Product datasheet

Anti-Cytokeratin 8 antibody ab59400

Rating: ★★★★★ 7 Abreviews  29 References  8 Images

Overview

Product name: Anti-Cytokeratin 8 antibody
Description: Rabbit polyclonal to Cytokeratin 8
Host species: Rabbit
Tested applications:
Suitable for: ICC/IF, IHC-Fr, WB, IHC-P
Species reactivity:
Reacts with: Mouse, Rat, Human
Immunogen:
Synthetic non-phosphopeptide derived from human Cytokeratin 8 around the phosphorylation site of serine 431 (L-T-S^P-P-G).

Properties

Form: Liquid
Storage instructions:
Shipped at 4°C. Store at -20°C. Stable for 12 months at -20°C.
Storage buffer:
pH: 7.40
Preservative: 0.02% Sodium azide
 Constituents: PBS, 50% Glycerol, 0.87% Sodium chloride
Purity:
Immunogen affinity purified
Clonality:
Polyclonal
Isotype:
IgG

Applications

Our Abpromise guarantee covers the use of ab59400 in the following tested applications.
The application notes include recommended starting dilutions; optimal dilutions/concentrations should be determined by the end user.

<table>
<thead>
<tr>
<th>Application</th>
<th>Abreviews</th>
<th>Notes</th>
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</thead>
<tbody>
<tr>
<td>ICC/IF</td>
<td></td>
<td>Use a concentration of 1 µg/ml.</td>
</tr>
<tr>
<td>IHC-Fr</td>
<td>★★★★★</td>
<td>Use at an assay dependent concentration.</td>
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<tr>
<td>WB</td>
<td>★★★★★★</td>
<td>1/500 - 1/1000. Detects a band of approximately 54 kDa (predicted molecular weight: 54 kDa).</td>
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Function
Together with KRT19, helps to link the contractile apparatus to dystrophin at the costameres of striated muscle.

Tissue specificity
Observed in muscle fibers accumulating in the costameres of myoplasm at the sarcolemma membrane in structures that contain dystrophin and spectrin. Expressed in gingival mucosa and hard palate of the oral cavity.

Involvement in disease
Cirrhosis

Sequence similarities
Belongs to the intermediate filament family.

Post-translational modifications
Phosphorylation on serine residues is enhanced during EGF stimulation and mitosis. Ser-74 phosphorylation plays an important role in keratin filament reorganization. O-glycosylated. O-GlcNAcylation at multiple sites increases solubility, and decreases stability by inducing proteasomal degradation. O-glycosylated (O-GlcNAcylated), in a cell cycle-dependent manner.

Cellular localization

Images
ab59400 staining Cytokeratin 8 in c57b6 mouse prostate tissue by Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections). Tissue was fixed in formaldehyde and a heat mediated antigen retrieval step was performed using citric acid. Samples were then blocked using 3% BSA for 1 hour at 25°C and then incubated with ab59400 at a 1/100 dilution for 18 hours at 4°C. The secondary used was a biotin conjugated goat anti-rabbit polyclonal used at a 1/200 dilution.
Immunochemistry/Immunofluorescence - Anti-Cytokeratin 8 antibody (ab59400)

ICC/IF image of ab59400 stained MCF7 cells. The cells were 100% methanol fixed (5 min) and then incubated in 1%BSA / 10% normal goat serum / 0.3M glycine in 0.1% PBS-Tween for 1h to permeabilise the cells and block non-specific protein-protein interactions. The cells were then incubated with the antibody (ab59400, 1µg/ml) overnight at +4°C. The secondary antibody (green) was Alexa Fluor® 488 goat anti-rabbit IgG (H+L) used at a 1/1000 dilution for 1h. Alexa Fluor® 594 WGA was used to label plasma membranes (red) at a 1/200 dilution for 1h. DAPI was used to stain the cell nuclei (blue) at a concentration of 1.43µM.

Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) - Anti-Cytokeratin 8 antibody (ab59400)

This image is courtesy of an Abreview provided by Megha Rajaram.

ab59400 staining Cytokeratin 8 in Human breast cancer cells xenografted into nude mice by Immunohistochemistry (Formalin/ PFA-fixed paraffin-embedded tissue sections). The sections were fixed in formalin and subjected to heat-mediated antigen retrieval in citrate buffer (0.1M Sodium Citrate) prior to blocking with 5% serum for 1 hour at 4°C. The primary antibody was diluted 1/200 in 5% goat serum in PBS and incubated with the sample for 12 hours at 4°C. A Biotin-conjugated Goat anti-Rabbit polyclonal was used as the secondary antibody, diluted 1/1000.

Immunohistochemistry (Frozen sections) - Anti-Cytokeratin 8 antibody (ab59400)

Image courtesy of an anonymous Abreview.

ab59400 staining Cytokeratin 8 in murine mammary gland duct tissue by Immunohistochemistry (Frozen sections). Tissue was fixed in paraformaldehyde, permeabilized using 0.1% Triton X-100, blocked with 1% BSA for 1 hour at 23°C and then incubated with ab59400 at a 1/250 dilution for 12 hours at 4°C. The secondary used was an Alexa-Fluor 568 conjugated goat anti-rabbit polyclonal used at a 1/1000 dilution.
Immunohistochemical analysis of paraffin embedded human colon carcinoma tissue using ab59400 at 1/50 dilution, in the presence (right) and absence (left) of immunising peptide. Then, a polymer secondary antibody was used for detection.

**Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) - Anti-Cytokeratin 8 antibody (ab59400)**

**Western blot - Anti-Cytokeratin 8 antibody (ab59400)**

**Immunocytochemistry/ Immunofluorescence - Anti-Cytokeratin 8 antibody (ab59400)**

**All lanes**: Anti-Cytokeratin 8 antibody (ab59400) at 1/500 dilution

**Lane 1**: EGF treated (200ng/ml, 30mins) 293 cell extracts

**Lane 2**: EGF treated (200ng/ml, 30mins) 293 cell extracts with immunising peptide

**Predicted band size**: 54 kDa

**Observed band size**: 54 kDa

Immunocytochemistry/ Immunofluorescence analysis of HeLa cells, labeling Cytokeratin 8 with ab59400. The picture on the right is blocked with the synthesized peptide.
Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) analysis of human kidney tissue sections labeling Cytokeratin 8 with ab59400 at 1/100 dilution. Tissue sections were fixed with formaldehyde; heat mediated antigen retrieval was performed using a citric acid. 2% BSA was used to block, followed by incubation with ab59400 in TBS/BSA/azide for 2 hours at 21°C. A polyclonal goat anti-rabbit IgG H&L (hrP) conjugated secondary antibody was used at 1/300 dilution.

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