abcam

Product datasheet

Anti-E Cadherin antibody [EPR16845-34] - BSA and Azide free ab251536

Recombinant RabMAb

4 Images

Overview

Product name	Anti-E Cadherin antibody [EPR16845-34] - BSA and Azide free
Description	Rabbit monoclonal [EPR16845-34] to E Cadherin - BSA and Azide free
Host species	Rabbit
Specificity	E-cadherin contains a number of cleavage sites which may yield a complex fragmentation pattern in WB. Multiple bands between ~80-120 kDa may be observed.
Tested applications	**Suitable for:** WB, IP
Species reactivity	**Reacts with:** Mouse, Rat
Immunogen	Recombinant fragment. This information is proprietary to Abcam and/or its suppliers.
General notes	ab251536 is the carrier-free version of **ab212059**. Our **carrier-free** antibodies are typically supplied in a PBS-only formulation, purified and free of BSA, sodium azide and glycerol. The carrier-free buffer and high concentration allow for increased conjugation efficiency. 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. 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. This product is a recombinant monoclonal antibody, which offers several advantages including: - High batch-to-batch consistency and reproducibility - Improved sensitivity and specificity - Long-term security of supply - Animal-free production For more information **see here**. Our RabMAb® technology is a patented hybridoma-based technology for making rabbit monoclonal antibodies. For details on our patents, please refer to **RabMAb® patents**.

Properties

Form	Liquid
Storage instructions	Shipped at 4°C. Store at +4°C. Do Not Freeze.
Storage buffer	pH: 7.2 Constituent: PBS
Carrier free	Yes
Purity	Protein A purified
Clonality	Monoclonal
Clone number	EPR16845-34
Isotype	IgG

Applications

The Abpromise guarantee Our **Abpromise guarantee** covers the use of ab251536 in the following tested applications.

The application notes include recommended starting dilutions; optimal dilutions/concentrations should be determined by the end user.

Application	Abreviews	Notes
WB		Use at an assay dependent concentration. Detects a band of approximately 80-120 kDa (predicted molecular weight: 98 kDa).
IP		Use at an assay dependent concentration.

Target

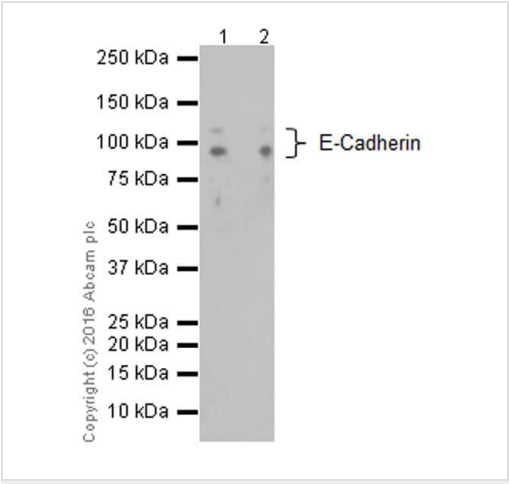
Function Cadherins are calcium-dependent cell adhesion proteins. They preferentially interact with themselves in a homophilic manner in connecting cells; cadherins may thus contribute to the sorting of heterogeneous cell types. CDH1 is involved in mechanisms regulating cell-cell adhesions, mobility and proliferation of epithelial cells. Has a potent invasive suppressor role. It is a ligand for integrin alpha-E/beta-7.
E-Cad/CTF2 promotes non-amyloidogenic degradation of Abeta precursors. Has a strong inhibitory effect on APP C99 and C83 production.

Tissue specificity Non-neural epithelial tissues.

Involvement in disease Defects in CDH1 are the cause of hereditary diffuse gastric cancer (HDGC) [MIM:137215]. An autosomal dominant cancer predisposition syndrome with increased susceptibility to diffuse gastric cancer. Diffuse gastric cancer is a malignant disease characterized by poorly differentiated infiltrating lesions resulting in thickening of the stomach. Malignant tumors start in the stomach, can spread to the esophagus or the small intestine, and can extend through the stomach wall to nearby lymph nodes and organs. It also can metastasize to other parts of the body. Note=Heterozygous germline mutations CDH1 are responsible for familial cases of diffuse gastric cancer. Somatic mutations in the has also been found in patients with sporadic diffuse gastric cancer and lobular breast cancer.
Defects in CDH1 are a cause of susceptibility to endometrial cancer (ENDMC) [MIM:608089].
Defects in CDH1 are a cause of susceptibility to ovarian cancer (OC) [MIM:167000]. Ovarian

	cancer common malignancy originating from ovarian tissue. Although many histologic types of ovarian neoplasms have been described, epithelial ovarian carcinoma is the most common form. Ovarian cancers are often asymptomatic and the recognized signs and symptoms, even of late-stage disease, are vague. Consequently, most patients are diagnosed with advanced disease.
Sequence similarities	Contains 5 cadherin domains.
Post-translational modifications	During apoptosis or with calcium influx, cleaved by a membrane-bound metalloproteinase (ADAM10), PS1/gamma-secretase and caspase-3 to produce fragments of about 38 kDa (E-CAD/CTF1), 33 kDa (E-CAD/CTF2) and 29 kDa (E-CAD/CTF3), respectively. Processing by the metalloproteinase, induced by calcium influx, causes disruption of cell-cell adhesion and the subsequent release of beta-catenin into the cytoplasm. The residual membrane-tethered cleavage product is rapidly degraded via an intracellular proteolytic pathway. Cleavage by caspase-3 releases the cytoplasmic tail resulting in disintegration of the actin microfilament system. The gamma-secretase-mediated cleavage promotes disassembly of adherens junctions.
Cellular localization	Cell junction. Cell membrane. Endosome. Golgi apparatus > trans-Golgi network. Colocalizes with DLGAP5 at sites of cell-cell contact in intestinal epithelial cells. Anchored to actin microfilaments through association with alpha-, beta- and gamma-catenin. Sequential proteolysis induced by apoptosis or calcium influx, results in translocation from sites of cell-cell contact to the cytoplasm. Colocalizes with RAB11A endosomes during its transport from the Golgi apparatus to the plasma membrane.

Images



Western blot - Anti-E Cadherin antibody [EPR16845-34] - BSA and Azide free (ab251536)

Lane 1 : Anti-E Cadherin antibody [EPR16845-34] (**ab212059**) at 1/20000 dilution
Lane 2 : Anti-E Cadherin antibody [EPR16845-34] (**ab212059**) at 1/100000 dilution

All lanes : Mouse E-Cadherin active protein (aa1-709)

Lysates/proteins at 0.01 µg per lane.

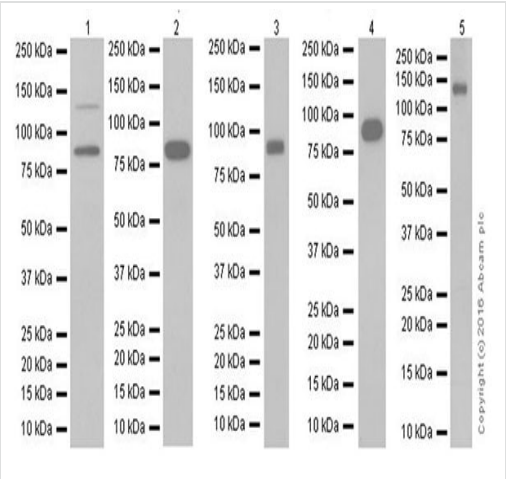
Secondary
All lanes : Goat Anti-Rabbit IgG H&L (HRP) (**ab97051**) at 1/20000 dilution

Predicted band size: 98 kDa
Observed band size: 120,84 kDa

Exposure time: 10 seconds

This data was developed using **ab212059**, the same antibody clone in a different buffer formulation.

Blocking and dilution buffer: 5% NFDM/TBST.



Western blot - Anti-E Cadherin antibody [EPR16845-34] - BSA and Azide free (ab251536)

Lanes 1-2 : Anti-E Cadherin antibody [EPR16845-34] (**ab212059**) at 1/5000 dilution
Lanes 3-5 : Anti-E Cadherin antibody [EPR16845-34] (**ab212059**) at 1/1000 dilution

Lane 1 : Rat spleen lysate
Lane 2 : Rat serum
Lane 3 : Mouse plasma
Lane 4 : Mouse serum
Lane 5 : Mouse brain lysate

Lysates/proteins at 10 µg per lane.

Secondary
Lanes 1 & 5 : Goat Anti-Rabbit IgG H&L (HRP) (**ab97051**) at 1/100000 dilution
Lanes 2-4 : Goat Anti-Rabbit IgG Peroxidase Conjugate, specific to the non-reduced form of IgG at 1/10000 dilution

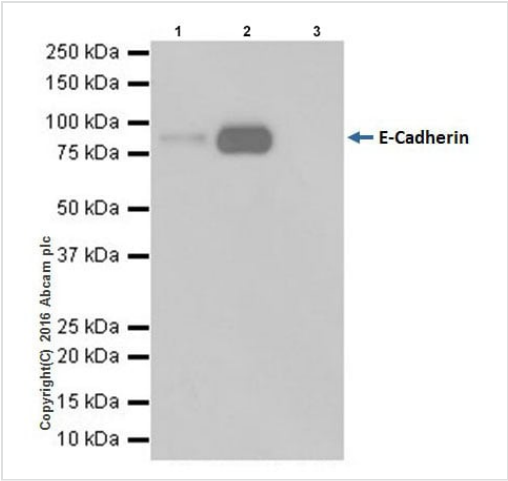
Predicted band size: 98 kDa
Observed band size: 120,84 kDa

This data was developed using **ab212059**, the same antibody clone in a different buffer formulation.

Blocking and dilution buffer: 5% NFDM/TBST.

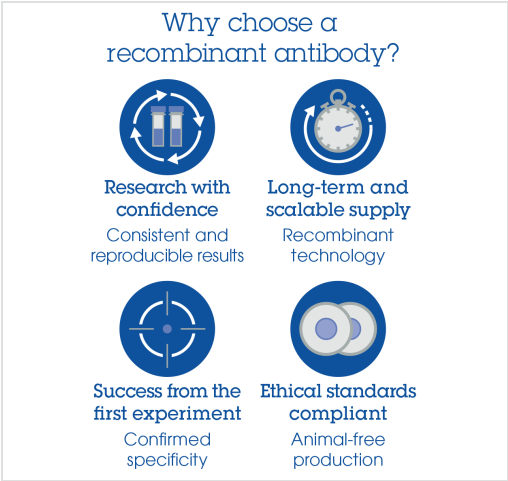
Exposure time: Lanes 1, 4, and 5: 30 seconds; Lane 2: 3 minutes; Lane 3: 15 seconds.

The expression profile observed is consistent with what has been described in the literature (PMID: 11076937; 11953314).



Immunoprecipitation - Anti-E Cadherin antibody [EPR16845-34] - BSA and Azide free (ab251536)

This data was developed using **ab212059**, the same antibody clone in a different buffer formulation.E Cadherin was immunoprecipitated from 1 mg of mouse serum with **ab212059** at 1/40 dilution. Western blot was performed from the immunoprecipitate using **ab212059** at 1/1000 dilution. VeriBlot for IP Detection Reagent (HRP) (**ab131366**), was used for detection at 1/10,000 dilution Lane 1: Mouse serum 10 µg (Input). Lane 2: **ab212059** IP in mouse serum. Lane 3: Rabbit monoclonal IgG (**ab172730**) instead of **ab212059** in mouse serum. Blocking and dilution buffer and concentration: 5% NFDM/TBST. Exposure time: 1 second.



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