

Product datasheet

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

KO VALIDATED 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 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 <u>see here</u> . Our RabMAb [®] technology is a patented hybridoma-based technology for making rabbit
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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	EPR2423(2)
Isotype	lgG

Applications

Properties

The Abpromise guarantee Our <u>Abpromise guarantee</u> 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-Lf inducing aggregation of recept surface coreceptor of Wnt/bet Wnt-induced Fzd/LRP6 corect which, in turn, recruits the AXI signalsomes and inhibiting AX catenin. Required for posterior	RP5-LRP6 complex that triggers beta-catenin signaling through otor-ligand complexes into ribosome-sized signalsomes. Cell- ca-catenin signaling, which plays a pivotal role in bone formation. The reptor complex recruits DVL1 polymers to the plasma membrane N1/GSK3B-complex to the cell surface promoting the formation of KIN1/GSK3-mediated phosphorylation and destruction of beta- or 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 clas Contains 20 LDL-receptor cla	s. s A domains. iss B repeats.
Domain	The YWTD-EGF-like domains The YWTD-EGF-like domains The PPPSP motifs play a cer	and 2 are required for the interaction with Wnt-frizzled complex. and 4 are required for the interaction with DKK1. atral role in signal transduction by being phosphorylated, leading to

	activate the Wnt signaling pathway.
Post-translational modifications	 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.
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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