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

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1 Works for me dx.doi.org/10.17504/protocols.io.bjg8kjzw

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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.

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#### PROTOCOL CITATION

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## MATERIALS

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

MATERIALS TEXT

**Pipettes** 

20ml to 1000 ml glass

Scale

Incubator

Refrigerator

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Freezer Centrifuges

SAFETY WARNINGS

Pay attention to all details as the times of reactions among the proteins involved in this preparation. It will prevent over-oxidation. The average time of preparation is 18 hours.

BEFORE STARTING

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

1	Horseradish peroxidase (500 $\mu$ g in 50 $\mu$ l 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 $\mu l$ 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 $\mu$ l 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 $\mu$ l of bovine serum albumin, BSA (20 mg/ ml).
10	The conjugate is then stored at -20°C until further used.

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