abcam

ab316223

Anti-Collagen I antibody [RM1131] - BSA and Azide free

Recombinant | RabMAb | KO Validated

Key facts

Isotype	IgG
Host species	Rabbit
Storage buffer	pH: 7.2 - 7.4 Constituents: PBS
Form	Liquid
Clonality	Multiclonal
Immunogens	The exact immunogen used to generate this antibody is proprietary information.
Carrier free	Yes
Clone number	RM1131
Purification technique	Affinity purification Protein A
Concentration	1.041 mg/mL The concentration of this product may be batch-dependent Batch concentration finder →

Reactivity data

Flow Cyt (Intra)

Tested

Species	Human, Mouse, Rat
Dilution info	-

Notes	-

WB

Tested

Species	Human
Dilution info	-
Notes	Negative control: HT-29. We are unsure how to define these extra bands below 100kDa.

Species	Mouse
Dilution info	-
Notes	Negative control: RAW 264.7. We are unsure how to define these extra bands below 100kDa.

Species	Rat
Dilution info	-
Notes	-

IHC-P

Tested

Species	Human, Mouse, Rat
Dilution info	-
Notes	-

ICC/IF

Tested

Species	Human, Mouse, Rat
Dilution info	-
Notes	-

IP

Tested

Species	Human, Mouse
Dilution info	-
Notes	-

Expected

Species	Rat
Dilution info	Use at an assay dependent concentration.
Notes	-

IHC-Fr

Tested

Species	Mouse, Rat
Dilution info	-
Notes	-

Expected

Species	Human
Dilution info	Use at an assay dependent concentration.
Notes	-

Target data

See full target information COL1A1

Function	Type I collagen is a member of group I collagen (fibrillar forming collagen).

Storage

Shipped at conditions	Blue Ice
Appropriate short-term storage conditions	+4°C

Appropriate long-term storage conditions	+4°C

Notes

ab316223 is the carrirer-free version of ab316222.

What are recombinant multiclonals?
Recombinant multiclonals are a mixture of recombinant antibodies co-expressed from a library of heavy and light chains. They offer several advantages including:

- The sensitivity of polyclonal antibodies by recognising multiple epitopes
- High batch-to-batch consistency and reproducibility
- Improved sensitivity and specificity
- Long-term security of supply
- Animal-free batch production

View our range of recombinant multiclonal antibodies.

Patented technology
Our RabMAb® technology is a patented hybridoma-based technology for making rabbit monoclonal antibodies. For details on our patents, please refer to RabMAb® patents.

Conjugation ready
Our carrier-free antibodies are typically supplied in a PBS-only formulation, purified and free of BSA, sodium azide and glycerol. This conjugation-ready format is designed for use with fluorochromes, metal isotopes, oligonucleotides, and enzymes, which makes them ideal for antibody labelling, functional and cell-based assays, flow-based assays (e.g. mass cytometry) and Multiplex Imaging applications.

Use our conjugation kits for antibody conjugates that are ready-to-use in as little as 20 minutes with 1 minute hands-on-time and 100% antibody recovery: available for fluorescent dyes, HRP, biotin and gold.

Compatibility
This product is compatible with the Maxpar® Antibody Labeling Kit from Fluidigm, without the need for antibody preparation. Maxpar® is a trademark of Fluidigm Canada Inc.

Supplementary info

This supplementary information is collated from multiple sources and compiled automatically.

Activity summary	Collagen type I also called collagen I is a structural protein expressed mainly in connective tissues such as skin tendon bone and ligaments. It serves as an important component in providing mechanical strength and integrity to these tissues. Collagen I is a fibrillar collagen known for its triple-helix structure composed of two alpha-1 chains and one alpha-2 chain and has a molecular mass of approximately 300 kDa. Researchers often employ collagen western blot and collagen ELISA techniques for its detection. Collagen suppliers offer various collagen antibodies used in these assays to study its distribution and function.
Biological function summary	Collagen type I plays a central role in maintaining the extracellular matrix and supporting cellular environments. It interacts with other matrix proteins and cells

	forming complexes that help in tissue development and repair. Type I collagen is especially important in bone matrix working alongside minerals like hydroxyapatite to provide rigidity and support. Anti-collagen antibodies aid in studying its biological functions and interactions which are critical to understanding tissue dynamics.
Pathways	Collagen type I interacts with multiple signaling cascades involved in tissue remodeling and repair. It is a significant player in the TGF-β pathway which regulates fibrosis and wound healing processes. In these pathways proteins such as fibronectin and integrins work in concert with collagen type I to orchestrate cellular responses to damage. Researchers often examine its role in these pathways to uncover therapeutic possibilities for disease interventions.
Associated diseases and disorders	Collagen type I has strong connections to conditions like osteogenesis imperfecta and fibrosis. Mutations or irregularities in collagen I production can lead to osteogenesis imperfecta a genetic disorder characterized by brittle bones. In fibrosis excessive collagen deposition disrupts normal tissue architecture contributing to organ dysfunction. In both conditions type I collagen interacts with other proteins like matrix metalloproteinases which modulate its breakdown and remodeling highlighting its importance in disease pathology.

Product promise

Tested

We have tested this species and application combination and it works. It is covered by our product promise.

Expected

We have not tested this specific species and application combination in-house, but expect it will work. It is covered by our product promise.

Predicted

This species and application combination has not been tested, but we predict it will work based on strong homology. However, this combination is not covered by our product promise.

Not recommended

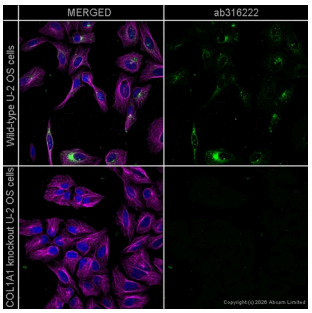
We do not recommend this combination. It is not covered by our product promise.

We are dedicated to supporting your work with high quality reagents and we are here for you every step of the way should you need us.

In the unlikely event of one of our products not working as expected, you are covered by our product promise.

Full details and terms and conditions can be found here:
Terms & Conditions.

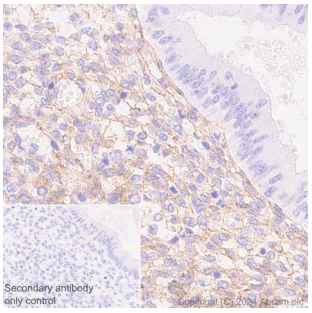
25 product images



Immunocytochemistry/ Immunofluorescence - Anti-Collagen I antibody [RM1131] - BSA and Azide free (ab316223)

Collagen I Immunocytochemistry/ Immunofluorescence staining using rabbit Anti-Collagen I antibody

This data was developed using ab316222 the same antibody clone in a different buffer formulation.
ab316222 staining COL1A1 in Wild-type U-2 OS (positive) and COL1A1 knockout U-2 OS (negative) cells (ab273846). The cells were fixed with 100% methanol (5 min), permeabilized with 0.1% PBS-Triton X-100 for 5 minutes and then blocked with 1% BSA/10% normal goat serum/0.3M glycine in 0.1%PBS-Tween for 1h. The cells were then incubated for 2 hours at room temperature with ab316222 at 1 ug/ml (shown in green) and ab7291, Anti-alpha Tubulin antibody [DM1A] - Loading Control (shown in pseudocolour magenta). Cells were then incubated with ab150081, Goat Anti-Rabbit IgG H&L (Alexa Fluor® 488) preadsorbed at 1/1000 dilution and ab150119, Goat Anti-Mouse IgG H&L (Alexa Fluor® 647) preadsorbed at 1/1000 dilution for 1 hour at room temperature. Nuclear DNA was labeled with DAPI (shown in blue). Also suitable in cells fixed with 4% paraformaldehyde (10 min). Image was acquired with a high-content analyser (Operetta CLS, Revvity) and a maximum intensity projection of confocal sections is shown.



Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) - Anti-Collagen I antibody [RM1131] - BSA and Azide free (ab316223)

This data was developed using ab316222, the same antibody clone in a different buffer formulation.

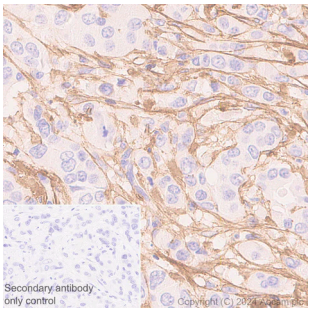
Immunohistochemical analysis of paraffin-embedded Human endometrium tissue labeling Collagen I with ab316222 at 1/500 (1.012 ug/ml) dilution, followed by a ready to use LeicaDS9800 (Bond™ Polymer Refine Detection).

Positive staining on stroma of human endometrium.
The section was incubated with ab316222 for 30 mins at room temperature.
The immunostaining was performed on a Leica Biosystems BOND® RX instrument

Counterstained with Hematoxylin.

Secondary antibody only control: Secondary antibody is a ready to use LeicaDS9800 (Bond™ Polymer Refine Detection).

Heat mediated antigen retrieval was performed with Tris-EDTA buffer (pH 9.0, Epitope Retrieval Solution2) for 20 mins



Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) - Anti-Collagen I antibody [RM1131] - BSA and Azide free (ab316223)

This data was developed using ab316222, the same antibody clone in a different buffer formulation.

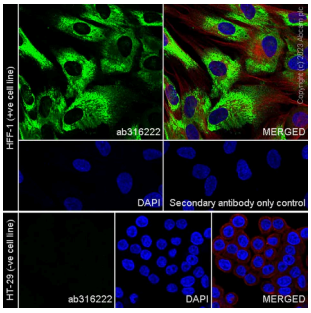
Immunohistochemical analysis of paraffin-embedded Human breast cancer tissue labeling Collagen I with ab316222 at 1/500 (1.012 ug/ml) dilution, followed by a ready to use LeicaDS9800 (Bond™ Polymer Refine Detection).

Positive staining on stroma of human breast cancer.
The section was incubated with ab316222 for 30 mins at room temperature.
The immunostaining was performed on a Leica Biosystems BOND® RX instrument

Counterstained with Hematoxylin.

Secondary antibody only control: Secondary antibody is a ready to use LeicaDS9800 (Bond™ Polymer Refine Detection).

Heat mediated antigen retrieval was performed with Tris-EDTA buffer (pH 9.0, Epitope Retrieval Solution2) for 20 mins



Immunocytochemistry/ Immunofluorescence - Anti-Collagen I antibody [RM1131] - BSA and Azide free (ab316223)

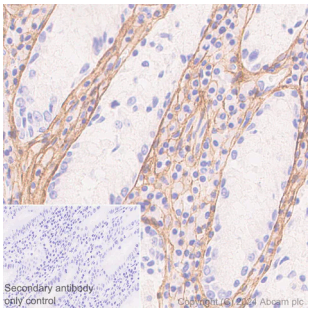
This data was developed using ab316222, the same antibody clone in a different buffer formulation.

Immunofluorescent analysis of 4% Paraformaldehyde-fixed, 0.1% TritonX-100 permeabilized HFF-1 (human skin fibroblast) cells labelling Collagen I with ab316222 at 1/100 (5.06 ug/ml) dilution, followed by ab150081 Goat Anti-Rabbit IgG H&L (Alexa Fluor® 488) preadsorbed antibody at 1/1000 (2 ug/ml) dilution (Green).

Confocal image showing cytoplasmic staining in HFF-1 cell line.
Negative control: HT-29.
Image was taken with a confocal microscope(Leica-Microsystems, TCS SP8).

ab195889 Anti-alpha Tubulin mouse monoclonal antibody - Microtubule Marker (Alexa Fluor® 594) was used to counterstain tubulin at 1/500 (1ug/ml) dilution (Red). The Nuclear counterstain was DAPI (Blue).

Secondary antibody only control: Secondary antibody is ab150081 Goat Anti-Rabbit IgG H&L (Alexa Fluor® 488) preadsorbed at 1/1000 (2 ug/ml) dilution.



Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) - Anti-Collagen I antibody [RM1131] - BSA and Azide free (ab316223)

This data was developed using ab316222, the same antibody clone in a different buffer formulation.

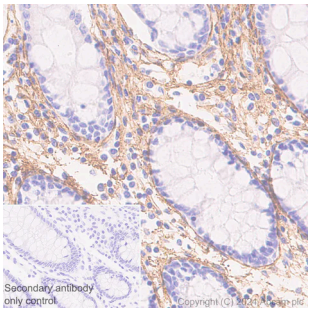
Immunohistochemical analysis of paraffin-embedded Human stomach tissue labeling Collagen I with ab316222 at 1/500 (1.012 ug/ml) dilution, followed by a ready to use LeicaDS9800 (Bond™ Polymer Refine Detection).

Positive staining on stroma of human stomach.
The section was incubated with ab316222 for 30 mins at room temperature.
The immunostaining was performed on a Leica Biosystems BOND® RX instrument

Counterstained with Hematoxylin.

Secondary antibody only control: Secondary antibody is a ready to use LeicaDS9800 (Bond™ Polymer Refine Detection).

Heat mediated antigen retrieval was performed with Tris-EDTA buffer (pH 9.0, Epitope Retrieval Solution2) for 20 mins



Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) - Anti-Collagen I antibody [RM1131] - BSA and Azide free (ab316223)

This data was developed using ab316222, the same antibody clone in a different buffer formulation.

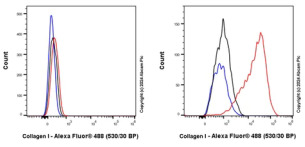
Immunohistochemical analysis of paraffin-embedded Human colon tissue labeling Collagen I with ab316222 at 1/500 (1.012 ug/ml) dilution, followed by a ready to use LeicaDS9800 (Bond™ Polymer Refine Detection).

Positive staining on stroma of human colon.
The section was incubated with ab316222 for 30 mins at room temperature.
The immunostaining was performed on a Leica Biosystems BOND® RX instrument

Counterstained with Hematoxylin.

Secondary antibody only control: Secondary antibody is a ready to use LeicaDS9800 (Bond™ Polymer Refine Detection).

Heat mediated antigen retrieval was performed with Tris-EDTA buffer (pH 9.0, Epitope Retrieval Solution2) for 20 mins



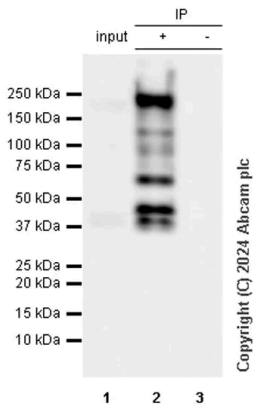
Flow Cytometry (Intracellular) - Anti-Collagen I antibody [RM1131] - BSA and Azide free (ab316223)

This data was developed using ab316222, the same antibody clone in a different buffer formulation.

Flow cytometric analysis of 4% paraformaldehyde fixed 90% methanol permeabilized HT-29 (human colorectal adenocarcinoma epithelial cell, Left) / HFF-1 (human skin fibroblast, Right) cells labelling Collagen I with ab316222 at 1/5000 dilution (0.01 ug)/Red compared with a Rabbit monoclonal IgG (ab172730) (Black) isotype control and an unlabelled control (cells without incubation with primary antibody and secondary antibody) (Blue).

Goat Anti-Rabbit IgG (Alexa Fluor® 488, ab150081) at 1/5000 dilution was used as the secondary antibody.

Negative control: HT-29.



Immunoprecipitation - Anti-Collagen I antibody [RM1131] - BSA and Azide free (ab316223)

This data was developed using ab316222, the same antibody clone in a different buffer formulation.

Collagen I was immunoprecipitated from 0.35 mg HFF-1 (human skin fibroblast) whole cell lysate with ab316222 at 1/30 dilution (2ug in 0.35mg lysates). Western blot was performed on the immunoprecipitate using ab316222 at 1/1000 dilution. VeriBlot for IP secondary antibody(HRP) (ab131366) was used at 1/5000 dilution.

Lane 1: HFF-1 (human skin fibroblast) whole cell lysate
Lane 2: ab316222 IP in HFF-1 (human skin fibroblast) whole cell lysate
Lane 3: Rabbit monoclonal IgG (ab172730) instead of ab316222 in HFF-1 whole cell lysate

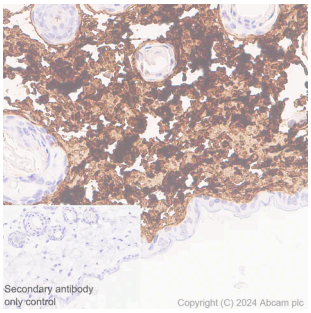
All lanes:
Immunoprecipitation - Anti-Collagen I antibody [RM1131] (ab316222) at 1/30 dilution

All lanes:
HFF-1 (human skin fibroblast) whole cell lysate

Secondary

All lanes:
Immunoprecipitation - VeriBlot for IP Detection Reagent (HRP) (ab131366) at 1/5000 dilution

Exposure time: 10s



Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) - Anti-Collagen I antibody [RM1131] - BSA and Azide free (ab316223)

This data was developed using ab316222, the same antibody clone in a different buffer formulation.

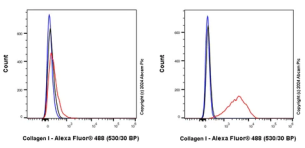
Immunohistochemical analysis of paraffin-embedded Mouse skin tissue labeling Collagen I with ab316222 at 1/500 (1.012 ug/ml) dilution, followed by a ready to use LeicaDS9800 (Bond™ Polymer Refine Detection).

Positive staining on stroma of mouse skin.
The section was incubated with ab316222 for 30 mins at room temperature.
The immunostaining was performed on a Leica Biosystems BOND® RX instrument

Counterstained with Hematoxylin.

Secondary antibody only control: Secondary antibody is a ready to use LeicaDS9800 (Bond™ Polymer Refine Detection).

Heat mediated antigen retrieval was performed with Tris-EDTA buffer (pH 9.0, Epitope Retrieval Solution2) for 20 mins



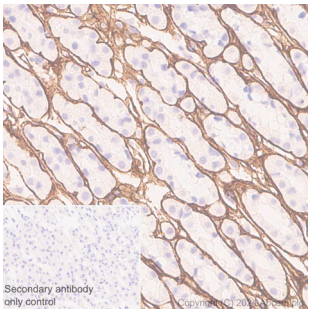
Flow Cytometry (Intracellular) - Anti-Collagen I antibody [RM1131] - BSA and Azide free (ab316223)

This data was developed using ab316222, the same antibody clone in a different buffer formulation.

Flow cytometric analysis of 4% paraformaldehyde fixed 90% methanol permeabilized C6 (rat glial tumor glial cell, Left) / L6 (rat skeletal muscle myoblast, Right) cells labelling Collagen I with ab316222 at 1/5000 dilution (0.01 ug)/Red compared with a Rabbit monoclonal IgG (ab172730) (Black) isotype control and an unlabelled control (cells without incubation with primary antibody and secondary antibody) (Blue).

Goat Anti-Rabbit IgG (Alexa Fluor® 488, ab150081) at 1/5000 dilution was used as the secondary antibody.

Negative control: C6.



Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) - Anti-Collagen I antibody [RM1131] - BSA and Azide free (ab316223)

This data was developed using ab316222, the same antibody clone in a different buffer formulation.

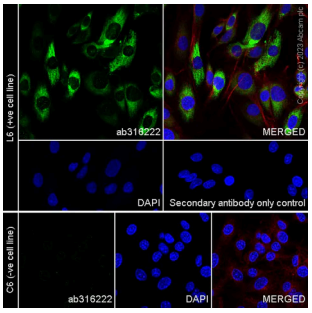
Immunohistochemical analysis of paraffin-embedded Rat stomach tissue labeling Collagen I with ab316222 at 1/500 (1.012 ug/ml) dilution, followed by a ready to use LeicaDS9800 (Bond™ Polymer Refine Detection).

Positive staining on stroma of rat stomach.
The section was incubated with ab316222 for 30 mins at room temperature.
The immunostaining was performed on a Leica Biosystems BOND® RX instrument

Counterstained with Hematoxylin.

Secondary antibody only control: Secondary antibody is a ready to use LeicaDS9800 (Bond™ Polymer Refine Detection).

Heat mediated antigen retrieval was performed with Tris-EDTA buffer (pH 9.0, Epitope Retrieval Solution2) for 20 mins



Immunocytochemistry/ Immunofluorescence - Anti-Collagen I antibody [RM1131] - BSA and Azide free (ab316223)

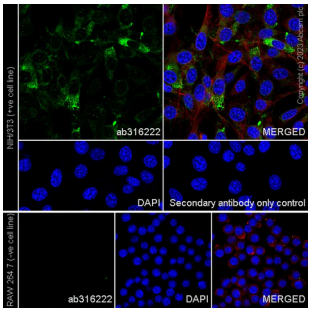
This data was developed using ab316222, the same antibody clone in a different buffer formulation.

Immunofluorescent analysis of 100% methanol-fixed, 0.1% TritonX-100 permeabilized L6 (rat skeletal muscle myoblast) cells labelling Collagen I with ab316222 at 1/100 (5.06 ug/ml) dilution, followed by ab150081 Goat Anti-Rabbit IgG H&L (Alexa Fluor® 488) preadsorbed antibody at 1/1000 (2 ug/ml) dilution (Green).

Confocal image showing cytoplasmic staining in L6 cell line.
Negative control: C6.
Image was taken with a confocal microscope(Leica-Microsystems, TCS SP8).

ab195889 Anti-alpha Tubulin mouse monoclonal antibody - Microtubule Marker (Alexa Fluor® 594) was used to counterstain tubulin at 1/500 (1ug/ml) dilution (Red). The Nuclear counterstain was DAPI (Blue).

Secondary antibody only control: Secondary antibody is ab150081 Goat Anti-Rabbit IgG H&L (Alexa Fluor® 488) preadsorbed at 1/1000 (2 ug/ml) dilution.



Immunocytochemistry/ Immunofluorescence - Anti-Collagen I antibody [RM1131] - BSA and Azide free (ab316223)

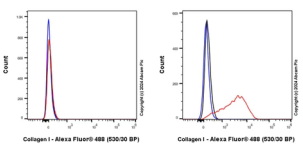
This data was developed using ab316222, the same antibody clone in a different buffer formulation.

Immunofluorescent analysis of 100% methanol-fixed, 0.1% TritonX-100 permeabilized NIH/3T3 (mouse embryonic fibroblast) cells labelling Collagen I with ab316222 at 1/100 (5.06 ug/ml) dilution, followed by ab150081 Goat Anti-Rabbit IgG H&L (Alexa Fluor® 488) preadsorbed antibody at 1/1000 (2 ug/ml) dilution (Green).

Confocal image showing cytoplasmic staining in NIH/3T3 cell line.
Negative control: RAW 264.7.
Image was taken with a confocal microscope(Leica-Microsystems, TCS SP8).

ab195889 Anti-alpha Tubulin mouse monoclonal antibody - Microtubule Marker (Alexa Fluor® 594) was used to counterstain tubulin at 1/500 (1ug/ml) dilution (Red). The Nuclear counterstain was DAPI (Blue).

Secondary antibody only control: Secondary antibody is ab150081 Goat Anti-Rabbit IgG H&L (Alexa Fluor® 488) preadsorbed at 1/1000 (2 ug/ml) dilution.



Flow Cytometry (Intracellular) - Anti-Collagen I antibody [RM1131] - BSA and Azide free (ab316223)

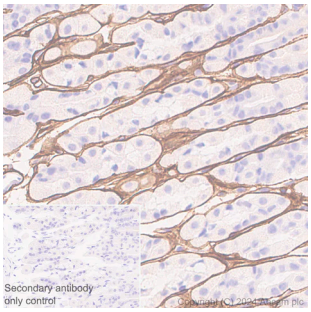
This data was developed using ab316222, the same antibody clone in a different buffer formulation.

Flow cytometric analysis of 4% paraformaldehyde fixed 90% methanol permeabilized Raw 264.7 (mouse Abelson murine leukemia virus-induced tumor macrophage, Left) / NIH/3T3 (mouse embryonic fibroblast, Right) cells labelling Collagen I with ab316222 at 1/5000 dilution (0.01 ug)/Red compared with a Rabbit monoclonal IgG (ab172730) (Black) isotype control and an

unlabelled control (cells without incubation with primary antibody and secondary antibody) (Blue).

Goat Anti-Rabbit IgG (Alexa Fluor® 488, ab150081) at 1/5000 dilution was used as the secondary antibody.

Negative control: Raw 264.7.



Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) - Anti-Collagen I antibody [RM1131] - BSA and Azide free (ab316223)

This data was developed using ab316222, the same antibody clone in a different buffer formulation.

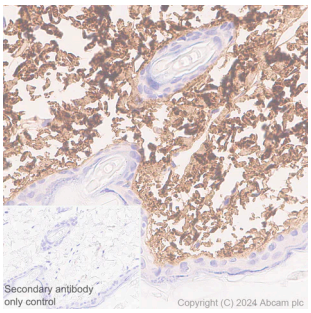
Immunohistochemical analysis of paraffin-embedded Mouse stomach tissue labeling Collagen I with ab316222 at 1/500 (1.012 ug/ml) dilution, followed by a ready to use LeicaDS9800 (Bond™ Polymer Refine Detection).

Positive staining on stroma of mouse stomach.
The section was incubated with ab316222 for 30 mins at room temperature.
The immunostaining was performed on a Leica Biosystems BOND® RX instrument

Counterstained with Hematoxylin.

Secondary antibody only control: Secondary antibody is a ready to use LeicaDS9800 (Bond™ Polymer Refine Detection).

Heat mediated antigen retrieval was performed with Tris-EDTA buffer (pH 9.0, Epitope Retrieval Solution2) for 20 mins



Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) - Anti-Collagen I antibody [RM1131] - BSA and Azide free (ab316223)

This data was developed using ab316222, the same antibody clone in a different buffer formulation.

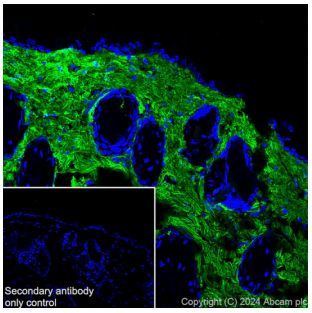
Immunohistochemical analysis of paraffin-embedded Rat skin tissue labeling Collagen I with ab316222 at 1/500 (1.012 ug/ml) dilution, followed by a ready to use LeicaDS9800 (Bond™ Polymer Refine Detection).

Positive staining on stroma of rat skin.
The section was incubated with ab316222 for 30 mins at room temperature.
The immunostaining was performed on a Leica Biosystems BOND® RX instrument

Counterstained with Hematoxylin.

Secondary antibody only control: Secondary antibody is a ready to use LeicaDS9800 (Bond™ Polymer Refine Detection).

Heat mediated antigen retrieval was performed with Tris-EDTA buffer (pH 9.0, Epitope Retrieval Solution2) for 20 mins



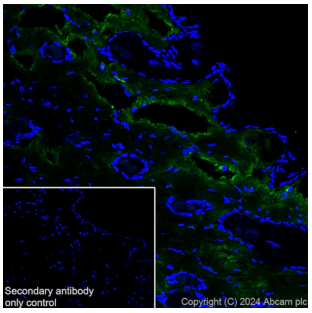
Immunohistochemistry (Frozen sections) - Anti-Collagen I antibody [RM1131] - BSA and Azide free (ab316223)

This data was developed using ab316222, the same antibody clone in a different buffer formulation.

Immunohistochemical analysis of 4% PFA-fixed, 0.2% Triton X-100 permeabilized frozen Mouse skin (fresh frozen) tissue labeling Collagen I with ab316222 at 1/500 (1.012 ug/ml) dilution followed by ab150081 Goat Anti-Rabbit IgG H&L (Alexa Fluor® 488) preadsorbed at 1/1000 (2 ug/mL) dilution (Green).

Confocal image showing positive staining on mouse skin. The nuclear counterstain was DAPI (Blue). The section was incubated with ab316222 for 60 mins at room temperature. The section was then mounted using Fluoromount®.The immunostaining was performed on a Leica Biosystems BOND® RX instrument. Image was taken with a confocal microscope (Leica-Microsystems, TCS SP8).

Secondary antibody control: Secondary antibody is ab150081 Goat Anti-Rabbit IgG H&L (Alexa Fluor® 488) preadsorbedat 1/1000 (2 ug/mL) dilution.



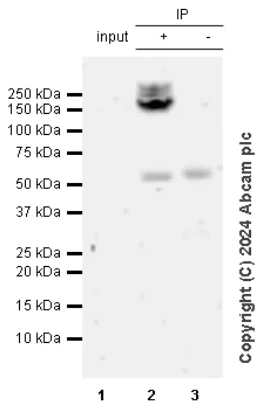
Immunohistochemistry (Frozen sections) - Anti-Collagen I antibody [RM1131] - BSA and Azide free (ab316223)

This data was developed using ab316222, the same antibody clone in a different buffer formulation.

Immunohistochemical analysis of 4% PFA-fixed, 0.2% Triton X-100 permeabilized frozen Rat skin (fresh frozen) tissue labeling Collagen I with ab316222 at 1/500 (1.012 ug/ml) dilution followed by ab150081 Goat Anti-Rabbit IgG H&L (Alexa Fluor® 488) preadsorbed at 1/1000 (2 ug/mL) dilution (Green).

Confocal image showing positive staining on rat skin. The nuclear counterstain was DAPI (Blue). The section was incubated with ab316222 for 60 mins at room temperature. The section was then mounted using Fluoromount®.The immunostaining was performed on a Leica Biosystems BOND® RX instrument. Image was taken with a confocal microscope (Leica-Microsystems, TCS SP8).

Secondary antibody control: Secondary antibody is ab150081 Goat Anti-Rabbit IgG H&L (Alexa Fluor® 488) preadsorbedat 1/1000 (2 ug/mL) dilution.



Immunoprecipitation - Anti-Collagen I antibody [RM1131] - BSA and Azide free (ab316223)

This data was developed using ab316222, the same antibody clone in a different buffer formulation.

Collagen I was immunoprecipitated from 0.35 mg Mouse skin tissue lysate with ab316222 at 1/30 dilution (2ug in 0.35mg lysates). Western blot was performed on the immunoprecipitate using ab316222 at 1/1000 dilution. VeriBlot for IP secondary antibody(HRP)(ab131366) was used at 1/5000 dilution.

Lane 1: Mouse skin tissue lysate
Lane 2: ab316222 IP in Mouse skin tissue lysate
Lane 3: Rabbit monoclonal IgG (ab172730) instead of ab316222 in mouse skin tissue lysate

All lanes:

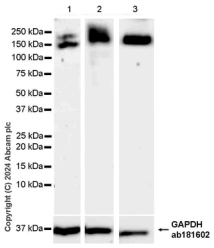
Immunoprecipitation - Anti-Collagen I antibody [RM1131] (ab316222) at 1/30 dilution

All lanes:
Mouse skin tissue lysate

Secondary

All lanes:
Immunoprecipitation - VeriBlot for IP Detection Reagent (HRP) (ab131366) at 1/5000 dilution

Exposure time: 180s



Western blot - Anti-Collagen I antibody [RM1131] - BSA and Azide free (ab316223)

Collagen I Western blot staining using rabbit Anti-Collagen I antibody

This data was developed using ab316222, the same antibody clone in a different buffer formulation.

The molecular weight observed is consistent with what has been described in the literature (PMID: 23940311)

In Western blot, Anti-GAPDH antibody [EPR16891] - Loading Control (ab181602) staining at 1/200000 dilution.

Exposure time: Lane 1: 180 seconds, Lane 2: 8 seconds, Lane 3: 81 seconds

All lanes:
Western blot - Anti-Collagen I antibody [RM1131] (ab316222) at 1/5000 dilution

Lane 1:
Mouse skin tissue lysate at 20 µg

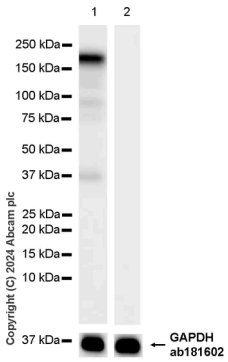
Lane 2:
Rat skin tissue lysate at 20 µg

Lane 3:
Human uterus tissue lysate at 20 µg

Secondary

All lanes:
Goat Anti-Rabbit IgG (HRP) with minimal cross-reactivity with human IgG at 1/2000 dilution

Observed band size: 139 kDa, 220 kDa, 36 kDa



Western blot - Anti-Collagen I antibody [RM1131] - BSA and Azide free (ab316223)

Collagen I Western blot staining using rabbit Anti-Collagen I antibody

This data was developed using ab316222, the same antibody clone in a different buffer formulation.

Collagen I might be easily degraded. Lysates were freshly made and used for Western blotting immediately to minimize protein degradation.

In Western blot, Anti-GAPDH antibody [EPR16891] - Loading Control (ab181602) staining at 1/200000 dilution.

All lanes:
Western blot - Anti-Collagen I antibody [RM1131] (ab316222) at 1/1000 dilution

Lane 1:
HFF-1 (human skin fibroblast) whole cell lysate at 20 µg

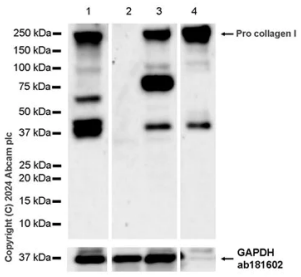
Lane 2:
HT-29 (human colorectal adenocarcinoma epithelial cell) whole cell lysate at 20 µg

Secondary

All lanes:
Goat Anti-Rabbit IgG (HRP) with minimal cross-reactivity with human IgG at 1/2000 dilution

Observed band size: 220 kDa, 36 kDa

Exposure time: 6s



Western blot - Anti-Collagen I antibody [RM1131] - BSA and Azide free (ab316223)

This data was developed using ab316222, the same antibody clone in a different buffer formulation.

Blocking and diluting buffer and concentration: 5% NFDM/TBST.
Negative control: HT-29.
We are unsure how to define these extra bands below 100kDa.

All lanes:
Western blot - Anti-Collagen I antibody [RM1131] (ab316222) at 1/1000 dilution

Lane 1:
HFF-1 (human skin fibroblast) whole cell lysate at 20 µg

Lane 2:
HT-29 (human colorectal adenocarcinoma epithelial cell) whole cell lysate at 20 µg

Lane 3:
NIH/3T3 (mouse embryonic fibroblast) whole cell lysate at 20 µg

Lane 4:
NIH/3T3 (mouse embryonic fibroblast) culture supernatant at 10 µL

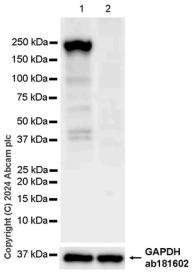
Secondary

All lanes:

Goat Anti-Rabbit IgG (HRP) with minimal cross-reactivity with human IgG at 1/2000 dilution

Observed band size: 220 kDa

Exposure time: 180s



Western blot - Anti-Collagen I antibody [RM1131] - BSA and Azide free (ab316223)

Collagen I Western blot staining using rabbit Anti-Collagen I antibody

This data was developed using ab316222, the same antibody clone in a different buffer formulation.

Negative control: C6.

In Western blot, Anti-GAPDH antibody [EPR16891] - Loading Control (ab181602) staining at 1/200000 dilution.

All lanes:
Western blot - Anti-Collagen I antibody [RM1131] (ab316222) at 1/1000 dilution

Lane 1:
L6 (rat skeletal muscle myoblast) whole cell lysate at 20 µg

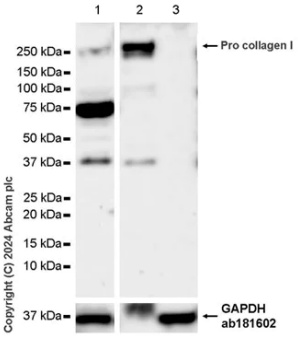
Lane 2:
C6 (rat glial tumor glial cell) whole cell lysate at 20 µg

Secondary

All lanes:
Goat Anti-Rabbit IgG (HRP) with minimal cross-reactivity with human IgG at 1/2000 dilution

Observed band size: 220 kDa, 36 kDa

Exposure time: 10s



Western blot - Anti-Collagen I antibody [RM1131] - BSA and Azide free (ab316223)

This data was developed using ab316222, the same antibody clone in a different buffer formulation.

Blocking and diluting buffer and concentration: 5% NFDM/TBST.
Negative control: RAW 264.7.
We are unsure how to define these extra bands below 100kDa.

All lanes:
Western blot - Anti-Collagen I antibody [RM1131] (ab316222) at 1/5000 dilution

Lane 1:
NIH/3T3 (mouse embryonic fibroblast) whole cell lysate at 20 µg

Lane 2:
NIH/3T3 (mouse embryonic fibroblast) culture supernatant at 10 µL

Lane 3:

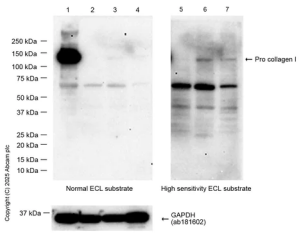
RAW 264.7 (mouse Abelson murine leukemia virus-induced tumor macrophage) whole cell lysate at 20 µg

Secondary

All lanes:
Goat Anti-Rabbit IgG (HRP) with minimal cross-reactivity with human IgG at 1/2000 dilution

Observed band size: 220 kDa

Exposure time: 81s



Western blot - Anti-Collagen I antibody [RM1131] - BSA and Azide free (ab316223)

This data was developed using ab316222, the same antibody clone in a different buffer formulation.

Blocking and diluting buffer and concentration: 5% NFDM/TBST.

This blot was developed using a high sensitivity ECL substrate.

Low expression tissue: normal mouse kidney (PMID: 33758176, PMID: 29102563)

All lanes:
Western blot - Anti-Collagen I antibody [RM1131] (ab316222) at 1/1000 dilution

Lane 1:
Mouse skin tissue lysate at 20 µg

Lanes 2 and 5:
Mouse kidney tissue lysate at 20 µg

Lanes 3 and 6:
Mouse lung tissue lysate at 20 µg

Lanes 4 and 8:
Mouse heart tissue lysate at 20 µg

Secondary

All lanes:
Western blot - Goat Anti-Rabbit IgG H&L (HRP) (ab97051) at 1/20000 dilution

Predicted band size: 139 kDa

Observed band size: 139 kDa

Exposure time: 180s

Please note: All products are 'FOR RESEARCH USE ONLY. NOT FOR USE IN DIAGNOSTIC OR THERAPEUTIC PROCEDURES'.