

NPRL2 Antibody (C-term)

Affinity Purified Rabbit Polyclonal Antibody (Pab) Catalog # AP16818B

Specification

NPRL2 Antibody (C-term) - Product Information

Application WB,E
Primary Accession O8WTW4

Other Accession <u>Q9WUE4</u>, <u>NP_006536.3</u>

Reactivity
Predicted
Host
Clonality
Isotype
Calculated MW
Antigen Region
Human
Mouse
Rabbit
Polyclonal
Rabbit IgG
A3658
A16-373

NPRL2 Antibody (C-term) - Additional Information

Gene ID 10641

Other Names

Nitrogen permease regulator 2-like protein, NPR2-like protein, Gene 21 protein, G21 protein, Tumor suppressor candidate 4, NPRL2, TUSC4

Target/Specificity

This NPRL2 antibody is generated from rabbits immunized with a KLH conjugated synthetic peptide between 346-373 amino acids from the C-terminal region of human NPRL2.

Dilution

WB~~1:1000

Format

Purified polyclonal antibody supplied in PBS with 0.09% (W/V) sodium azide. This antibody is purified through a protein A column, followed by peptide affinity purification.

Storage

Maintain refrigerated at 2-8°C for up to 2 weeks. For long term storage store at -20°C in small aliquots to prevent freeze-thaw cycles.

Precautions

NPRL2 Antibody (C-term) is for research use only and not for use in diagnostic or therapeutic procedures.

NPRL2 Antibody (C-term) - Protein Information

Name NPRL2 {ECO:0000303|PubMed:18616680, ECO:0000312|HGNC:HGNC:24969}



Function Catalytic component of the GATOR1 complex, a multiprotein complex that functions as an inhibitor of the amino acid-sensing branch of the mTORC1 pathway (PubMed:23723238, PubMed:29590090, PubMed:35338845, PubMed:38006878). In response to amino acid depletion, the GATOR1 complex has GTPase activating protein (GAP) activity and strongly increases GTP hydrolysis by RagA/RRAGA (or RagB/RRAGB) within heterodimeric Rag complexes, thereby turning them into their inactive GDP-bound form, releasing mTORC1 from lysosomal surface and inhibiting mTORC1 signaling (PubMed:23723238, PubMed:29590090, PubMed:35338845). In the presence of abundant amino acids, the GATOR1 complex is ubiquitinated and inhibited by GATOR2 (PubMed:23723238, PubMed:36528027). Within the GATOR1 complex, NPRL2 constitutes the catalytic subunit that mediates the GTPase activator activity and under methionine-sufficient conditions, the GTPase activator activity is inhibited by PRMT1 through methylation and consequently inducing timely mTORC1 activation (PubMed:30651352, PubMed:35338845, PubMed:27173016).

Cellular Location

Lysosome membrane. Note=Localization to lysosomes is mediated by the KICSTOR complex and is amino acid-independent.

Tissue Location

Most abundant in skeletal muscle, followed by brain, liver and pancreas, with lower amounts in lung, kidney, placenta and heart. Expressed in the frontal lobe cortex as well as in the temporal, parietal, and occipital lobes (PubMed:27173016, PubMed:26505888). Expressed in most lung cancer cell lines tested

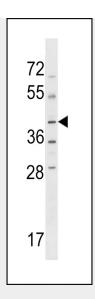
NPRL2 Antibody (C-term) - Protocols

Provided below are standard protocols that you may find useful for product applications.

- Western Blot
- Blocking Peptides
- Dot Blot
- Immunohistochemistry
- Immunofluorescence
- <u>Immunoprecipitation</u>
- Flow Cytomety
- Cell Culture

NPRL2 Antibody (C-term) - Images





NPRL2 Antibody (C-term) (Cat. #AP16818b) western blot analysis in HL-60 cell line lysates (35ug/lane). This demonstrates the NPRL2 antibody detected the NPRL2 protein (arrow).

NPRL2 Antibody (C-term) - Background

Suppresses Src-dependent tyrosine phosphorylation and activation of PDPK1 and its downstream signaling. Down-regulates PDPK1 kinase activity by interfering with tyrosine phosphorylation at the Tyr-9 Tyr-373 and Tyr-376 residues. May act as a tumor suppressor. Suppresses cell growth and enhanced sensitivity to various anticancer drugs.

NPRL2 Antibody (C-term) - References

Spielewoy, N., et al. Eukaryotic Cell 9(4):592-601(2010) Otani, S., et al. J Surg Oncol 100(5):358-363(2009) Neklesa, T.K., et al. PLoS Genet. 5 (6), E1000515 (2009) : Anedchenko, E.A., et al. Mol. Biol. (Mosk.) 42(6):965-976(2008) Kurata, A., et al. Cancer Sci. 99(9):1827-1834(2008)