

## Product datasheet for TP308708

### 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)
	<p>MAAVVAATRWWQLLLLVLSAAGMGASGAPQPPNILLLLMDDMGWGDLGVYGEPSRETPNIDRMAAEGLLFP  NFYSANPLCSPSRAALLTGRPLIRNGFYTTNAHARNAYTPQEIVGGIPDSEQLLPELLKKAGYVSKIVGK  WHLGHRPQFHPLKHGFDEWFGSPNCHFGPYDNKARNIPVYRDWEMVGRYEEFPINLKTGEANLTQIYL  QEALDFIKRQARHHPFFLYWAVDATHAPVYASKPFLGTSQRGRYGDVREIDDSIGKILELLQDLHVADN  TFVFFTSNDGAALISAPEQGGSNPFLCGKQTTFEGGMREPALAWWPGHVHTAGQVSHQLGSIMDLFTTSL  ALAGLTPPSDRAIDGLNLLPTLLQGRLMDRPIFYRGDTLMAATLGQHKAFWWTNSWENFRQGIDFCP  GQNVSGVTTHNLEDHTKLPLIFHLGRDPGERFPLSFASAEYQEALSRTSVVQQHQEALVPAQPQLNVCN  WAVMNVWAPPGCEKLGKCLTPPESIPKKCLWSH</p> <p>TRTRPLEQKLISEEDLAANDILDYKDDDDKV</p>
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.



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RefSeq: [NP\\_000503](#)

Locus ID: 2588

UniProt ID: [P34059](#)

RefSeq Size: 2380

Cytogenetics: 16q24.3

RefSeq ORF: 1566

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 6-sulfate. 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]).