold).

and emission spectrum.

a. Perform control reactions.

or disprove the findings.

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® 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. Al liquids are completely miscible in water and biological buffers.

FIRE AND EXPLOSION HAZARDS

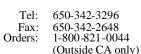
Not considered to be a bire hazard. At high concentrations the chemicals may emit toxic fumes. Such high concentrations are not normally found in a research laboratory.

EXTINGUISHING MEDIA: SPECIAL FIRE FIGHTING PRECAUTIONS:

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

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.

environmental hazards. All of these proteins are fully biodegradable.

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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