

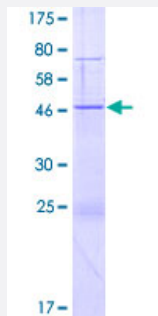
Full-Length

HGSNAT (Human) Recombinant Protein (P01)

Catalog # H00138050-P01

Size 25 ug, 10 ug

Applications



Specification

Product Description	Human HGSNAT full-length ORF (AAH12452.1, 1 a.a. - 206 a.a.) recombinant protein with GST-tag at N-terminal.
Sequence	MALGLCRCFHPRHSMAAFGLFPALPSALNSHPACTCLLDPSTWRPAHVSGPALASSPQILSVFS LGFPGFVNGSCVSRYPDIIFPPGLPPDLPSSVSIFYLQLLCSHGHCITESGILLSFSNWPPSL VPHFLKSPVHCHQIKLSPARSPLSEKPPLTWKHHCLAHILTYSPSRDPHTSFQPPPLHSLPPPP PPHPLVSPPL
Host	Wheat Germ (in vitro)
Theoretical MW (kDa)	48.7
Preparation Method	in vitro wheat germ expression system
Purification	Glutathione Sepharose 4 Fast Flow
Quality Control Testing	12.5% SDS-PAGE Stained with Coomassie Blue.
Storage Buffer	50 mM Tris-HCl, 10 mM reduced Glutathione, pH=8.0 in the elution buffer.
Storage Instruction	Store at -80°C. Aliquot to avoid repeated freezing and thawing.
Note	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 — HGSNAT

Entrez GeneID [138050](#)

GeneBank Accession# [BC012452.1](#)

Protein Accession# [AAH12452.1](#)

Gene Name HGSNAT

Gene Alias DKFZp686G24175, FLJ22242, FLJ32731, HGNAT, MPS3C, TMEM76

Gene Description heparan-alpha-glucosaminide N-acetyltransferase

Omim ID [252930 610453](#)

Gene Ontology [Hyperlink](#)

Gene Summary This gene encodes a lysosomal acetyltransferase, which is one of several enzymes involved in the lysosomal degradation of heparin sulfate. Mutations in this gene are associated with Sanfilippo syndrome C, one type of the lysosomal storage disease mucopolysaccharidosis III, which results from impaired degradation of heparan sulfate. [provided by RefSeq]

Other Designations transmembrane protein 76

Pathway

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