Detection of CC10 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

Avidin / Biotin Blocking Kit

Vector Laboratories, Inc.
Burlingame, CA 94010
www.vectorlabs.com
1-800-227-6666

Catalog # SP-2001

Primary Antibody: Goat Anti-CC10 (T-18) Polyclonal Antibody

Santa Cruz Biotechnology, Inc.

Santa Cruz, CA 95060

www.scbt.com 1-800-457-3801

Catalog # sc-9772

Negative Control Serum: Normal Goat Serum

Jackson Immunoresearch Laboratories, Inc.

West Grove, PA 19390

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: Bronchiolar airways of the lung (identifies Clara cells)

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.

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|----|--|--|--|--|
| 1. | 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. <i>Temperature Before Cooling Slides</i> | | | |
| | Rinse the slides in 2 changes of distilled water for 3 minutes each time. | | | |
| 5. | 7. Rinse the slides in 2 changes of 1X Wash Buffer for 5 minutes each time. | | | |
| 5. | 6. Block with the Dako Protein Blocking Reagent and incubate 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 #New 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 1:100 dilution and incubate for 30 minutes at room temperature. Lot #______ Date Aliquoted ______

For negative control slides, dilute the protein concentration of the normal goat serum to match that of the primary antibody. Make a 1:100 dilution from this normalized serum and apply to the slides.

| Incubate 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 30 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:

| 95% Ethanol | 1 time | 3 minutes |
|--------------|---------|-----------|
| 100% Ethanol | 3 times | 3 minutes |
| Xylene | 2 times | 5 minutes |

20. Coverslip

Updated 10/25/04