

Rab 3 GAP p150 Polyclonal Antibody

Catalog No: YT3927

Reactivity: Human; Mouse; Rat; Monkey

Applications: WB;IHC;IF;ELISA

Target: Rab 3 GAP p150

Gene Name: RAB3GAP2

Protein Name: Rab3 GTPase-activating protein non-catalytic subunit

Human Gene Id: 25782

Human Swiss Prot

No:

Mouse Gene ld: 98732

Mouse Swiss Prot

No:

Rat Gene Id: 289350

Rat Swiss Prot No: Q5U1Z0

Immunogen: The antiserum was produced against synthesized peptide derived from human

RAB3GAP2. AA range:417-466

Specificity: Rab 3 GAP p150 Polyclonal Antibody detects endogenous levels of Rab 3 GAP

p150 protein.

Q9H2M9

Q8BMG7

Formulation: Liquid in PBS containing 50% glycerol, 0.5% BSA and 0.02% sodium azide.

Source: Polyclonal, Rabbit, IgG

Dilution: WB 1:500 - 1:2000. IHC 1:100 - 1:300. ELISA: 1:40000.. IF 1:50-200

Purification: The antibody was affinity-purified from rabbit antiserum by affinity-

chromatography using epitope-specific immunogen.

Concentration: 1 mg/ml

Storage Stability: -15°C to -25°C/1 year(Do not lower than -25°C)

Observed Band: 156kD

Background: The protein encoded by this gene belongs to the RAB3 protein family, members

of which are involved in regulated exocytosis of neurotransmitters and hormones. This protein forms the Rab3 GTPase-activating complex with RAB3GAP1, where it constitutes the regulatory subunit, whereas the latter functions as the catalytic subunit. This gene has the highest level of expression in the brain, consistent with it having a key role in neurodevelopment. Mutations in this gene are associated

with Martsolf syndrome.[provided by RefSeg, Oct 2009],

Function: disease:Defects in RAB3GAP2 are the cause of Martsolf syndrome

[MIM:212720]. Martsolf syndrome is characterized by congenital cataracts,

mental retardation, and hypogonadism. Inheritance is autosomal

recessive.,function:Regulatory subunit of a GTPase activating protein that has specificity for Rab3 subfamily (RAB3A, RAB3B, RAB3C and RAB3D). Rab3 proteins are involved in regulated exocytosis of neurotransmitters and hormones. Rab3 GTPase-activating complex specifically converts active Rab3-GTP to the inactive form Rab3-GDP. Required for normal eye and brain development. May participate in neurodevelopmental processes such as proliferation, migration and differentiation before synapse formation, and non-synaptic vesicular release of neurotransmitters., similarity:Belongs to the Rab3-GAP regulatory subunit

family.,subcellular location:In neurons, it is enriched in the synaptic soluble fracti

Subcellular Location:

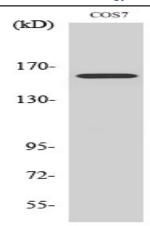
Cytoplasm. In neurons, it is enriched in the synaptic soluble fraction.

Expression: Ubiquitous.

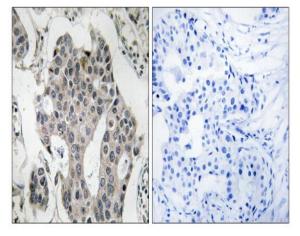
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No4:

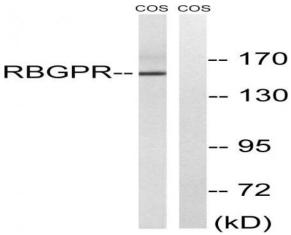
Products Images



Western Blot analysis of various cells using Rab 3 GAP p150 Polyclonal Antibody



Immunohistochemistry analysis of paraffin-embedded human breast carcinoma tissue, using RAB3GAP2 Antibody. The picture on the right is blocked with the synthesized peptide.



Western blot analysis of lysates from COS cells, using RAB3GAP2 Antibody. The lane on the right is blocked with the synthesized peptide.