

A Reliable Research Partner in Life Science and Medicine

Recombinant Wnt5a Monoclonal Antibody

catalog number: E-AB-81621

Note: Centrifuge before opening to ensure complete recovery of vial contents.

Description

Reactivity Human

Immunogen A synthetic peptide of human Wnt5a

HostRabbitIsotypeIgGCloneR08-4A3

Purification Affinity Purified

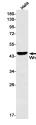
Buffer 50mM Tris-Glycine(pH 7.4), 0.15M NaCl, 40% Glycerol, 0.05% stabilizer and 0.05%

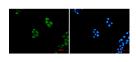
protective protein.

Applications Recommended Dilution

WB 1:500-1:1000 **IF** 1:50-1:100

Data





Western blot detection of Wnt5a in Hela cell lysates using Wnt5a Rabbit mAb(1:500 diluted).Predicted band size:42kDa.Observed band size:45kDa.

Immunofluorescence of Wnt5a (green) in hela using Wnt5a Rabbit mAb at dilution 1:50, and DAPI(blue)

Observed-MW:45 kDa Calculated-MW:42 kDa

Preparation & Storage

Storage Storage Store at -20°C Valid for 12 months. Avoid freeze / thaw cycles.

Shipping The product is shipped with ice pack, upon receipt, store it immediately at the

temperature recommended.

Background

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Elabscience Bionovation Inc.



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The WNT gene family consists of structurally related genes which encode secreted signaling proteins. These proteins have been implicated in oncogenesis and in several developmental processes, including regulation of cell fate and patterning during embryogenesis. This gene encodes a member of the WNT family that signals through both the canonical and non-canonical WNT pathways. This protein is a ligand for the seven transmembrane receptor frizzled-5 and the tyrosine kinase orphan receptor 2. This protein plays an essential role in regulating developmental pathways during embryogenesis. This protein may also play a role in oncogenesis. Mutations in this gene are the cause of autosomal dominant Robinow syndrome. Alternate splicing results in multiple transcript variants. WNT5A (Wnt Family Member 5A) is a Protein Coding gene. Diseases associated with WNT5A include Robinow Syndrome, Autosomal Dominant 1 and Autosomal Dominant Robinow Syndrome. Among its related pathways are Validated targets of C-MYC transcriptional repression and Wnt Signaling Pathways: beta-Catenin-independent Wnt/Ca2+ Signaling and Other Noncanonical Wnt Signaling Pathways. GO annotations related to this gene include transcription factor activity, sequence-specific DNA binding and protein domain specific binding. An important paralog of this gene is WNT5B.

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