## Identification of IL1 beta in PLP-Fixed, Paraffin-Embedded Mouse Tissue

## **Reagents:**

1X Automation Buffer
3% Hydrogen Peroxide
Antibody Diluent
Citrate Buffer
DAB Chromagen
Hematoxylin
PLP fixative

## **Antibody Information**

Primary antibody: Biotin-conjugated Rabbit anti-IL1 beta Antigenix America Huntington Sta, NY 11746 1-800-558-1008 Catalog # RMF326B

Negative Control: Biotin conjugated normal rabbit serum Vector Laboratories 30 Ingold Rd Burlingame CA 94010 1-800-227-6666 Catalog # BI-1005

Label: Biogenex supersensitive label Biogenex San Ramon CA 94583 Catalog # HK-330-9K

Comment: the following protocol with the listed antibody works best in tissues fixed overnight in PLP (periodate-lysine-paraformaldehyde) fixative.

Bouin's-fixed tissue are applicable for this procedure. Formalin and zinc-formalin fixation is not acceptable for this commercial antibody.

## **Staining Procedure**

-Positive Control Tissue: tissue treated with an immunogen such as LPS

-Stain Localization: Cytoplasmic

Deparaffinize and hydrate slides through the following solutions.

Xylene	2 times	5 minutes
100% EtOH	2 times	3 minutes
95% EtOH	2 times	3 minutes
1X Automation Buffer	2 times	5 minutes

1. Quench endogenous peroxidase by placing slides in 3% hydrogen peroxide for 15 minutes.

2.	Rinse	slides	in 2	changes	of	1X	Automation	Buffer	for:	5 minutes	each.

2. Rinse slides in 2 changes of 1X Automation Buffer for 5 minutes each.
3. Unmasking Techniques: Steamer Place slides in 1X Citrate Buffer and steam for 35 minutes. Remove slides from steamer and cool for 20 minutes.Temp Stop reaction by rinsing slides in D/W. Place slides in 1X Automation buffer for 5 minutes.
4. Apply primary antibody (Biotinylated rabbit anti-IL beta) at a 1:30 dilution and incubate for 1 hr at room temperature.  Lot# Exp Date
For negative control slides, normalize the protein concentration of biotin-conjugated normal rabbit serum to the protein concentration of the primary antibody.  Lot# Reconstituted Date (note: if you are reconstituting a new bottle of the serum DO NOT VORTEX)
5. Rinse slides in 2 changes of 1X Automation Buffer for 5 minutes each.
6. Apply Biogenex super sensitive label and incubate for 30 minutes.  Lot# Exp. Date

7. Rinse slides in 2 changes of 1X Automation Buffer for 5 minutes each.

8. Apply liquid Dako	DAB Chromagen for 6 mi	nutes in the	dark.	
(Add 1 drop of DAB	per ml of substrate)			
Lot#	Exp. Date	New Kit:	yes /	no

- 9. Rinse in tap water 3 minutes.
- 10. Counterstain with Modified Harris Hematoxylin for 1 min.
- 11. Rinse in tap water until water is clear.
- 12. Place slides in 1X Automation Buffer for 1 minute with gentle agitation to blue slides.
- 13. Dehydrate through the following solutions.

95% Ethanol	1 change	3 minutes
100% EtOH	3 changes	3 minutes
Xylene	2 changes	5 minutes

14. Coverslip

updated 03/23/04