Detection of MMP-9 in Formalin-Fixed, Paraffin-Embedded Human 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: Rabbit Polyclonal MMP-9 Antibody Lab Vision / Thermo Fisher Scientific Fremont, CA 94539 www.labvision.com 1-800-828-1628

1-800-828-1628 Catalog # RB-9234-P1

Negative Control Serum: Normal Goat Serum Jackson Immunoresearch Laboratories, Inc. West Grove, PA 19390 http://www.jacksonimmuno.com 1-800-367-5296 Catalog # 005-000-121

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: Placenta Stain Localization: Cytoplasmic

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.

Lot #_____ Date Aliquoted_____

4. <u>Heat-Induced Epitope Retrieval Using The Decloaker</u>				
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. <i>Temperature Before Cooling Slides</i>				
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.				
6. 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. Anidia / Distin Distring Vit				
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.				
ONLT WILL EACESS DUTTER.				
8. Apply the primary antibody at a 1:25 dilution, and incubate for 30 minutes at room temperature	د			
o. Typis the primary antibody at a 1.25 dilution, and incubate for 50 minutes at room temperature.				

For negative control slides, dilute the protein concentration of the normal rabbit serum to match that of the primary antibody. Make a 1:25 dilution from this normalized serum, and apply to the slides. Incubate for 30 minutes at room temperature. Lot # Date Reconstituted
9. Rinse the slides in 2 changes of 1X Wash Buffer for 5 minutes each.
LSAB+ Kit Lot # Exp Date
10. Apply the Link (yellow bottle) from the LSAB+ Kit, and incubate for 30 minutes at room temperature.
11. Rinse the slides in 2 changes of 1X Wash Buffer for 5 minutes each.
12. Apply the Label (red bottle) from the LSAB+ Kit, and incubate for 30 minutes at room temperature.
13. Rinse the slides in 2 changes of 1X Wash Buffer for 5 minutes each.
14. Apply the DAB chromagen, and incubate in the dark for 6 minutes at room temperature. (Add 1 drop of DAB per ml of substrate) Lot # Exp Date New Kit: yes / no
15. Rinse the slides in tap water 3 minutes.
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:
---------------	-------------	----------------------

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

20. Coverslip