

Method for ABC Technique using Monoclonal Antibodies on Frozen Sections

The following method is recommended for mouse monoclonal antibodies from Novocastra Laboratories Ltd. Customers should determine optimal dilutions for primary and secondary antibodies as these may vary according to the application.

REAGENTS

1. Standard solvents used in immunohistochemistry.
2. Hydrogen peroxide 30% w/v.
3. 50mM Tris buffered saline (TBS) pH7.6.
4. Normal rabbit serum.
5. Biotinylated secondary antibody.
6. ABC reagents.
7. DAB reagent.

EQUIPMENT

1. Appropriate incubators.
2. Standard immunohistochemistry facilities.
3. Pressure cooker or other appliance used for unmasking antigens - if required.

PROCEDURE

1. Air dry frozen sections for 60 minutes or overnight.
2. Fix sections in acetone for 10 minutes.
3. Air dry sections for 10 minutes.
4. Wash sections in 50mM Tris buffered saline pH7.6 for 5 minutes.
5. Cover sections with blocking reagent eg 10% v/v normal rabbit serum in TBS for 10 minutes.
6. Remove excess blocking reagent and replace with primary antiserum diluted in blocking reagent as required (see datasheet), for 60 minutes at 25°C or overnight at 4°C.
7. Wash in TBS buffer for 2 x 5 minutes.
8. Remove excess TBS buffer and incubate sections with biotinylated rabbit anti-mouse secondary diluted in blocking reagent for 30 minutes at 25°C.
9. Wash in TBS buffer for 2 x 5 minutes.
10. Remove excess TBS buffer and incubate sections with ABCComplex/HRP for 30 minutes at 25°C.
11. Wash in TBS buffer for 2 x 5 minutes.
12. Develop with 3 3' diaminobenzidine tetrahydrochloride (DAB).
13. Rinse slides in water.
14. Counterstain with Haematoxylin (if required).
15. Dehydrate, clear and mount sections with DPX mountant.

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