Detection of Chromogranin A in Formalin-Fixed, Paraffin-Embedded Rat 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 Code No. X0909

A '1' /D' /' D1 1' T

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

Primary Antibody: Mouse Anti-Chromogranin A Antibody

Lab Vision / Thermo Fisher Scientific
Fremont, CA 94539
www.labvision.com
1-800-828-1628
Catalog # MS-324-P

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: Islets of pancreas

Stain Localization: Cytoplasmic

1. Deparaffinize and hydrate slides through the following solutions:

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. <i>Maximum Pressure</i>							
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.	5. Rinse the slides in 2 changes of 1X Wash Buffer for 5 minutes each time.							
6. Block with the Dako Protein Blocking Reagent and incubate for 10 minutes at room ter								
	Lot # Exp Date							
DO NOT DINGE THE GUIDEG CONTINUE TO AMDIN DIOTIN DUOCK								
DO NOT RINSE THE SLIDES. CONTINUE TO AVIDIN-BIOTIN BLOCK.								
7	Avidin / Biotin Blocking Kit							
′ •	Lot # Exp DateNew Kit: yes / no							
	Apply avidin block - 15 minutes at room temperature.							
	Quick rinse in 1X Wash Buffer.							
	Apply biotin block - 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 1:25 dilution and incubate for 30 minutes at room temperature.							
	Lot #Exp Date							

primate for 30	ary antibody. Make 0 minutes at room t	des, dilute the protein concentration of the normal mouse serum to the a 1:25 dilution from this normalized serum and apply to the slides temperature. Date Reconstituted					
9. Rinse	se the slides in 2 changes of 1X Wash Buffer for 5 minutes each.						
LSA	B+ Kit						
Lot #	#	Exp Dat	e				
10. Арр	oly the Link (yellow	v bottle) from t	he LSAB+ Ki	and incubate f	or 15 minutes at room to	emperature	
11. Rir	se the slides in 2	changes of 1	X Wash Buff	er for 5 minute	es each.		
12. App	oly the Label (red b	ottle) from the	LSAB+ Kit an	nd incubate for	15 minutes at room tem	perature.	
13. Rin	se the slides in 2 ch	anges of 1X W	Vash Buffer for	5 minutes eacl	1		
(Ad	oly the DAB chrom d 1 drop of DAB p #	er ml of substr	ate)		at room temperature. yes / no		
15. Rin	se the slides in tap	water 3 minute	es.				
16. Cou	ınterstain with Harr	ris Hematoxyli	n for 30 secon	ds.			
17. Rin	se the slides in tap	water until wat	er is clear.				
18. Gen	atly agitate slides in	1X Wash Buf	fer until they t	urn blue.			
19. Deh	ydrate through the	following solu	ations:				
	95% Ethanol	1 time	3 minutes				
	100% Ethanol	3 times	3 minutes				
	Xylene	2 times	5 minutes				

20. Coverslip

Updated 04/06/06