

Product datasheet for TP308708

OriGene Technologies, Inc.

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GALNS (NM 000512) Human Recombinant Protein

Product data:

Product Type: Recombinant Proteins

Description: Recombinant protein of human galactosamine (N-acetyl)-6-sulfate sulfatase (GALNS), 20 μg

Species: Human Expression Host: HEK293T

Expression cDNA Clone or AA Sequence: >RC208708 protein sequence Red=Cloning site Green=Tags(s)

MAAVVAATRWWQLLLVLSAAGMGASGAPQPPNILLLLMDDMGWGDLGVYGEPSRETPNLDRMAAEGLLFP NFYSANPLCSPSRAALLTGRLPIRNGFYTTNAHARNAYTPQEIVGGIPDSEQLLPELLKKAGYVSKIVGK WHLGHRPQFHPLKHGFDEWFGSPNCHFGPYDNKARPNIPVYRDWEMVGRYYEEFPINLKTGEANLTQIYL

QEALDFIKRQARHHPFFLYWAVDATHAPVYASKPFLGTSQRGRYGDAVREIDDSIGKILELLQDLHVADN TFVFFTSDNGAALISAPEQGGSNGPFLCGKQTTFEGGMREPALAWWPGHVTAGQVSHQLGSIMDLFTTSL ALAGLTPPSDRAIDGLNLLPTLLQGRLMDRPIFYYRGDTLMAATLGQHKAHFWTWTNSWENFRQGIDFCP GQNVSGVTTHNLEDHTKLPLIFHLGRDPGERFPLSFASAEYQEALSRITSVVQQHQEALVPAQPQLNVCN

WAVMNWAPPGCEKLGKCLTPPESIPKKCLWSH

TRTRPLEQKLISEEDLAANDILDYKDDDDK**V**

Tag: C-Myc/DDK
Predicted MW: 55.4 kDa

Concentration: >0.05 µg/µL as determined by microplate BCA method

Purity: > 80% as determined by SDS-PAGE and Coomassie blue staining

Buffer: 25 mM Tris-HCl, 100 mM glycine, pH 7.3, 10% glycerol

Preparation: Recombinant protein was captured through anti-DDK affinity column followed by conventional

chromatography steps.

Note: For testing in cell culture applications, please filter before use. Note that you may experience

some loss of protein during the filtration process.

Storage: Store at -80°C.

Stability: Stable for 12 months from the date of receipt of the product under proper storage and handling

conditions. Avoid repeated freeze-thaw cycles.



ORIGENE

RefSeq: NP 000503

Locus ID: 2588 **UniProt ID:** P34059

RefSeq Size: 2380

Cytogenetics: 16q24.3 1566 RefSeq ORF:

Synonyms: GalN6S; GALNAC6S; GAS; MPS4A

Summary: This gene encodes N-acetylgalactosamine-6-sulfatase which is a lysosomal exohydrolase

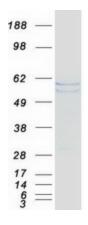
> required for the degradation of the glycosaminoglycans, keratan sulfate, and chondroitin 6sulfate. Sequence alterations including point, missense and nonsense mutations, as well as those that affect splicing, result in a deficiency of this enzyme. Deficiencies of this enzyme lead to

Morquio A syndrome, a lysosomal storage disorder. [provided by RefSeq, Jul 2008]

Protein Families: Druggable Genome

Protein Pathways: Glycosaminoglycan degradation, Lysosome, Metabolic pathways

Product images:



Coomassie blue staining of purified GALNS protein (Cat# TP308708). The protein was produced from HEK293T cells transfected with GALNS cDNA clone (Cat# [RC208708]) using

MegaTran 2.0 (Cat# [TT210002]).