Cytokeratin 19 (KRT19) Rabbit mAb

Catalog No.: A25546 Recombinant



Basic Information

Observed MW

Refer to figures

Calculated MW

44kDa

Category

Primary antibody

Applications

IF/ICC,IF-P,IHC-P,FC (intra),ELISA

Cross-Reactivity

Human, Mouse, Rat

CloneNo number

ARC54260

Background

The protein encoded by this gene is a member of the keratin family. The keratins are intermediate filament proteins responsible for the structural integrity of epithelial cells and are subdivided into cytokeratins and hair keratins. The type I cytokeratins consist of acidic proteins which are arranged in pairs of heterotypic keratin chains. Unlike its related family members, this smallest known acidic cytokeratin is not paired with a basic cytokeratin in epithelial cells. It is specifically expressed in the periderm, the transiently superficial layer that envelopes the developing epidermis. The type I cytokeratins are clustered in a region of chromosome 17q12-q21.

Recommended Dilutions

IF/ICC 1:200 - 1:800

IF-P 1:200 - 1:800

IHC-P 1:5000 - 1:20000

FC (intra) 1:1000 - 1:5000

ELISA Recommended starting

concentration is 1 µg/mL.

Please optimize the
concentration based on
your specific assay
requirements.

Immunogen Information

Gene ID3880

Swiss Prot
P08727

Immunogen

Synthetic peptide. This information is considered to be commercially sensitive.

Synonyms

K19; CK19; K1CS

Contact

a	400-999-6126
×	cn.market@abclonal.com.cn
<u> </u>	www.abclonal.com.cn

Product Information

SourceIsotypePurificationRabbitIgGAffinity purification

Storage

Store at -20°C. Avoid freeze / thaw cycles.

Buffer: PBS containing 50% glycerol and 0.05% BSA, preserved with proclin300 or sodium azide (as specified on the Certificate of Analysis), pH 7.3.



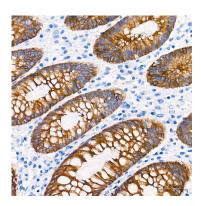




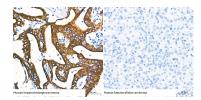


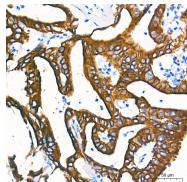
Flow cytometry: 1X10^6 Jurkat cells (negative control,left) and MCF7 cells (right) were intracellularly-stained with Cytokeratin 19 (KRT19) Rabbit mAb (A25546,2 μg/mL,orange line) or Rabbit IgG isotype control (AC042,2 µg/mL,blue line), followed by FITC conjugated goat anti-Rabbit pAb staining. Non-fluorescently stained cells were used as blank control (red line).

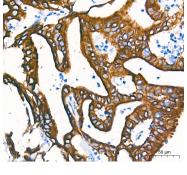
Flow cytometry: 1X10^6 MCF7 cells were intracellularly-stained with Rabbit IgG isotype control (AC042,2 µg/mL,left) or Cytokeratin 19 (KRT19) Rabbit mAb (A25546,2 μg/mL,right), followed by FITC conjugated goat anti-Rabbit pAb staining.

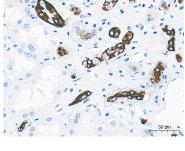


Immunohistochemistry analysis of paraffinembedded Human colon tissue using Cytokeratin 19 (KRT19) Rabbit mAb (A25546) at a dilution of 1:10000 (40x lens). High pressure antigen retrieval performed with 0.01M Tris-EDTA Buffer (pH 9.0) prior to IHC staining.





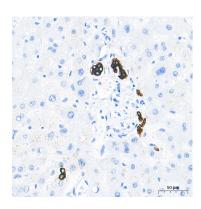




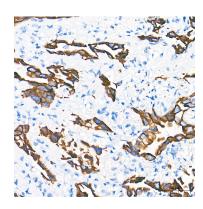
Immunohistochemistry analysis of paraffinembedded Human hepatocholangiocarcinoma and hepatocellular carcinoma tissue using Cytokeratin 19 (KRT19) Rabbit mAb (A25546) at a dilution of 1:10000 (40x lens). High pressure antigen retrieval performed with 0.01M Tris-EDTA Buffer (pH 9.0) prior to IHC staining.

Immunohistochemistry analysis of paraffinembedded Human hepatocholangiocarcinoma tissue using Cytokeratin 19 (KRT19) Rabbit mAb (A25546) at a dilution of 1:10000 (40x lens). High pressure antigen retrieval performed with 0.01M Tris-EDTA Buffer (pH 9.0) prior to IHC staining.

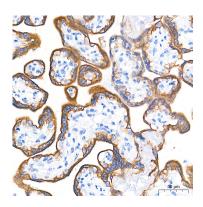
Immunohistochemistry analysis of paraffinembedded Human kidney tissue using Cytokeratin 19 (KRT19) Rabbit mAb (A25546) at a dilution of 1:10000 (40x lens). High pressure antigen retrieval performed with 0.01M Tris-EDTA Buffer (pH 9.0) prior to IHC staining.



Immunohistochemistry analysis of paraffinembedded Human liver tissue using Cytokeratin 19 (KRT19) Rabbit mAb (A25546) at a dilution of 1:10000 (40x lens). High pressure antigen retrieval performed with 0.01M Tris-EDTA Buffer (pH 9.0) prior to

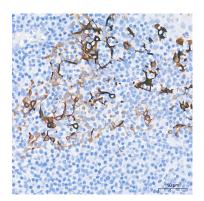


Immunohistochemistry analysis of paraffinembedded Human lung cancer tissue using Cytokeratin 19 (KRT19) Rabbit mAb (A25546) at a dilution of 1:10000 (40x lens). High pressure antigen retrieval performed with 0.01M Tris-EDTA Buffer (pH 9.0) prior to

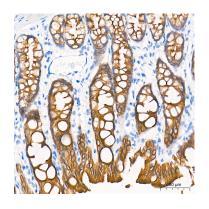


Immunohistochemistry analysis of paraffinembedded Human placenta tissue using Cytokeratin 19 (KRT19) Rabbit mAb (A25546) at a dilution of 1:10000 (40x lens). High pressure antigen retrieval performed with 0.01M Tris-EDTA Buffer (pH 9.0) prior to

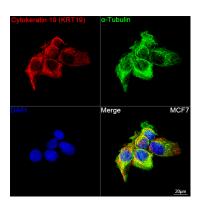
IHC staining.



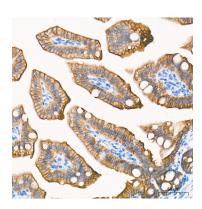
Immunohistochemistry analysis of paraffinembedded Human tonsil tissue using Cytokeratin 19 (KRT19) Rabbit mAb (A25546) at a dilution of 1:10000 (40x lens). High pressure antigen retrieval performed with 0.01M Tris-EDTA Buffer (pH 9.0) prior to IHC staining.



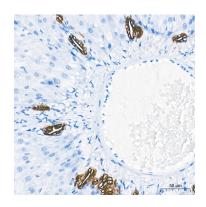
Immunohistochemistry analysis of paraffinembedded Rat colon tissue using Cytokeratin 19 (KRT19) Rabbit mAb (A25546) at a dilution of 1:10000 (40x lens). High pressure antigen retrieval performed with 0.01M Tris-EDTA Buffer (pH 9.0) prior to IHC staining.



Confocal imaging of MCF7 cells using Cytokeratin 19 (KRT19) Rabbit mAb (A25546, dilution 1:500) followed by a further incubation with Cy3 Goat Anti-Rabbit IgG (H+L) (AS007, dilution 1:500) (Red). The IHC staining.

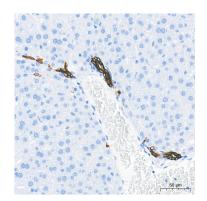


Immunohistochemistry analysis of paraffinembedded Mouse intestin tissue using Cytokeratin 19 (KRT19) Rabbit mAb (A25546) at a dilution of 1:10000 (40x lens). High pressure antigen retrieval performed with 0.01M Tris-EDTA Buffer (pH 9.0) prior to IHC staining.

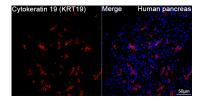


Immunohistochemistry analysis of paraffinembedded Rat liver tissue using Cytokeratin 19 (KRT19) Rabbit mAb (A25546) at a dilution of 1:10000 (40x lens). High pressure antigen retrieval performed with 0.01M Tris-EDTA Buffer (pH 9.0) prior to IHC staining.

IHC staining.



Immunohistochemistry analysis of paraffinembedded Mouse liver tissue using Cytokeratin 19 (KRT19) Rabbit mAb (A25546) at a dilution of 1:10000 (40x lens). High pressure antigen retrieval performed with 0.01M Tris-EDTA Buffer (pH 9.0) prior to IHC staining.



Confocal imaging of paraffin-embedded Human pancreas tissue using Cytokeratin 19 (KRT19) Rabbit mAb (A25546, dilution 1:500) followed by a further incubation with Cy3 Goat Anti-Rabbit IgG (H+L) (AS007, dilution 1:500) (Red). DAPI was used for nuclear staining (Blue). High pressure antigen retrieval performed with 0.01M Citrate Buffer (pH 6.0) prior to IF staining. Objective: 40x.

Validation Data

cells were counterstained with α -Tubulin Mouse mAb (AC012, dilution 1:400) followed by incubation with ABflo® 488-conjugated Goat Anti-Mouse IgG (H+L) Ab (AS076, dilution 1:500) (Green). DAPI was used for nuclear staining (Blue). Objective: 100x.