**Product datasheet**

**Anti-alpha smooth muscle Actin antibody ab5694**

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<td><strong>Purity</strong></td>
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Clonality: Polyclonal
Isotype: IgG

Applications

Our Abpromise guarantee covers the use of ab5694 in the following tested applications. The application notes include recommended starting dilutions; optimal dilutions/concentrations should be determined by the end user.

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<td>IHC-FoFr</td>
<td>✔️ ✔️ ✔️ ✔️</td>
<td>Use at an assay dependent concentration.</td>
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<tr>
<td>ICC</td>
<td>✔️ ✔️ ✔️ ✔️</td>
<td>Use at an assay dependent concentration.</td>
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<tr>
<td>ICC/IF</td>
<td>✔️ ✔️ ✔️ ✔️</td>
<td>1/100.</td>
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<tr>
<td>WB</td>
<td>✔️ ✔️ ✔️ ✔️</td>
<td>Use a concentration of 0.5 - 2 µg/ml. Predicted molecular weight: 42 kDa.</td>
</tr>
<tr>
<td>ELISA</td>
<td>✔️ ✔️ ✔️ ✔️</td>
<td>Use a concentration of 0.1 - 1 µg/ml.</td>
</tr>
<tr>
<td>IHC-P</td>
<td>✔️ ✔️ ✔️ ✔️</td>
<td>1/50 - 1/200. Perform heat mediated antigen retrieval before commencing with IHC staining protocol.</td>
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Target

Function: Actins are highly conserved proteins that are involved in various types of cell motility and are ubiquitously expressed in all eukaryotic cells.

Involvement in disease: Defects in ACTA2 are the cause of aortic aneurysm familial thoracic type 6 (AAT6) [MIM:611788]. AATs are characterized by permanent dilation of the thoracic aorta usually due to degenerative changes in the aortic wall. They are primarily associated with a characteristic histologic appearance known as 'medial necrosis' or 'Erdheim cystic medial necrosis' in which there is degeneration and fragmentation of elastic fibers, loss of smooth muscle cells, and an accumulation of basophilic ground substance.

Sequence similarities: Belongs to the actin family.

Cellular localization: Cytoplasm > cytoskeleton.

Images
ab5694 at 1/500 staining rat myofibroblast cells by Immunocytochemistry/Immunofluorescence. The cells were formaldehyde fixed and blocked with 5% serum prior to incubation with the antibody for 2 hours. A FITC conjugated goat anti-rabbit IgG was used as the secondary. Nuclei were counterstained with propidium iodide.

Lanes 1-3: Anti-alpha smooth muscle Actin antibody (ab5694) at 1 µg/ml

Lane 1: HEK293 cell lysate - overexpressing alpha-Actin
Lanes 2 & 4: 3T3 cell lysate
Lanes 3 & 5: Mouse heart tissue homogenate

Lysates/proteins at 20 µg per lane.

Secondary
All lanes: Fluor 750-conjugated goat anti-rabbit IgG (H+L) at 1/12500 dilution

Predicted band size: 42 kDa
Observed band size: 42 kDa

Incubated with the primary antibody at 4°C overnight.

Incubated with the secondary antibody at room temperature for 1 hour.
ab5694 staining Human fetal heart cells by ICC/IF. Cells were fixed with PFA and permeabilized in 0.1% saponin prior to blocking in 10% serum for 45 minutes at 37°C. The primary antibody was diluted 1/400 and incubated with the sample for 1 hour at 37°C. A Alexa Fluor® 594 conjugated goat polyclonal to rabbit IgG (H+L), diluted 1/600 was used as secondary antibody.

This picture shows formalin-fixed, paraffin embedded mouse intestine and mesentery, the optimal dilution is 1:1600 to 1:3200, incubation overnight at 4°C, counterstained with Hematoxylin.

This image was kindly supplied as part of the review by JQ Zhang.
Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) analysis of human tonsil tissue labelling alpha smooth muscle Actin with ab5694 at a dilution of 1/1000. Heat mediated antigen retrieval was performed for 35 minutes followed by cooling for 20 minutes. Sections were incubated with the primary antibody for 1 hour followed by incubation with a biotinylated secondary antibody for 30 minutes then HRP-Streptavidin for 30 minutes. Developed using DAB chromogen substrate (5-10 minutes). Counter stained with hematoxylin.

Magnification: left - 10X, right - 40X.

Immunohistochemistry (Formalin-fixed paraffin-embedded sections) analysis of skeletal muscle tissue (left) incubated with ab5694 at 1/100 at room temperature for 1 hour showing no specific staining. Right - human tonsil tissue secondary only control.

Heat mediated antigen retrieval was performed for 35 minutes followed by cooling for 20 minutes. A biotinylated secondary antibody was used for 30 minutes followed by incubation with HRP-Streptavidin for 30 minutes. Developed using DAB chromogen substrate (5-10 minutes). Counter stained with hematoxylin. Magnification 10X.

Human Leiomyoma stained with ab5694.
All lanes: Anti-alpha smooth muscle Actin antibody (ab5694) at 1 µg/ml

Lane 1: HeLa Nuclear
Lane 2: HeLa whole cell
Lane 3: A431 cell lysate
Lane 4: Jurkat cell lysate
Lane 5: HEK293 cell lysate

Lysates/proteins at 20 µg per lane.

Secondary
All lanes: Alexa Fluor anti-rabbit at 1/5000 dilution

Performed under reducing conditions.

Predicted band size: 42 kDa
Observed band size: 42 kDa
Additional bands at: 30 kDa, 35 kDa, 37 kDa, 50 kDa, 75 kDa. We are unsure as to the identity of these extra bands.

Please note that ab5694 does not appear to be specific to smooth muscle.
Ab5694 positively staining smooth muscle cells in blood vessels and myoepithelial cells in the frozen tissue of cancerous human mammary gland (pink) at 1/100 dilution. Secondary: CY5 conjugated goat anti rabbit (1/100). Co immunostaining of glandular cell cytokeratin can be seen stained by FITC (green). Auto fluorescent erythrocytes that are present within blood vessels are shown (red), whilst the DAPI counter stain may clearly be seen staining nuclei (blue).

This image is courtesy of an Abreview submitted by on 22 August 2005. We do not have any further information relating to this image.
Western blot - Anti-alpha smooth muscle Actin antibody (ab5694)

**All lanes**: Anti-alpha smooth muscle Actin antibody (ab5694) at 1/500 dilution

**Lane 1**: Rat2 myofibroblasts (untreated before treatment-0 days)
**Lane 2**: Rat2 myofibroblasts (untreated for 5 days)
**Lane 3**: Rat2 myofibroblasts (treated with 1ng/mL TGF beta)
**Lane 4**: Rat2 myofibroblasts (treated with 10ng/mL TGF beta)
**Lane 5**: Positive control (NIH3T3)
**Lane 6**: Negative control (MDA-MB-469 breast carcinoma cells)

Lysates/proteins at 10 µg per lane.

**Secondary**

**All lanes**: Donkey anti rabbit (HRP) at 1/2500 dilution

Performed under reducing conditions.

**Predicted band size**: 42 kDa

This image is an edited version of an image submitted courtesy of an Abreview on **20 September 2005**. We do not have any further information relating to this image.
Western blot - Anti-alpha smooth muscle Actin antibody (ab5694)

All lanes: Anti-alpha smooth muscle Actin antibody (ab5694) at 1/1000 dilution

Lanes 1-2: Lystates prepared from pig heart tissue from normal control animals
Lanes 3-4: Lystates prepared from pig heart tissue from experimental animals

Lysates/proteins at 4 µg per lane.

Secondary

All lanes: HRP-conjugated goat polyclonal to rabbit IgG at 1/20000 dilution

Performed under reducing conditions.

Predicted band size: 42 kDa
Observed band size: 45 kDa

Exposure time: 1 minute

Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) - Anti-alpha smooth muscle Actin antibody (ab5694)

ab5694 staining alpha smooth muscle Actin in human skin tissue section by Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections). Tissue underwent formaldehyde fixation before heat mediated antigen retrieval in Citrate pH 6.0 and then blocked with 10% serum for 1 hour at RT. The primary antibody was diluted 1/300 and incubated with sample in 2% serum for 15 hours at 4°C. A Biotin conjugated goat polyclonal to rabbit IgG was used at dilution at 1/500 as secondary antibody.
ab5694 staining alpha smooth muscle Actin in rat lung tissue by Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections).

Tissue was fixed in formaldehyde and a heat mediated antigen retrieval step was performed using citrate buffer pH 6.0. Samples were then permeabilized using 0.5 Triton X-100 for 20 minutes, blocked with 1% BSA for 30 minutes at 20°C and then incubated with ab5694 at a 1/100 dilution for 16 hours at 4°C. The secondary used was an Alexa-Fluor 568 conjugated goat anti-rabbit polyclonal used at a 1/250 dilution.

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