SuperLight[™] Atto HRP Reagent

Ultra-Sensitive Luminescent Detection of HRP Conjugates

DESCRIPTION

PEROXIDASES (or peroxide reductases, EC number 1.11.1.x) are a large group of enzymes that break up peroxides and play important roles in various biological processes. Horse Radish Peroxidase (HRP) is an enzyme commonly used for conjugation. One example is antibody-HRP conjugate that is frequently and widely used in immunoassays such as ELISA and Western blots. BioAssay Systems' SuperLight[™] Atto HRP Reagent provides an ultra-sensitive luminescent means for detecting HRP activity and its conjugates. The improved reagent rapidly reacts with peroxidase or the HRP in an HRP-conjugate, producing highly stable and intense luminescence for maximum detection of the peroxidase activity in solution assavs.

KEY FEATURES

Single reagent: Single, stable, and ready to use.

High sensitivity: Detection of HRP in the atto- to low femtogram/well range

APPLICATIONS

HRP determination in ELISA, Southern Blot, Western Blot.

KIT CONTENTS (200 TESTS IN 96-WELL PLATES)

SLAP Reagent: 20 mL. Cat#: SLAP-200

For 200 tests in 96well plate or 800 tests in 384well plate.

For bulk orders of >500mL reagent, please inquire by email order@bioassaysys.com.

Storage conditions. The kit is shipped at room temperature. Store reagent at -20°C upon receiving. Use within 6 months after receipt.

Precautions: reagents are for research use only. Normal precautions for laboratory reagents should be exercised while using the reagents. Please refer to Material Safety Data Sheet for detailed information.

PROCEDURES

Important: Prior to assays, let the SLAP Reagent equilibrate to room temperature. Although not essential, use of a multi-channel pipette is recommended for adding Reagent to wells.

A.96-Well Assays

1. For ELISAs, after washing off free HRP detection antibody, remove any remaining wash buffer. Add 90 µL of the SLAP Reagent to each well.

For liquid samples, add 10 µL of sample per well. Add 90 µL of the SLAP Reagent to each well.

2. Immediately tap plate to mix. Read luminescence after 10min.

B. 384-Well Assays

1. For ELISAs, after washing off free HRP detection antibody, remove any remaining wash buffer. Add 25 µL of the SLAP Reagent to each well.

For liquid samples, add 5 µL sample per well. Add 25 µL of the SLAP Reagent to each well.

2. Immediately tap plate to mix. Read luminescence at 10min

MATERIALS REQUIRED, BUT NOT PROVIDED

Pipetting devices and accessories (e.g. multi-channel pipettor), solid white flat-bottom 96-well plates, centrifuge tubes, and plate reader capable of measuring light intensity.



Free HRP Standard Curve in A 96-Well Plate Assay

RELATED PRODUCTS

- QuantiFluo™ HRP Reagent (QFHRP-25mL), single-reagent, OD570nm, FL530/585nm.
- Quantichrom™ HRP Detection Reagent (DTMB-200), colorimetric, OD 450nm.
- QuantiFluo™ ALP Reagent (QFALP-6mL), single reagent for Alkaline Phosphatase detection, FL360/450nm.

LITERATURE

- 1. Lin, A.V. (2015). Direct ELISA. Methods Mol Biol. 1318:61-7.
- 2. Tobos, C.I. et al (2019). Sensitivity and binding kinetics of an ultrasensitive chemiluminescent enzyme-linked immunosorbent assay at arrays of antibodies. J Immunol Methods. 474:112643.
- 3. Dong, J. et al. (2019). Enhanced chemiluminescence enzyme-linked immunoassay for the determination of DNA methyltransferase 1 in human serum. Luminescence. 34(3):368-374.