

### **Product datasheet for TA397070S**

#### 9620 Medical Center Drive, Ste 200 Rockville, MD 20850, US Phone: +1-888-267-4436 https://www.origene.com

OriGene Technologies, Inc.

https://www.origene.com techsupport@origene.com EU: info-de@origene.com CN: techsupport@origene.cn

# GDF15 Rat Monoclonal Antibody [Clone ID: 6D12.H10.E4]

#### **Product data:**

**Product Type:** Primary Antibodies

Clone Name: 6D12.H10.E4
Applications: ELISA, WB

Recommended Dilution: WB: 1:1,000

**ELISA**: 1:2,000

Reactivity: Human, Mouse

**Host:** Rat

Isotype: lgG2a, lambda
Clonality: Monoclonal

Immunogen: This Protein-A purified antibody was prepared by repeated immunizations with an MBP-

tagged recombinant protein produced in E.coli corresponding to C-term mouse NAG-1

protein. Cross reactivity to MBP is negative.

**Specificity:** This product was purified from concentrated tissue culture supernatant Protein G

chromatography. This antibody reacts with the C-terminus of endogenous NAG-1 protein from mouse tissues. A BLAST analysis suggests reactivity with NAG-1 from chimpanzee and macaque based on a 100% homology. Partial reactivity is expected against rat based on an 86% homology with the immunizing sequence. Cross-reactivity with NAG-1 from other

sources has not been determined.

**Formulation:** 0.02 M Potassium Phosphate, 0.15 M Sodium Chloride, pH 7.2

**Concentration:** 1.0 mg/mL - lot specific

Conjugation: Unconjugated

Storage: Store vial at -20° C or below prior to opening. This vial contains a relatively low volume of

reagent (25  $\mu$ L). To minimize loss of volume dilute 1:10 by adding 225  $\mu$ L of the buffer stated above directly to the vial. Recap, mix thoroughly and briefly centrifuge to collect the volume at the bottom of the vial. Use this intermediate dilution when calculating final dilutions as recommended below. Store the vial at -20°C or below after dilution. Avoid cycles of freezing

and thawing.

**Stability:** Expiration date is three (3) months from date of receipt.

Database Link: Q99988





Background:

Non-steroidal anti-inflammatory drug (NSAID) activated gene (NAG-1) is a member of the transforming growth factor-beta (TGF-beta) superfamily. NAG-1 is also known as Macrophage Inhibitory Cytokine-1 (MIC-1), Growth Differentiation Factor 15 (GDF15), Placental Bone Morphogenetic Protein (PLAB), or Prostate Derived Factor (PDF). NAG-1 is expressed in human placenta, prostate and colon. It possesses antitumorigenic and proapoptotic activities. NAG-1 expression is dramatically increased in inflammation, injury and malignancy. Increase of NAG-1 expression is a feature of many cancers including breast, colon, pancreas and prostate. In a number of studies, NAG-1 expression was increased by a number of NSAIDs. This increase in expression may correlate with the chemopreventive effect NSAIDs seem to have with certain cancers. NAG-1 expression is also induced by PPAR gamma ligands and by several dietary compounds such as conjugated linoleic acids (CLAs), naturally occurring fatty acids in ruminant food products, indoles, epicatechin gallate, and genistein. Induced expression of NAG-1 results in stimulation of apoptosis and inhibition of cell growth. Inhibition of NAG-1 induced expression by small interference RNA (siRNA) results in repression of induced apoptosis. NAG-1 expression is regulated by a numbers of transcription factors such as ERG-1 and Sp1. EGR-1 may be necessary for NSAID-induced NAG-1 expression. The study of expression of NAG-1 proteins, including variants, is important to define their potential role as serum biomarkers for cancer diagnosis, treatment monitoring, epidemiology study, and nutrition surveys.

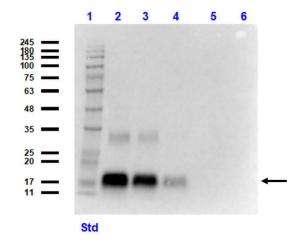
Synonyms:

rat anti-NAG1 antibody, NAG-1, GDF15, MIC-1, nonsteroidal anti-inflammatory drug-activated gene, NSAID-activated gene 1 protein, growth differentiation factor 15, macrophage inhibitory compound 1, prostate-derived factor

Note:

This Protein G Anti-NAG1 purified antibody has been tested by ELISA and western blot for mouse NAG-1 protein. Specific conditions for reactivity should be optimized by the end user. Expect bands in Western blots of native protein of approximately ~58kDa using the appropriate cell lysate or extract.

# **Product images:**



Western Blot of Rat Anti-NAG1 C-term Antibody. Lane 1: Opal Prestained Molecular Weight Marker (p/n MB-210-0500). Lane 2: HeLa Lysate  $(W09-000-364)[10\mu L] + recomb. NAG1[0.05\mu g].$ Lane 3: HeLa Lysate (W09-000-364) [10µL] + recomb. NAG1 [0.02µg]. Lane 4: HeLa Lysate  $(W09-000-364)[10\mu L] + recomb. NAG1[0.01\mu g].$ Lane 5: HeLa Lysate (W09-000-364) [10µL]. Lane 6: HeLa Lysate (W09-000-364) [10µL] + MBP (000-001-385-1) [0.05µg]. Primary Antibody: Anti-NAG1-C-term at 1µg/mL overnight at 2-8°C. Secondary Antibody: Goat Anti-Rat IgG HRP (612-103-120) at 1:40,000 for 30 mins at RT. Block: BlockOut Buffer (MB-073) for 30 mins at RT. Predicted MW: 17kDa. Observed MW: ~17kDa. Exposure: 10 sec. Gel: 12%.