

## HRP labeled PNA

### Product Description

<b>Name :</b>	<b>Pure <i>Arachis hypogaea</i> lectin (PNA), Horseradish Peroxidase conjugated</b>
<b>Catalog Number :</b>	FP-MS9210      2 mg purified PNA HRP / 2 ml Buffer
<b>Carbohydrate Specificity :</b>	Terminal β-Galactose
<b>Inhibitory Carbohydrate :</b>	Lactose > β-Galactose
<b>Activity :</b>	Less than 1 µg/ml will agglutinate human erythrocytes neuraminidase human erythrocytes neuraminidase
<b>Buffer :</b>	0.01M Phosphate - 0.15M NaCl, pH 7.2 - 7.4

**Storage:** Store liquid material frozen in aliquots in amber vials or covered with foil. Avoid freeze thaw cycles. Clarify by centrifugation. Protect from light and moisture. No preservatives have been added. Sodiumazide will inactivate the enzyme, peroxidase.

**Stability:** The liquid material is stable for at least 1 year when stored frozen in aliquots.

### Directions for use

#### Chemical Principle

Peroxidase + H<sub>2</sub>O<sub>2</sub> → Complex  
 Complex + AH<sub>2</sub> (donor) → Peroxidase + H<sub>2</sub>O + A (colored)

#### Assays Reagents

**BUFFER:** 0.01M Sodium phosphate, pH 6.0.  
**ENZYME:** Dilute with Buffer. Acceptable dilution: 1-2 µg/ml.  
**DYE:** 1% o-dianisidine in methanol prepared fresh daily. Store in amber bottle or wrapped in foil.  
**SUBSTRATE:** Prepare 0.3% H<sub>2</sub>O<sub>2</sub> solution in deionized or distilled water from stock H<sub>2</sub>O<sub>2</sub> solution Prior to use dilute to a final concentration 0.003% in Buffer.

#### Procedure

1. Add 0.05 ml of DYE to 6.0 ml of SUBSTRATE. Add 2.9 ml to Reaction test tube and 2.9 ml to Control test tube.
2. At time=0, add 100µl of diluted ENZYME to Reaction tube and 100µl PBS to Control tube. Mix thoroughly.
3. Measure and record optical density at 460nm (OD460) every 15 seconds for 3 minutes, or take the end point reading after 3 minutes by stopping the reaction with 100µl of concentrated NaN<sub>3</sub>.
4. Use this value to determine the rate of change in absorbance per minute.

#### Enzyme Activity Calculations

One unit of peroxidase activity is that amount of enzyme decomposing 1 µmole of peroxide/minute at 25°C. 11.3 x 10<sup>3</sup> cm<sup>-1</sup> is the molar absorbance of H<sub>2</sub>O<sub>2</sub>.

$$OD460/min = \frac{OD460/3min - OD\ Control/3minutes}{3minutes}$$

$$mg\ enzyme/ml\ reactionmixture = \frac{[enzyme\ dilution]}{30}$$

$$units/mg = \frac{OD460/min}{11.3 \times mg\ enzyme/ml\ reactionmixture}$$

FT-MS9210

## Caution

Due to inhibitory sugar present in the conjugates solution, to dilute the Conjugate 50-100 times with buffer before assay.

## References

- Cooper H. *et al.*, *Human Pathology*, **15**:904-906 (1984)
- Moller P. *et al.*, *Virchows Arch.*, **396**:313-317 (1982)
- Vierbuchen M. *et al.*, *Laboratory Inv.*, **48**:(2):181 (1983)
- Ree H. *et al.*, *Cancer.*, **51**:1631 (1983)

## Related / associated products and documents

See [BioSciences Innovations catalogue](#) and [e-search tool](#).

- [Lectin List](#) (or search [conjugated lectins](#)):
- ConA-Biotin, [FP-MS9690](#); -FITC, [FP-47496A](#);
- -Cy3, [FP-WT8680](#),
- GS-I-FITC; [FP-MS9020](#)
- PNA-FITC, [FP-BV4181](#)
- WGA-biotin, [FP-MS5730](#); -SR101, [FP-MS9540](#); -FITC, [FP-CE8070](#)

## Ordering information

[Catalog size quantities and prices may be found at 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

**Disclaimer :** Materials from FluoProbes® are sold **for research use only**, and are not intended for food, drug, household, or cosmetic use. FluoProbes® is not liable for any damage resulting from handling or contact with this product.