

## **Product datasheet for TA356470**

# **GNS Rabbit Polyclonal Antibody**

**Product data:** 

**Product Type: Primary Antibodies** 

**Applications:** IHC, WB Reactivity: Human Rabbit Host:

Clonality: Polyclonal

Immunogen: The immunogen is a synthetic peptide directed towards the C terminal region of human GNS

Expected reactivity: Cow, Dog, Goat, Guinea Pig, Horse, Human, Mouse, Rabbit, Rat, Specificity:

Zebrafish

Homology: Cow: 100%; Dog: 93%; Goat: 100%; Guinea Pig: 100%; Horse: 100%; Human:

100%; Mouse: 100%; Rabbit: 100%; Rat: 100%; Zebrafish: 85%

Formulation: Liquid. Purified antibody supplied in 1x PBS buffer with 0.09% (w/v) sodium azide and 2%

sucrose.

Note that this product is shipped as lyophilized powder to China customers.

Concentration: lot specific

**Purification: Affinity Purified** Conjugation: Unconjugated

Storage: For short term use, store at 2-8°C up to 1 week. For long term storage, store at -20°C in small

aliquots to prevent freeze-thaw cycles.

Stability: Shelf life: one year from despatch.

**Predicted Protein Size:** 62kDa

Gene Name: glucosamine (N-acetyl)-6-sulfatase

Database Link: NP 002067

Entrez Gene 2799 Human

P15586



OriGene Technologies, Inc. 9620 Medical Center Drive, Ste 200

CN: techsupport@origene.cn

Rockville, MD 20850, US Phone: +1-888-267-4436 https://www.origene.com techsupport@origene.com EU: info-de@origene.com



#### Background:

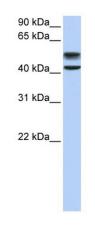
GNS is a lysosomal enzyme found in all cells. It is involved in the catabolism of heparin, heparin sulphate, and keratan sulphate. Deficiency of this enzyme results in the accumulation of undegraded substrate and the lysosomal storage disorder ucopolysaccharidosis type IIID (Sanfilippo D syndrome). Mucopolysaccharidosis type IIID is the least common of the four subtypes of Sanfilippo syndrome. 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 IIID (Sanfilippo D syndrome). Mucopolysaccharidosis type IIID is the least common of the four subtypes of Sanfilippo syndrome.

Synonyms: G6S; Glucosamine-6-sulfatase; MGC21274; N-acetylglucosamine-6-sulfatase

**Protein Families:** Druggable Genome, Transmembrane

**Protein Pathways:** Glycosaminoglycan degradation, Lysosome, Metabolic pathways

### **Product images:**

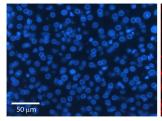


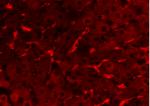
WB Suggested Anti-GNS Antibody Titration: 0.2-1

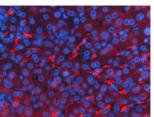
ug/ml

ELISA Titer: 1:312500

Positive Control: Human Liver







Rabbit Anti-GNS Antibody
Catalog Number: TA356470

Formalin Fixed Paraffin Embedded Tissue:

**Human Liver Tissue** 

Observed Staining: Cytoplasm in Kupffer cells in

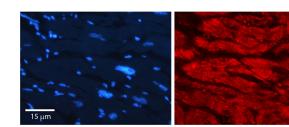
sinusoids

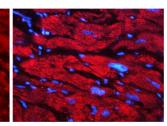
Primary Antibody Concentration: 1:100
Other Working Concentrations: N/A
Secondary Antibody: Donkey anti-Rabbit-Cy3

Secondary Antibody Concentration: 1:200

Magnification: 20X Exposure Time: 0.5–2.0 sec







Rabbit Anti-GNS Antibody
Catalog Number: TA356470
Formalin Fixed Paraffin Embedded Tissue:
Human heart Tissue
Observed Staining: Cytoplasmic
Primary Antibody Concentration: 1:100
Other Working Concentrations: N/A
Secondary Antibody: Donkey anti-Rabbit-Cy3
Secondary Antibody Concentration: 1:200

Magnification: 20X Exposure Time: 0.5–2.0 sec