

# Biotin Labeled Lectin

## Product Description

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|---|---|
| <b>Name :</b>                                   | <b>Pure Griffonia simplicifolia lectin (GS-I ), Biotin conjugated.</b>  |
| <b>Catalog Number:</b>                          | GS-I Biotin<br>FP-MS8800, 2mg   |
| <b>Protein Concentration (based on OD 280):</b> | 2 mg purified GS-I Biotin/vial. Reconstitute with Buffer to a final concentration of 1mg/ml if lyophilized  |
| <b>Purification Procedure:</b>                  | Gel filtration performed after conjugation to remove free FITC.   |
| <b>Carbohydrate Specificity:</b>                | Melibiose, $\alpha$ -D-Galactose  |
| <b>Inhibitory Carbohydrate:</b>                 | $\alpha$ -D-Glucose.  |
| <b>Activity:</b>                                | 20-30 $\mu$ g/ml is required to agglutinate fresh type B blood cells. Lectin activity against all blood types increases after neuraminidase treatment of the cells. |
| <b>Buffer:</b>                                  | 0.01M Phosphate- 0.15M NaCl containing 0.5 mM CaCl <sub>2</sub> , pH 7.2 - 7.4  |
| <b>Chemical Used for Conjugation:</b>           | Biotinyl N - hydroxysuccinimide ester (BNOHSE).   |

**Storage:** Store lyophilized powder refrigerated at 5-8°C or frozen. Store liquid frozen in aliquots. Avoid freeze-thaw cycles.

**Stability:** The lyophilized material is stable for several years when stored frozen. After reconstitution the material is stable for at least 1 year when stored frozen in aliquots with 0.05% sodium azide added as a preservative.

## Directions for use

### Guidelines for use

The following is a general Procedure and Trouble-Shooting Guide. The information is provided only for your convenience.

1. Wash and block tissue section. It is recommended that 1% purified Bovine Serum Albumin (BSA) or defatted milk powder be used for blocking to prevent non-specific binding. Do not use serum products, they contain glycoproteins which may lead to high levels of non specific background. After blocking, rinse briefly with recommended Buffer.
2. Dilute **Biotin Labeled Lectin** to desired concentration of 5-50  $\mu$ g/ml using recommended Buffer. Incubate section or blot for 30-90 minutes at room temperature in a moist chamber. Slightly longer incubation times may be required if incubation is done at 2-8°C. Rinse 3 times, 5 minutes each time with recommended Buffer.
3. Dilute and incubate Avidin Conjugate according to manufacturer directions.  
**Notes:** Inhibition of lectin binding may be accomplished by using one of two procedures:  
 A. Before proceeding to **Step #3** incubate lectin treated section or blot with inhibitory carbohydrate for 30-60 minutes at room temperature. NOTE: Complete inhibition may not occur.  
 B. Preincubate diluted **Biotin Labeled Lectin** with inhibitory carbohydrate for 30-60 minutes at room temperature before applying to section or blot.

| Problem                     | Cause  | Solution  |
|-----------------------------|--|---|
| Weak or no Staining         | <ol style="list-style-type: none"> <li>1. Low concentration of specific oligosaccharide on sample.</li> <li>2. Low concentration of lectin conjugate.</li> <li>3. Low concentration of avidin conjugate.</li> <li>4. Insufficient incubation time.</li> <li>5. Inappropriate treatment of sample prior to labeling.</li> </ol> | Causes #1 - #4<br><ol style="list-style-type: none"> <li>a. Increase incubation time.</li> <li>b. Increase concentration of sample (on blot) lectin conjugate and/or avidin conjugate.</li> <li>a. Treat section or blot with a different blocking reagent.</li> </ol>  |
| High Background             | <ol style="list-style-type: none"> <li>1. Lectin conjugate and/or avidin conjugate is too concentrated.</li> <li>2. Insufficient washing.</li> <li>3. Insufficient blocking.</li> <li>4. Sample contains endogenous enzymatic activity.</li> </ol>   | <ol style="list-style-type: none"> <li>a. Decrease concentration of respective reagents.</li> <li>b. Shorten incubation times.</li> <li>a. Perform multiple washings and prolong washing time.</li> <li>a. Treat section or blot with a different blocking reagent.</li> <li>a. Determine if sample contains activity which would give background staining in the absence of the avidin conjugate.</li> </ol> |
| Unexpected Staining Pattern | Multiple causes  | <ol style="list-style-type: none"> <li>a. Perform control reactions.</li> <li>b. Use other cytochemical technique to prove or disprove the findings.</li> </ol>   |

## References

1. Murphy, L. A. and Goldstein, I. J. (1977). J. Biol. Chem. 252 : 4739-4742.
2. Judd, W. J., et al. (1978). Transfusion (Philadelphia). 18: 274-280.
3. Eckhardt, A. E., et al. (1982). Cancer Res. 42 : 2977-2979.
4. Maddox, D. E., et al. (1982). PNAS. 79 : 166-170.

## Related products

- ConA-FITC, [FP-47496A](#)
- ConA-Cy3, [FP-WT8680](#)
- ConA-Biotin, [FP-MS9690](#)
- GS-I-FITC, [FP-MS9020](#)
- PNA-FITC, [FP-BV4181](#)
- WGA-biotin, [FP-MS5730](#)
- WGA-SR101, [FP-MS9540](#)
- WGA-FITC, [FP-CE8070](#)
- Other reagents: BSA, [UPQ84170](#)
- Streptavidin – FluoProbes 547H, [FP-CA5570](#)

## Ordering information

[Catalog size quantities and prices may be found at www.interchim.com/](http://www.interchim.com/)

Please inquire for higher quantities (availability, shipment conditions).

For any information, please ask : FluoProbes® / Interchim; Hotline : +33(0)4 70 03 73 06

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