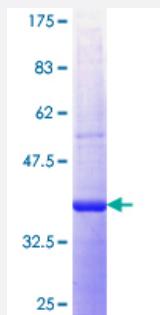


# GNS (Human) Recombinant Protein (Q01)

Catalog # H00002799-Q01

Size 25 ug, 10 ug

## Applications



## Specification

<b>Product Description</b>	Human GNS partial ORF ( NP_002067.1, 455 a.a. - 551 a.a.) recombinant protein with GST-tag at N-terminal.
<b>Sequence</b>	VRTMSALWNLQYCEFDQEVFVEVYNLTADPDQITNIAKTIDPELLGKMNYRLMMLQSCSGPTCR TPGVFDPGYRFDPRLMFSNRGSVRTRRFSKHL
<b>Host</b>	Wheat Germ (in vitro)
<b>Theoretical MW (kDa)</b>	36.41
<b>Preparation Method</b>	<a href="#">in vitro wheat germ expression system</a>
<b>Purification</b>	Glutathione Sepharose 4 Fast Flow
<b>Quality Control Testing</b>	12.5% SDS-PAGE Stained with Coomassie Blue.
<b>Storage Buffer</b>	50 mM Tris-HCl, 10 mM reduced Glutathione, pH=8.0 in the elution buffer.
<b>Storage Instruction</b>	Store at -80°C. Aliquot to avoid repeated freezing and thawing.
<b>Note</b>	Best use within three months from the date of receipt of this protein.

## Applications

- Enzyme-linked Immunoabsorbent Assay
- Western Blot (Recombinant protein)
- Antibody Production
- Protein Array

## Gene Info — GNS

Entrez GeneID	<a href="#">2799</a>
GeneBank Accession#	<a href="#">NM_002076</a>
Protein Accession#	<a href="#">NP_002067.1</a>
Gene Name	GNS
Gene Alias	G6S, MGC21274
Gene Description	glucosamine (N-acetyl)-6-sulfatase
Omim ID	<a href="#">252940 607664</a>
Gene Ontology	<a href="#">Hyperlink</a>
Gene Summary	The product of this gene is a lysosomal enzyme found in all cells. It is involved in the catabolism of heparin, heparan sulphate, and keratan sulphate. Deficiency of this enzyme results in the accumulation of undegraded substrate and the lysosomal storage disorder mucopolysaccharidosis type III D (Sanfilippo D syndrome). Mucopolysaccharidosis type IIID is the least common of the four subtypes of Sanfilippo syndrome. [provided by RefSeq]
Other Designations	N-acetylglucosamine-6-sulfatase glucosamine-6-sulfatase

## Pathway

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