

DNAPab

Hard-to-Find
Antibody

OPHN1 DNAPab

Catalog # H00004983-W01P

Size 200 ug

Specification

Product Description	Rabbit polyclonal antibody raised against a full-length human OPHN1 DNA using DNAX™ Immune technology.
Technology	DNAX™ Immune
Immunogen	Full-length human DNA
Sequence	MGHPPLEFSDCYLDSPDFRERLKCYEQELERTNKFIDVIKDGNALISAMRNYSSAVQKFSQTLQ SFQDFIGDITLTDDEINIAESFKEFAELLNEVENRMMMVMHNASDLLIKPLENFRKEQIGFTKERKK KFEKDGERFYSLDRHLHLSSKKKESQLQEADLQVDERHNFFESSLDYVYQIQEVQESKKFNV EPVLAFLHSLFISNSLTVELTQDFLPYKQQLQLSLQNTRNHFSSSTREEMEELKKRMKEAPQTCKLP GQPTIEGYLYTQEKWALGISWVKYYCQYEKETKTLTMTPEQKPGAKQGPLDLTKYCVRRKTESI DKRFCFDIETNERPGTITLQALSEANRRLWMEAMDGKEPYHSPITKQQEMELNEVGFKFVRKCINII ETKGIKTEGLYRTVGSNIQVQKLLNAFFDPKCPGDVDFHNSDWDIKTITSSLKFYLRNLSEPVMTYR LHKELVSAAKSDNLDYRLGAIHSLVYKLPEKNREMLELLIRHLVNVCEHSKENLMTPSNMGVIFGP TLMRAQEDTVAAMNIKFQNVVEILIEHFVKIYLPPEESAAPPVPPPRVTARRHKPITISKRLRLRER TVFYTSSLDESEDEIQHQTPNGTITSSIEPPKPPQHPKLPQIRSGETDPGRKSPSRPILDGKLEPCP EVDVGKLVSRQLDGGTKITPKATNGPMPGSGPTKTPSFHIKRPAPRPLAHHKEGDADSFSSKVRP PGEKPTIIRPPVRPPDPPCRAATPQKPEPKPDVAGNAGEITSSVVASRTRFFETASRKTGSSQGR LPGDES
Host	Rabbit
Reactivity	Human
Purification	Protein A
Quality Control Testing	Antibody reactive against mammalian transfected lysate.
Storage Buffer	In 1x PBS, pH 7.4
Storage Instruction	Store at -20°C or lower. Aliquot to avoid repeated freezing and thawing.

Applications

- Western Blot (Transfected lysate)

[Protocol Download](#)

- Immunofluorescence (Transfected cell)

- Flow Cytometry (Transfected cell)

Gene Info — OPHN1

Entrez GeneID [4983](#)

GeneBank Accession# [NM_002547.1](#)

Protein Accession# [NP_002538.1](#)

Gene Name OPHN1

Gene Alias MRX60, OPN1

Gene Description oligophrenin 1

Omim ID [300127](#) [300486](#)

Gene Ontology [Hyperlink](#)

Gene Summary This gene encodes a Rho-GTPase-activating protein that promotes GTP hydrolysis of Rho subfamily members. Rho proteins are important mediators of intracellular signal transduction, which affects cell migration and cell morphogenesis. Mutations in this gene are responsible for OPHN1-related X-linked mental retardation with cerebellar hypoplasia and distinctive facial dysmorphism. [provided by RefSeq]

Other Designations OTTHUMP00000023451|mental retardation, X-linked 60|oligophrenin-1, Rho-GTPase activating protein

Disease

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- [Cerebellar Diseases](#)
- [Coronary Vasospasm](#)
- [Facial Asymmetry](#)
- [Genetic Predisposition to Disease](#)

- [Mental Retardation](#)
- [Nervous System Malformations](#)