abcam

Product datasheet

Anti-LRP6 antibody [EPR2423(2)] - BSA and Azide free ab232484



Recombinant

RabMAb

2 Images

Overview

Product name Anti-LRP6 antibody [EPR2423(2)] - BSA and Azide free

Description Rabbit monoclonal [EPR2423(2)] to LRP6 - BSA and Azide free

Host species Rabbit

Tested applications Suitable for: WB

Unsuitable for: Flow Cyt,ICC/IF,IHC-P or IP

Species reactivity Reacts with: Human

Predicted to work with: Mouse, Rat

Immunogen Synthetic peptide. This information is proprietary to Abcam and/or its suppliers.

Positive control WB: HAP1, HeLa, HepG2, 293T, and Jurkat whole cell lysate (ab7899).

General notes ab232484 is the carrier-free version of **ab134146**.

Our <u>carrier-free</u> 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 <u>conjugation kits</u> 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

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

ClonalityMonoclonalClone numberEPR2423(2)

Isotype IgG

Applications

The Abpromise guarantee Our Abpromise guarantee covers the use of ab232484 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 220 kDa (predicted molecular weight: 180 kDa).

Application notes Is unsuitable for Flow Cyt,ICC/IF,IHC-P or IP.

Target

Function Component of the Wnt-Fzd-LRP5-LRP6 complex that triggers beta-catenin signaling through

inducing aggregation of receptor-ligand complexes into ribosome-sized signalsomes. Cell-surface coreceptor of Wnt/beta-catenin signaling, which plays a pivotal role in bone formation. The Wnt-induced Fzd/LRP6 coreceptor complex recruits DVL1 polymers to the plasma membrane which, in turn, recruits the AXIN1/GSK3B-complex to the cell surface promoting the formation of signalsomes and inhibiting AXIN1/GSK3-mediated phosphorylation and destruction of beta-

catenin. Required for posterior patterning of the epiblast during gastrulation.

Tissue specificity Widely co-expressed with LRP5 during embryogenesis and in adult tissues.

Involvement in disease Defects in LRP6 are the cause of autosomal dominant coronary artery disease type 2 (ADCAD2)

[MIM:610947].

Sequence similarities Belongs to the LDLR family.

Contains 4 EGF-like domains.

Contains 3 LDL-receptor class A domains. Contains 20 LDL-receptor class B repeats.

Domain The YWTD-EGF-like domains 1 and 2 are required for the interaction with Wnt-frizzled complex.

The YWTD-EGF-like domains 3 and 4 are required for the interaction with DKK1.

The PPPSP motifs play a central role in signal transduction by being phosphorylated, leading to

Post-translational modifications

activate the Wnt signaling pathway.

Dual phosphorylation of cytoplasmic PPPSP motifs sequentially by GSK3 and CK1 is required for AXIN1-binding, and subsequent stabilization and activation of beta-catenin via preventing GSK3-mediated phosphorylation of beta-catenin. Phosphorylated, in vitro, by GRK5/6 within and outside the PPPSP motifs. Phosphorylation at Ser-1490 by CDK14 during G2/M phase leads to regulation of the Wnt signaling pathway during the cell cycle. Phosphorylation by GSK3B is induced by RPSO1 binding and inhibited by DKK1. Phosphorylated, in vitro, by casein kinase I on Thr-1479.

Undergoes gamma-secretase-dependent regulated intramembrane proteolysis (RIP). The extracellular domain is first released by shedding, and then, through the action of gamma-secretase, the intracellular domain (ICD) is released into the cytoplasm where it is free to bind to GSK3B and to activate canonical Wnt signaling.

Palmitoylation on the two sites near the transmembrane domain leads to release of LRP6 from the endoplasmic reticulum.

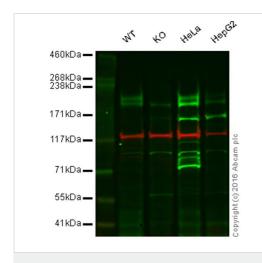
Mono-ubiquitinated which retains LRP6 in the endoplasmic reticulum.

N-glycosylation is required for cell surface location.

Cellular localization

Membrane. Endoplasmic reticulum. On Wnt signaling, undergoes a cycle of caveolin- or clathrin-mediated endocytosis and plasma membrane location. Released from the endoplasmic reticulum on palmitoylation. Mono-ubiquitination retains it in the endoplasmic reticulum in the absence of palmitoylation. On Wnt signaling, phosphorylated, aggregates and colocalizes with AXIN1 and GSK3B at the plasma membrane in LRP6-signalsomes. Chaperoned to the plasma membrane by MESD.

Images



Western blot - Anti-LRP6 antibody [EPR2423(2)] - BSA and Azide free (ab232484)

Lane 1: Wild-type HAP1 cell lysate (20 µg)

Lane 2: LRP6 knockout HAP1 cell lysate (20 µg)

Lane 3: HeLa (human epithelial cell line from cervix adenocarcinoma) cell lysate (20 µg)

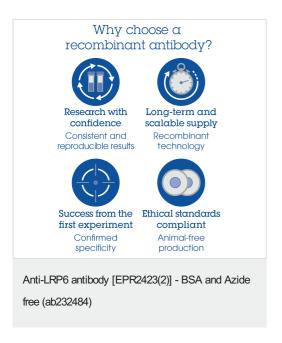
Lane 4: HepG2 (human liver hepatocellular carcinoma cell line) cell lysate (20 µg)

Lanes 1 - 4: Merged signal (red and green). Green - <u>ab134146</u> observed at 220 kDa. Red - loading control, <u>ab18058</u>, observed at 124 kDa.

ab134146 was shown to recognize LRP6 when LRP6 knockout samples were used, along with additional cross-reactive bands. Wild-type and LRP6 knockout samples were subjected to SDS-PAGE. ab134146 and ab18058 (loading control to Vinculin) were diluted 1/10000 and 1/1000 respectively, and incubated overnight at 4°C. Blots were developed with Goat anti-Rabbit IgG H&L (IRDye® 800CW) preadsorbed (ab216773) and Goat anti-Mouse IgG H&L (IRDye® 680RD) preadsorbed (ab216776) secondary antibodies at 1/10000 dilution for 1 hour at room temperature before imaging.

This data was developed using the same antibody clone in a

different buffer formulation containing PBS, BSA, glycerol, and sodium azide (<u>ab134146</u>).



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