



PCSK9 Rabbit mAb

Catalog # AP75878

Product Information

ApplicationWB, IPPrimary AccessionQ8NBP7ReactivityHumanHostRabbit

Clonality Monoclonal Antibody

Calculated MW 74286

Additional Information

Gene ID 255738

Other Names PCSK9

Dilution WB~~1/500-1/1000 IP~~N/A

Format Liquid

Protein Information

Name PCSK9

Synonyms NARC1

Function Crucial player in the regulation of plasma cholesterol homeostasis. Binds to

low-density lipid receptor family members: low density lipoprotein receptor (LDLR), very low density lipoprotein receptor (VLDLR), apolipoprotein E receptor (LRP1/APOER) and apolipoprotein receptor 2 (LRP8/APOER2), and

promotes their degradation in intracellular acidic compartments

(PubMed:<u>18039658</u>). Acts via a non- proteolytic mechanism to enhance the degradation of the hepatic LDLR through a clathrin LDLRAP1/ARH-mediated pathway. May prevent the recycling of LDLR from endosomes to the cell surface or direct it to lysosomes for degradation. Can induce ubiquitination of

LDLR leading to its subsequent degradation (PubMed: 17461796,

PubMed: 18197702, PubMed: 18799458, PubMed: 22074827). Inhibits intracellular degradation of APOB via the autophagosome/lysosome pathway in a LDLR-independent manner. Involved in the disposal of non-acetylated intermediates of BACE1 in the early secretory pathway (PubMed: 18660751). Inhibits epithelial Na(+) channel (ENaC)-mediated Na(+) absorption by reducing ENaC surface expression primarily by increasing its proteasomal degradation. Regulates neuronal apoptosis via modulation of LRP8/APOER2

levels and related anti-apoptotic signaling pathways.

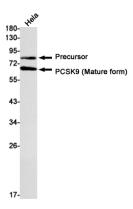
Cellular Location Cytoplasm. Secreted. Endosome. Lysosome. Cell surface. Endoplasmic

reticulum. Golgi apparatus. Note=Autocatalytic cleavage is required to transport it from the endoplasmic reticulum to the Golgi apparatus and for the secretion of the mature protein Localizes to the endoplasmic reticulum in the absence of LDLR and colocalizes to the cell surface and to the endosomes/lysosomes in the presence of LDLR. The sorting to the cell surface and endosomes is required in order to fully promote LDLR degradation

Tissue Location

Expressed in neuro-epithelioma, colon carcinoma, hepatic and pancreatic cell lines, and in Schwann cells

Images



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