

DNAxPAb

Hard-to-Find
Antibody

GNB1L DNAxPab

Catalog # H00054584-W01P

Size 200 ug

Specification

Product Description Rabbit polyclonal antibody raised against a full-length human GNB1L DNA using DNAx™ Immune technology.

Technology [DNAx™ Immune](#)

Immunogen Full-length human DNA

Sequence MTAPCPPPPDPQFVLRGTQSPVHALHFCEGAQAQGRPLLFSGSQSGLVHWSLQTRRAVTTLD
GHGGQCVTWLQTLPPQGRQLLSQGRDLKLCLWDLAEGRSAVVDSVCLESVGFRCRSSILAGGQPR
WTLAVPGRGSDEVQILEMPSKTSCALKPKADAKLGMPMCLRLWQADCSSRPLL LAGYEDGSV
VLWDVSEQKVCRIACHEEPVMDLDFDSQKARGISGSAGKALAVWSLDWQQALQVRGTHELTN
PGIAEVTIRPDRKILATAGWDHRIRVFHWRTMQPLAVLAFHSAAVQCVAFTADGLLAAGSKDQRIS
LWSLYPRA

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 — GNB1L

Entrez GeneID [54584](#)

GeneBank Accession# [NM_053004.1](#)

Protein Accession# [NP_443730.1](#)

Gene Name GNB1L

Gene Alias DGCRK3, FKSG1, GY2, KIAA1645, WDR14, WDVCF

Gene Description guanine nucleotide binding protein (G protein), beta polypeptide 1-like

Omim ID [610778](#)

Gene Ontology [Hyperlink](#)

Gene Summary

This gene encodes a G-protein beta-subunit-like polypeptide which is a member of the WD repeat protein family. WD repeats are minimally conserved regions of approximately 40 amino acids typically bracketed by gly-his and trp-aspartate (GH-WD), which may facilitate formation of heterotrimeric or multiprotein complexes. Members of this family are involved in a variety of cellular processes, including cell cycle progression, signal transduction, apoptosis, and gene regulation. This protein contains 6 WD repeats and is highly expressed in the heart. The gene maps to the region on chromosome 22q11, which is deleted in DiGeorge syndrome, trisomic in derivative 22 syndrome and tetrasomic in cat-eye syndrome. Therefore, this gene may contribute to the etiology of those disorders. Transcripts from this gene share exons with some transcripts from the C22orf29 gene. [provided by RefSeq]

Other Designations G-protein beta subunit-like protein|guanine nucleotide binding protein beta-subunit-like polypeptide

Disease

- [Chromosome Deletion](#)
- [Schizophrenia](#)
- [Schizophrenic Psychology](#)