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Product Information

SIGMA*FAST*™ OPD

Tablet set

P9187

Product Description

The SIGMAFAST[™] OPD (*o*-phenylenediamine dihydrochloride) tablet set is designed for use as a soluble substrate to detect horseradish peroxidase (HRP) activity in Enzyme Immunoassays (EIA and ELISA).¹⁻³ EIA/ELISA applications utilizing OPD may be read in timed assays or stopped with dilute acid solutions for delayed readings. The SIGMAFAST[™] OPD tablet set requires only the addition of water, with no additional buffers or steps, to prepare an active substrate solution.

Several dissertations⁴⁻⁹ have cited use of product P9187 in their research protocols.

Precautions and Disclaimer

For R&D use only. Not for drug, household, or other uses. Please consult the Safety Data Sheet for information regarding hazards and safe handling practices.

Components

Each SIGMAFAST[™] OPD tablet set consists of one OPD tablet (silver foil) and one urea hydrogen peroxide tablet (gold foil). The tablets are individually packaged in foil packets.

Storage/Stability

Store the tablet sets at 2-8 °C. The tablets are very hygroscopic and light-sensitive, and should be stored in the protective foil packets.

The Substrate Solution is light-sensitive and should be protected from direct sunlight or UV sources in a tightly capped amber bottle.

Preparation Instructions

Each tablet set, when dissolved in 20 mL of water, provides 20 mL of ready-to-use substrate with final concentrations of:

- 0.4 mg/mL OPD
- 0.4 mg/mL urea hydrogen peroxide
- 0.05 M phosphate-citrate, pH 5.0
- 1. Remove the required number of OPD and urea hydrogen peroxide tablets for the assay.
- 2. Return the box to the refrigerator.
- 3. Allow the tablets to reach room temperature.
- 4. Open one OPD tablet package (silver foil) and one urea hydrogen peroxide tablet package (gold foil).
- Drop the tablets into an amber bottle containing 20 mL of water. Avoid skin contact with the tablets.
- 6. Vortex until dissolved.

The Substrate Solution is now ready for use. For best results, the Substrate Solution should be used within one hour.

Procedure

- 1. After the EIA/ELISA reaction with the HRP-conjugated antibody is completed, wash the plate thoroughly to remove unbound conjugate.
- 2. Add 200 μ L of the Substrate Solution to each well. Incubate the plate, in the dark, for 30 minutes at room temperature.
- 3. After the incubation period, read the plate at 450 nm on a multiwell plate reader.
- 4. If the plate cannot be read immediately, add 50 μL of 3 M HCl or 3 M H_2SO_4 solution per 200 μL of solution. Read stopped reactions at 492 nm.



Troubleshooting

If the background is too high:

- Use a blocking step prior to the application of the primary antibody. Normal serum (5% v/v) from the same species as the host of the second antibody generally produces the best results.
- 2. Additional blocking agents for an ELISA are:
 - 0.05% TWEEN[®] 20 in 50 mM TBS, pH 8.0 (Cat. No. T9039).
 - 1% BSA containing 0.05% TWEEN[®] 20 in 50 mM TBS, pH 8.0.
 - 3% nonfat-dried milk in 0.01 M TBS (Cat. No. P2194). Do not use milk as a blocking agent when using avidin-biotin systems.
- 3. Use 0.05% TWEEN[®] 20 in all washing and antibody diluent buffers.
- 4. Run control wells without the primary antibody to check for nonspecific reactivity of the secondary antibody.
- 5. Titer the primary antibody and the conjugate to optimize working dilutions.

If no color develops or the color is too faint:

- 1. Adjust the concentration of the primary antibody.
- 2. Adjust the concentration of the secondary antibody.
- 3. Determine if the enzyme conjugate is active by mixing a small sample of substrate and conjugate together in a tube.
- 4. Increase the reaction time or temperature.
- 5. Adjust the concentration of the coating antigen.
- 6. Consider using an amplifying system such as avidin-biotin.

References

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