

DNAxPAb

Hard-to-Find
Antibody

SNTG1 DNAxPab

Catalog # H00054212-W01P

Size 200 ug

Specification

Product Description	Rabbit polyclonal antibody raised against a full-length human SNTG1 DNA using DNAx™ Immune technology.
Technology	DNAx™ Immune
Immunogen	Full-length human DNA
Sequence	MDFRTACEETKTGICLLQDGNQEPFKVRLHLAKDILMIQEQDVICVSGEPFYSGERTVTIRRQTVGG FGLSIKGGAEHNIPVVVSKISKEQRAELSGLLFIGDAILQINGINVRKCRHEEVVQVLRNAGEEVTLT VSFLKRAPAFLKLPLNEDCACAPSDQSSGTSSPLCDSGLHLNYHPNNTDTLSCSSWPTSPGLR WEKRWCDLRLIPLLHSRFSQYVPGTDLSRQNAFQVIAVDGVCTGIIQCLSAEDCVDWLQAIATNIS NLTKHNIKKINRNFVNQQVMGWCEAREQDPLQDRVYSPTFLALRGSCLYKFLAPPVTTWDWT RAEKTFSVYEIMCKILKDSDLLDRRKQCFTVQSEGEDLYFSVELESDLAQWERAFQTATFLEVE RIQCKTYACVLESHLMGLTIDFSTGFICFDAATKAVLWRYKFSQLKGSSDDGKSKIKFLFQNPDTK QIEAKELEFSNLF AVLHCHISFFAAKVACLDPLFLGNQATASTAASSATTSKAKYTT
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 — SNTG1

Entrez GeneID [54212](#)

GeneBank Accession# [AM393671.1](#)

Protein Accession# [CAL38547.1](#)

Gene Name SNTG1

Gene Alias G1SYN, SYN4

Gene Description syntrophin, gamma 1

Omim ID [608714](#)

Gene Ontology [Hyperlink](#)

Gene Summary The protein encoded by this gene is a member of the syntrophin family. Syntrophins are cytoplasmic peripheral membrane proteins that typically contain 2 pleckstrin homology (PH) domains, a PDZ domain that bisects the first PH domain, and a C-terminal domain that mediates dystrophin binding. This gene is specifically expressed in the brain. Transcript variants for this gene have been described, but their full-length nature has not been determined. [provided by RefSeq]

Other Designations gamma1-syntrophin|syntrophin 4

Disease

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- [Duchenne muscular dystrophy](#)
- [Genetic Predisposition to Disease](#)
- [Inversion](#)
- [Scoliosis](#)
- [Tobacco Use Disorder](#)