

DDR2 Antibody(Ascites)

Mouse Monoclonal Antibody (Mab) Catalog # AM2058a

Product Information

Application WB, IHC-P, E Primary Accession Q16832

Other Accession NP_001014796.1

Reactivity Human

Host Mouse

Clonality Monoclonal

Isotype IgG1

Clone Names 484CT5.5.4

Calculated MW 96736

Antigen Region 290-320

Additional Information

Gene ID 4921

Other Names Discoidin domain-containing receptor 2, Discoidin domain receptor 2, CD167

antigen-like family member B, Discoidin domain-containing receptor tyrosine

kinase 2, Neurotrophic tyrosine kinase, receptor-related 3, Receptor

protein-tyrosine kinase TKT, Tyrosine-protein kinase TYRO10, CD167b, DDR2,

NTRKR3, TKT, TYRO10

Target/SpecificityThis DDR2 antibody is generated from mice immunized with a KLH conjugated

synthetic peptide between 290-320 amino acids from human DDR2.

Dilution WB~~1:100~1600 IHC-P~~1:200 E~~Use at an assay dependent concentration.

Format Mouse monoclonal antibody supplied in crude ascites with 0.09% (W/V)

sodium azide.

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 DDR2 Antibody(Ascites) is for research use only and not for use in diagnostic

or therapeutic procedures.

Protein Information

Name DDR2

Synonyms NTRKR3, TKT, TYRO10

Function

Tyrosine kinase involved in the regulation of tissues remodeling (PubMed:30449416). It functions as a cell surface receptor for fibrillar collagen and regulates cell differentiation, remodeling of the extracellular matrix, cell migration and cell proliferation. Required for normal bone development. Regulates osteoblast differentiation and chondrocyte maturation via a signaling pathway that involves MAP kinases and leads to the activation of the transcription factor RUNX2. Regulates remodeling of the extracellular matrix by up- regulation of the collagenases MMP1, MMP2 and MMP13, and thereby facilitates cell migration and tumor cell invasion. Promotes fibroblast migration and proliferation, and thereby contributes to cutaneous wound healing.

Cellular Location

Cell membrane; Single-pass type I membrane protein

Tissue Location

Detected in osteocytes, osteoblastic cells in subchondral bone, bone lining cells, tibia and cartilage (at protein level). Detected at high levels in heart and lung, and at low levels in brain, placenta, liver, skeletal muscle, pancreas, and kidney

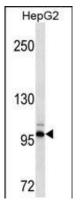
Background

Receptor tyrosine kinases (RTKs) play a key role in the communication of cells with their microenvironment. These molecules are involved in the regulation of cell growth, differentiation, and metabolism. In several cases the biochemical mechanism by which RTKs transduce signals across the membrane has been shown to be ligand induced receptor oligomerization and subsequent intracellular phosphorylation. This autophosphorylation leads to phosphorylation of cytosolic targets as well as association with other molecules, which are involved in pleiotropic effects of signal transduction. RTKs have a tripartite structure with extracellular, transmembrane, and cytoplasmic regions. This gene encodes a member of a novel subclass of RTKs and contains a distinct extracellular region encompassing a factor VIII-like domain. Alternative splicing in the 5' UTR results in multiple transcript variants encoding the same protein. [provided by RefSeq].

References

Rose, J.E., et al. Mol. Med. 16 (7-8), 247-253 (2010): Ali, B.R., et al. Hum. Mol. Genet. 19(11):2239-2250(2010) Johnatty, S.E., et al. PLoS Genet. 6 (7), E1001016 (2010): Ehret, G.B., et al. Eur. J. Hum. Genet. 17(12):1650-1657(2009): Su, J., et al. Mol. Cell. Biochem. 330 (1-2), 141-152 (2009):

Images



DDR2 Antibody (Cat. #AM2058a) western blot analysis in HepG2 cell line lysates (35µg/lane). This demonstrates the DDR2 antibody detected the DDR2 protein (arrow).

Immunohistochemical analysis of paraffin-embedded



Human lung section using Pink1(Cat#am2058a). am2058a was diluted at 1:200 dilution. A undiluted biotinylated goat polyvalent antibody was used as the secondary, followed by DAB staining.

Please note: All products are 'FOR RESEARCH USE ONLY. NOT FOR USE IN DIAGNOSTIC OR THERAPEUTIC PROCEDURES'.