

CCDC134 Antibody (C-term)

Affinity Purified Rabbit Polyclonal Antibody (Pab) Catalog # AP13242b

Product Information

Application WB, IHC-P, E Primary Accession Q9H6E4

Other Accession <u>Q5M862</u>, <u>NP 079097.1</u>

Reactivity Human **Predicted** Rat Host Rabbit Clonality Polyclonal Isotype Rabbit IgG **Clone Names** RB33111 **Calculated MW** 26561 **Antigen Region** 197-225

Additional Information

Gene ID 79879

Other Names Coiled-coil domain-containing protein 134, CCDC134

Target/SpecificityThis CCDC134 antibody is generated from rabbits immunized with a KLH

conjugated synthetic peptide between 197-225 amino acids from the

C-terminal region of human CCDC134.

Dilution WB~~1:1000 IHC-P~~1:100~500 E~~Use at an assay dependent concentration.

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 CCDC134 Antibody (C-term) is for research use only and not for use in

diagnostic or therapeutic procedures.

Protein Information

Name CCDC134 {ECO:0000303 | PubMed:39509507,

ECO:0000312 | HGNC:HGNC:26185}

Function Molecular adapter required to prevent protein hyperglycosylation of

HSP90B1: during translation, associates with nascent HSP90B1 and the STT3A

catalytic component of the OST-A complex and tethers them to a specialized translocon that forms a microenvironment for HSP90B1 folding (PubMed:38670073, PubMed:39509507). In the CCDC134-containing translocon, STT3A associates with the SRT pseudosubstrate motif of HSP90B1, preventing access to facultative glycosylation sites until folding is completed, preventing hyperglycosylation and subsequent degradation of HSP90B1 (PubMed:39509507). In extracellular secreted form, promotes proliferation and activation of CD8(+) T-cells, suggesting a cytokine- like function (PubMed: 25125657). May inhibit ERK and JNK signaling activity (PubMed: 18087676, PubMed: 23070808). May suppress cell migration and invasion activity, via its effects on ERK and JNK signaling (PubMed: 23070808). May also localize in the nucleus: enhances stability of the PCAF histone acetyltransferase (HAT) complex member TADA2A and thus promotes PCAF-mediated histone acetyltransferase activity (PubMed: 22644376). Has a critical role in the regulation of osteogenesis and bone development (PubMed:32181939).

Cellular Location

Endoplasmic reticulum lumen. Secreted. Cytoplasm Nucleus. Note=Mainly localizes to the endoplasmic reticulum (PubMed:39509507). Accumulates in the nucleus in response to UV irradiation (PubMed:22644376)

Tissue Location

Expressed in cervical gland, cervical squamous epithelium, endometrium, stomach, kidney distal convoluted tubule, spermatogenic cells in testis, mammary gland, liver and striated muscle (at protein level) (PubMed:18087676, PubMed:23070808). Also detected in placenta (PubMed:18087676). Highest expression in testis relative to other tissues (PubMed:18087676). Detected in T cells and dendritic cells; highly expressed in activated CD8(+) T cells, and also expressed at lower levels in CD4(+) T cells (PubMed:25125657)

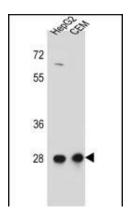
Background

The specific function of this protein remains unknown.

References

Huang, J., et al. Cell. Mol. Life Sci. 65(2):338-349(2008) Lim, J., et al. Cell 125(4):801-814(2006) Collins, J.E., et al. Genome Biol. 5 (10), R84 (2004): Dunham, I., et al. Nature 402(6761):489-495(1999)

Images



CCDC134 Antibody (C-term) (Cat. #AP13242b) western blot analysis in HepG2,CEM cell line lysates (35ug/lane).This demonstrates the CCDC134 antibody detected the CCDC134 protein (arrow).



CCDC134 Antibody (C-term) (Cat. #AP13242b)immunohistochemistry analysis in formalin fixed and paraffin embedded human brain tissue followed by peroxidase conjugation of the secondary antibody and DAB staining. This data demonstrates the use of CCDC134 Antibody (C-term) for immunohistochemistry. Clinical relevance has not been evaluated.

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