# NEUROMICS

# Collagen I

# **Data Sheet**

Catalog Number:	RA25100	Host:	Rabbit
Product Type:	Affinity Purified	Species Reactivity:	Human, Mouse
Immunogen Sequence:	A synthetic peptide made to an internal portion of the human Collagen I protein (between residues 150-200) [UniProt P02452]	Format:	Liquid. PBS, 30% glycerol with 0.05% Sodium Azide. Concentration 1.1mg/ml.
Applications:	Immunohistochemistry-Paraffin 1:200 *Dilutions listed as a recommendation. Optimal dilution should be determined by investigator.		
Storage:	Store frozen. Aliquot as undiluted antisera and immediately place at -20°C. Antisera may have become trapped in top of vial during shipping. Centrifugation of vial is recommended before opening. Stable for at least 6 months at -20°C. Repeated freeze/thaw cycles compromise the integrity of the antiserum.		

### **Application Notes**

This Collagen 1 antibody is useful for IHC-paraffin embedded sections.

#### Immunohistochemistry-Paraffin Embedded Sections:

#### Antigen Unmasking:

Bring slides to a boil in 10 mM sodium citrate buffer (pH 6.0) then maintain at a sub-boiling temperature for 10 minutes. Cool slides on bench-top for 30 minutes.

Staining:

1. Wash sections in deionized water three times for 5 minutes each.

- 2. Wash sections in wash buffer for 5 minutes.
- 3. Block each section with 100-400 ul blocking solution for 1 hour at room temperature.
- 4. Remove blocking solution and add 100-400 ul diluted primary antibody. Incubate overnight at 4C.
- 5. Remove antibody solution and wash sections in wash buffer three times for 5 minutes each.
- 6. Add 100-400 ul biotinylated diluted secondary antibody. Incubate 30 minutes at room temperature.
- 7. Remove secondary antibody solution and wash sections three times with wash buffer for 5 minutes each.
- 8. Add 100-400 ul Streptavidin-HRP reagent to each section and incubate for 30 minutes at room temperature.
- 9. Wash sections three times in wash buffer for 5 minutes each.
- 10. Add 100-400 ul DAB substrate to each section and monitor staining closely.
- 11. As soon as the sections develop, immerse slides in deionized water.
- 12. Counterstain sections in hematoxylin.
- 13. Wash sections in deionized water two times for 5 minutes each.
- 14. Dehydrate sections.
- 15. Mount coverslips.

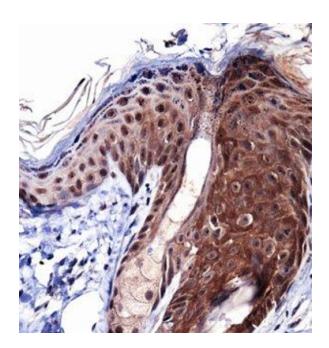
Note: The above information is only intended as a guide. The researcher should determine what protocol best meets their needs. Please follow safe laboratory procedures.

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Image: Collagen 1 staining of mouse epidermis tissue using DAB with hematoxylin counterstain.



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Neuromics Antibodies • 5325 West 74<sup>th</sup> Street, Suite 8 • Edina, MN 55439 phone 866-350-1500 • fax 612-677-3976 • e-mail: <u>pshuster@neuromics.com</u>