

DNAxPAb

Hard-to-Find
Antibody

NAGLU DNAxPab

Catalog # H00004669-W01P

Size 200 ug

Specification

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

Technology [DNAx™ Immune](#)

Immunogen Full-length human DNA

Sequence MEAVAVAAAVGVLLLAGAGGAAGDEAREAAAVRALVARLLGPGPAADFSVSVERALAAKPGLD
TYSLGGGGAARVRVRGSTGVAAAAGLHRYLRDFCGCHVAWSGSQLRLPRPLPAVPGELTEATP
NRYRYQNVCTQSYSFVWWDWARWEREIDWMALNGINLALAWSGQEAWQRVYLALGLTQAEIN
EFFTGP AFLAWGRMG NLHTWDG PLPPSWHIKQLYLQHRVLDQMRSFGMTPVLP AFAGHVPEAV
TRVFPQVNVTKMG SWGHFNCSYSCS FLLAPEDPIFPIIGSLFLRELIFEGTDHIYGADTFNEMQPP
SSEPSYLA AATTAVYEAMTAVDTEAVWLLQGWL FQHQPQFWGPAQIRAVLGAVPRGRLLVLDLF
AESQPVYTRTASFQGPFIWCMLHNF GGNHGLFGALEAVNGGPEAARLFPNSTMVGTGMAPEGI
SQNEVVYSLMAELGWRKDPVPDLAAWVTSFAARRYGVSHPDAGAAWRLLLRSVYNCSGEACR
GHNRSPLVRRPSLQMNTSIWYNRSDVFEAWRLLLTSAPSLATSPAFRYDLLDLTRQAVQELVSLY
YEEARSAYLSKELASLLRAGGVLAYELL PALDEVLASDSRFL LGSWLEQARAAAVSEAEADFYE
QNSRYQLTLWGPEGNILDYANKQLAGLVANYTTPRWRLFLEALVDSVAQGIPFQQHQFDKNVFQL
EQAFVLSKQRYPSQPRGDTVDLAKKIFLKYYPGWVAGSW

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 — NAGLU

Entrez GeneID [4669](#)

GeneBank Accession# [BC053991.1](#)

Protein Accession# [AAH53991.1](#)

Gene Name NAGLU

Gene Alias MPS-IIIB, MPS3B, NAG, UFHSD

Gene Description N-acetylglucosaminidase, alpha-

Omim ID [252920 609701](#)

Gene Ontology [Hyperlink](#)

Gene Summary This gene encodes an enzyme that degrades heparan sulfate by hydrolysis of terminal N-acetyl-D-glucosamine residues in N-acetyl-alpha-D-glucosaminides. Defects in this gene are the cause of mucopolysaccharidosis type IIIB (MPS-IIIB), also known as Sanfilippo syndrome B. This disease is characterized by the lysosomal accumulation and urinary excretion of heparan sulfate. [provided by RefSeq]

Other Designations N-acetyl-alpha-glucosaminidase|alpha-N-acetylglucosaminidase

Pathway

- [Glycosaminoglycan degradation](#)
- [Lysosome](#)
- [Metabolic pathways](#)