E-Cadherin Rabbit mAb

Catalog No.: A20798 Recombinant 80 Publications



Basic Information

Observed MW 125kDa/135kDa

Calculated MW 97kDa

Category Primary antibody

Applications WB,IHC-P,IF/ICC,ELISA

Cross-Reactivity Human, Mouse, Rat

CloneNo number ARC51012

Background

This gene encodes a classical cadherin of the cadherin superfamily. Alternative splicing results in multiple transcript variants, at least one of which encodes a preproprotein that is proteolytically processed to generate the mature glycoprotein. This calcium-dependent cellcell adhesion protein is comprised of five extracellular cadherin repeats, a transmembrane region and a highly conserved cytoplasmic tail. Mutations in this gene are correlated with gastric, breast, colorectal, thyroid and ovarian cancer. Loss of function of this gene is thought to contribute to cancer progression by increasing proliferation, invasion, and/or metastasis. The ectodomain of this protein mediates bacterial adhesion to mammalian cells and the cytoplasmic domain is required for internalization. This gene is present in a gene cluster with other members of the cadherin family on chromosome 16.

Recommended Dilutions

WB	1:1000 - 1:2000
IHC-P	1:400 - 1:2000
IF/ICC	1:100 - 1:1000
ELISA	Recommended starting concentration is 1 µg/mL. Please optimize the concentration based on your specific assay requirements.

Immunogen Information

Gene ID 999

Swiss Prot P12830

Immunogen

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

Synonyms

UVO; CDHE; ECAD; LCAM; Arc-1; BCDS1; CD324; E-Cadherin

Contact

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Product Information

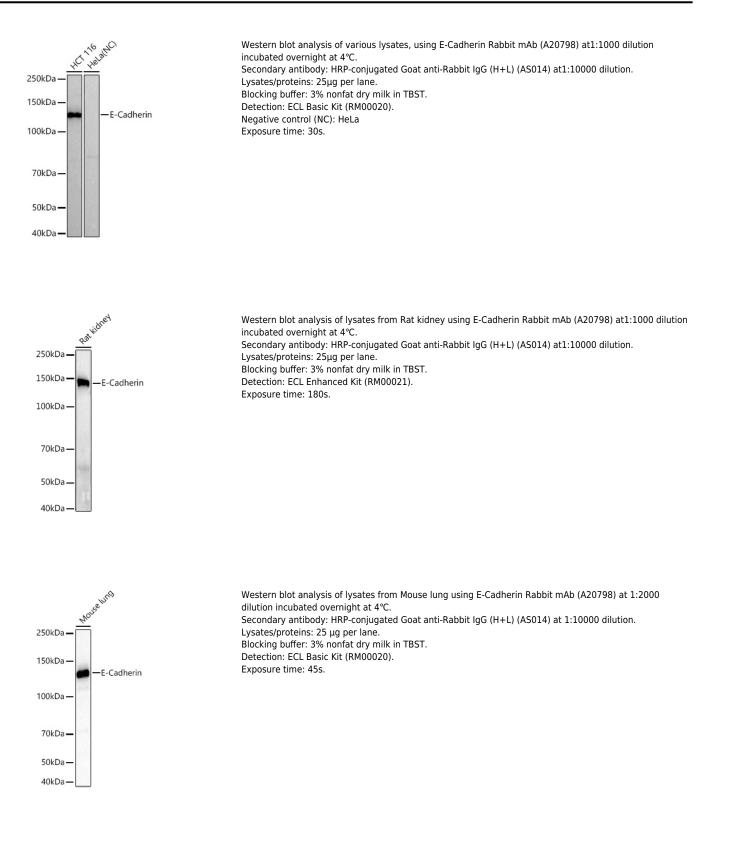
Source Rabbit

Isotype lgG

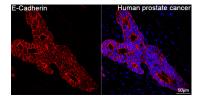
Purification Affinity purification

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.



Validation Data



E-Cadherin Merge Mouse pancreas

Confocal imaging of paraffin-embedded

Mouse pancreas tissue using E-Cadherin

followed by a further incubation with Cy3

1:500) (Red). DAPI was used for nuclear

staining (Blue). High pressure antigen

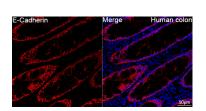
Goat Anti-Rabbit IgG (H+L) (AS007, dilution

retrieval performed with 0.01M Citrate Buffer

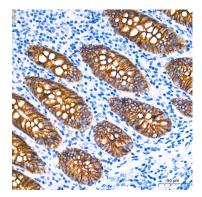
(pH 6.0) prior to IF staining. Objective: 40x.

Rabbit mAb (A20798, dilution 1:100)

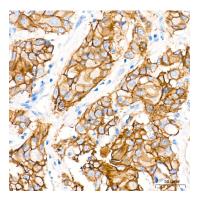
Confocal imaging of paraffin-embedded Human prostate cancer tissue using E-Cadherin Rabbit mAb (A20798, dilution 1:100) followed by a further incubation with Cy3 Goat Anti-Rabbit IgG (H+L) (AS007, dilution 1:500) (Red). DAPI was used for nuclear staining (Blue). High pressure antigen retrieval performed with 0.01M Citrate Buffer (pH 6.0) prior to IF staining. Objective: 40x.



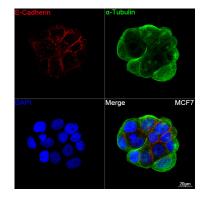
Confocal imaging of paraffin-embedded Humanc colon tissue using E-Cadherin Rabbit mAb (A20798, dilution 1:200) followed by a further incubation with Cy3 Goat Anti-Rabbit IgG (H+L) (AS007, dilution 1:500) (Red). DAPI was used for nuclear staining (Blue). High pressure antigen retrieval performed with 0.01M Citrate Buffer (pH 6.0) prior to IF staining. Objective: 40x.



Immunohistochemistry analysis of paraffinembedded Human colon tissue using E-Cadherin Rabbit mAb (A20798) at a dilution of 1:500 (40x lens). High pressure antigen



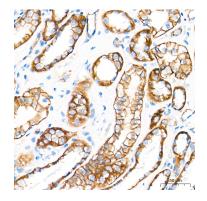
Immunohistochemistry analysis of paraffinembedded Human breast cancer tissue using E-Cadherin Rabbit mAb (A20798) at a dilution of 1:500 (40x lens). High pressure antigen retrieval performed with 0.01M Tris-EDTA Buffer(pH 9.0) prior to IHC staining.



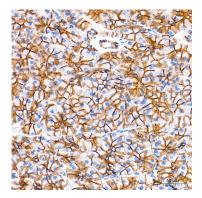
Confocal imaging of MCF7 cells using E-Cadherin Rabbit mAb (A20798, dilution 1:200) followed by a further incubation with Cy3 Goat Anti-Rabbit IgG (H+L) (AS007, dilution 1:500) (Red). The cells were counterstained with α -Tubulin Mouse mAb (AC012, dilution 1:400) followed by incubation with ABflo® 488-conjugated Goat Anti-Mouse IgG (H+L) Ab (AS076, dilution 1:500) (Green). DAPI was used for nuclear staining (Blue). Objective: 100x.



Immunohistochemistry analysis of paraffinembedded Human colon carcinoma tissue using E-Cadherin Rabbit mAb (A20798) at a dilution of 1:500 (40x lens). High pressure antigen retrieval performed with 0.01M Tris-EDTA Buffer(pH 9.0) prior to IHC staining.

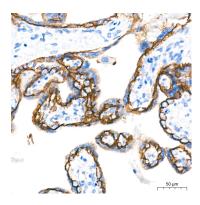


Immunohistochemistry analysis of paraffinembedded Human kidney tissue using E-Cadherin Rabbit mAb (A20798) at a dilution of 1:500 (40x lens). High pressure antigen

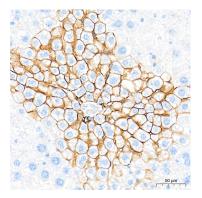


Immunohistochemistry analysis of paraffinembedded Human pancreas tissue using E-Cadherin Rabbit mAb (A20798) at a dilution of 1:500 (40x lens). High pressure antigen

retrieval performed with 0.01M Tris-EDTA Buffer(pH 9.0) prior to IHC staining.



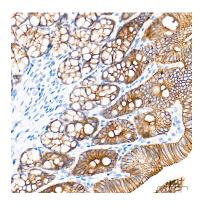
Immunohistochemistry analysis of paraffinembedded Human placenta tissue using E-Cadherin Rabbit mAb (A20798) at a dilution of 1:500 (40x lens). High pressure antigen retrieval performed with 0.01M Tris-EDTA Buffer(pH 9.0) prior to IHC staining.



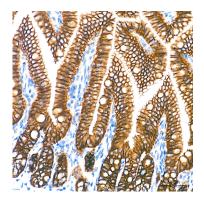
Immunohistochemistry analysis of paraffinembedded Mouse liver tissue using E-Cadherin Rabbit mAb (A20798) at a dilution of 1:500 (40x lens). High pressure antigen retrieval performed with 0.01M Tris-EDTA Buffer(pH 9.0) prior to IHC staining. retrieval performed with 0.01M Tris-EDTA Buffer(pH 9.0) prior to IHC staining.



Immunohistochemistry analysis of paraffinembedded Human thyroid cancer tissue using E-Cadherin Rabbit mAb (A20798) at a dilution of 1:500 (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 E-Cadherin Rabbit mAb (A20798) at a dilution of 1:500 (40x lens). High pressure antigen retrieval performed with 0.01M Tris-EDTA Buffer(pH 9.0) prior to IHC staining. retrieval performed with 0.01M Tris-EDTA Buffer(pH 9.0) prior to IHC staining.



Immunohistochemistry analysis of paraffinembedded Mouse intestin tissue using E-Cadherin Rabbit mAb (A20798) at a dilution of 1:500 (40x lens). High pressure antigen retrieval performed with 0.01M Tris-EDTA Buffer(pH 9.0) prior to IHC staining.