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# Preparation of staphylococcal protein-A conjugated to horseradish peroxidase by the periodate method.

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Protocol status: Working We use this protocol and it's working

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### Abstract

This reagent can be used in ELISA, Western blotting and Dot blot to detect antigens and antibodies. It is important in the immunodiagnosis of infectious diseases and other problems. I find this useful in the detection of anti-HIV antibodies by ELISA.

## Guidelines

All reagents but specially the enzyme and sodium periodate solution has to be prepared freshly before mixing it with the enzyme.

## Materials

MATERIALS

- X Ammonium Sulfate P212121
- Sodium periodate Bio Basic Inc. Catalog #SB0875.SIZE.100g
- Sodium borohydride Sigma Aldrich Catalog #452882
- X Horseradish Peroxidase (HRP) type IV Sigma Aldrich Catalog #P8375-25KU
- Staphylococcal Protein-A Sigma Aldrich

Pipettes 20ml to 1000 ml glass Scale Incubator Refrigerator Freezer Centrifuges

- 1 Horseradish peroxidase (500 μg in 50 μl NaCO3 , pH 9.6) is mixed with freshly made sodium periodate solution (1.71 mg/ml) followed by incubation in the dark for 2 h.
- 2 Mix 500 μg of staphylococcal protein-A (SpA) with an equal amount (500 micrograms) of a mix of horseradish peroxidase-sodium periodate.
- 3 The mixture is incubated for 3 hours at 4°C with gentle agitation.
- 4 Forty μl of freshly prepared NaBH4 solution (5 mg NaBH4 /ml 0.1 mM NaOH) is then added to the preparation.
- 5 The preparation is incubated for 90 min at 4°C in the dark with gentle agitation.
- 6 Cold 50% saturated ammonium sulphate solution (pH 7.4) is added drop by drop in the ratio 1:1 (v/v).
- 7 The mixture is then centrifuged for 25 min at 4°C and recover the pellet at the bottom of the tube.
- 8 The pellets is re-suspended in 200 μl of PBS pH=7.4 and dialysed against 1L of PBS for 24 h with 3 buffer changes.
- 9 An equal volume of glycerol is added to the dialysate followed by 100 μl of bovine serum albumin, BSA (20 mg/ ml).
- 10 The conjugate is then stored at -20°C until further used.

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