## Detection of PKC Theta in Formalin-Fixed, Paraffin- Embedded Mouse Tissue

## **Reagent and Antibody Information**

1X Wash Buffer
3% Hydrogen Peroxide
1% BSA Diluent
1X Citrate Buffer
DAB Chromagen
Hematoxylin

Blocking Solution: Dakocytomation Protein Block Serum-Free Ready-To-Use Dakocytomation Corporation
Carpinteria CA 93013
www.dakousa.com
1-800-235-5763

Avidin / Biotin Blocking Kit Vector Laboratories, Inc. Burlingame, CA 94010 www.vectorlabs.com 1-800-227-6666 Catalog # SP-2001

Code No. X0909

Primary Antibody: Mouse Anti-PKCØ Antibody
BD Pharmingen
Distributed by Transduction Labs
Lexington, KY 40511
www.bdpharma.com
1-800-227-4063
Catalog # 610090

Negative Control Serum: Normal Mouse Serum Jackson Immunoresearch Laboratories, Inc. West Grove, PA 19390 www.jacksonimmuno.com 1-800-367-5296 Catalog # 015-000-001

Staining Kit: LSAB+ System-HRP
Dakocytomation Corporation
Carpinteria CA 93013
www.dakousa.com
1-800-235-5763
Code No. K0690

**Note**: This kit includes reagents needed for the secondary antibody (link) and label complex.

## **Staining Procedure**

Positive Control Tissue: Spleen

Stain Localization: Cytoplasmic and cell membrane

1. Deparaffinize and hydrate slides through the following solutions:

Solution	Repetitions	Time
Xylene	2 times	5 minutes
100% Ethanol	2 times	3 minutes
95% Ethanol	2 times	3 minutes
1X Wash Buffer	2 times	5 minutes

- 2. Quench endogenous peroxidase by placing the slides in 3% hydrogen peroxide for 15 minutes.
- 3. Rinse the slides in 2 changes of 1X Wash Buffer for 5 minutes each.

4.	Heat-Induced Epitope Retrieval Using The Decloaker Add 500 ml of distilled water to the pan inside the decloaker.  Place a full rack of slides into a Tissue Tek® container with 200 ml of 1X citrate buffer (Insert blank slides into any empty slots in the rack to ensure even heating of slides)  Place the container stably inside the pan and decloak for 5 minutes. Maximum Pressure  Depressurize for 10 minutes.  Remove pan top and cool for 10 minutes. Temperature Before Cooling Slides  Rinse the slides in 2 changes of distilled water for 3 minutes each time.
5.	Rinse the slides in 2 changes of 1X Wash Buffer for 5 minutes each time.
	Block with the Dako Protein Blocking Reagent for 10 minutes at room temperature.  Lot # Exp Date  DO NOT RINSE THE SLIDES. CONTINUE TO AVIDIN-BIOTIN BLOCK.
7.	Avidin / Biotin Blocking Kit  Lot # Exp Date New Kit: yes / no  Apply avidin block for 15 minutes at room temperature.  Quick rinse in 1X Wash Buffer.  Apply biotin block for 15 minutes at room temperature.  DO NOT RINSE SLIDES WITH BUFFER BEFORE ADDING PRIMARY ANTIBODY.  ONLY WIPE EXCESS BUFFER.
8.	Apply the primary antibody at a 1:10 dilution, and incubate for 1 hour at room temperature.  Lot # Date Aliquoted

For negative control slides, dilute the protein concentration of the normal rabbit serum to match that of the primary antibody. Make a 1:10 dilution from this normalized serum, and apply to the slides.

9. Rinse the slides in 2 changes of 1X Wash Buffer for 5 minutes each	ch.
LSAB+ Kit	
Lot # Exp Date	
10. Apply the Link (yellow bottle) from the LSAB+ Kit, and incubat temperature.	e for 30 minutes at room
11. Rinse the slides in 2 changes of 1X Wash Buffer for 5 minutes ea	ach.
12. Apply the Label (red bottle) from the LSAB+ Kit, and incubate f	for 30 minutes at room temperature.
13. Rinse the slides in 2 changes of 1X Wash Buffer for 5 minutes ea	ach.
14. Apply the DAB chromagen, and incubate in the dark for 6 minute (Add 1 drop of DAB per ml of substrate)	es at room temperature.
Lot # Exp Date New Ki	it: yes / no

- 16. Counterstain with Harris Hematoxylin for 20 seconds.
- 17. Rinse the slides in tap water until water is clear.
- 18. Gently agitate slides in 1X Wash Buffer until they turn blue.
- 19. Dehydrate through the following solutions:

Solution	Repetitions	Time
95% Ethanol	1 time	3 minutes
100% Ethanol	3 times	3 minutes
Xylene	2 times	5 minutes

20. Coverslip

Updated 07/07/05