

Anti-VPS54 Antibody

Rabbit polyclonal antibody to VPS54 Catalog # AP60827

Product Information

Application WB, IHC Primary Accession Q9P1Q0

Reactivity Human, Mouse, Rat

HostRabbitClonalityPolyclonalCalculated MW110589

Additional Information

Gene ID 51542

Other Names HCC8; Vacuolar protein sorting-associated protein 54; Hepatocellular

carcinoma protein 8; Tumor antigen HOM-HCC-8; Tumor antigen SLP-8p

Target/Specificity KLH-conjugated synthetic peptide encompassing a sequence within the center

region of human VPS54. The exact sequence is proprietary.

Dilution WB~~WB (1/500 - 1/2000), IHC (1/50 - 1/200) IHC~~WB (1/500 - 1/2000), IHC

(1/50 - 1/200)

Format Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30%

glycerol, and 0.09% (W/V) sodium azide.

Storage Store at -20 °C.Stable for 12 months from date of receipt

Protein Information

Name VPS54

Synonyms HCC8

Function Acts as a component of the GARP complex that is involved in retrograde

transport from early and late endosomes to the trans-Golgi network (TGN). The GARP complex is required for the maintenance of the cycling of mannose 6-phosphate receptors between the TGN and endosomes, this cycling is necessary for proper lysosomal sorting of acid hydrolases such as CTSD (PubMed:18367545). Within the GARP complex, required to tether the complex to the TGN. Not involved in endocytic recycling (PubMed:25799061).

Cellular Location Golgi apparatus, trans-Golgi network. Membrane

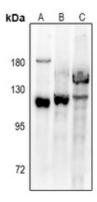
{ECO:0000250|UniProtKB:Q9JMK8}. Note=Associates with membranes in an

EIPR1-independent manner. {ECO:0000250 | UniProtKB:Q9JMK8}

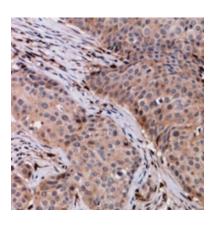
Background

KLH-conjugated synthetic peptide encompassing a sequence within the center region of human VPS54. The exact sequence is proprietary.

Images



Western blot analysis of VPS54 expression in mouse testis (A), rat testis (B), HEK293T (C) whole cell lysates.



Immunohistochemical analysis of VPS54 staining in human breast cancer formalin fixed paraffin embedded tissue section. The section was pre-treated using heat mediated antigen retrieval with sodium citrate buffer (pH 6.0). The section was then incubated with the antibody at room temperature and detected using an HRP conjugated compact polymer system. DAB was used as the chromogen. The section was then counterstained with haematoxylin and mounted with DPX.

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