

## **Product Datasheet**

# GNG2 Antibody (monoclonal, 7C13) (orb865586)

Catalog Number orb865586

**Category** Antibodies

**Description** Anti-GNG2 Antibody (monoclonal, 7C13). Tested in IF, ICC, WB applications. This

antibody reacts with Human, Mouse, Rat.

**Clonality** Monoclonal

Species/Host Mouse

**Isotype** Mouse IgG2b

**Conjugation** Unconjugated

**Reactivity** Human, Mouse, Rat

Form/Appearance Lyophilized

**Concentration** Adding 0.2 ml of distilled water will yield a concentration of 500 μg/ml.

**Purification** Immunogen affinity purified.

**Immunogen** E.coli-derived human GNG2 recombinant protein (Position: A2-D48).

UniProt ID P59768

MW 12 kDa

**Tested applications** ICC, IF, WB

**Application notes** Western blot, 0.25-0.5 μg/ml, Mouse, Rat

Immunocytochemistry/Immunofluorescence, 5 µg/ml, Human. Adding 0.2 ml of

distilled water will yield a concentration of 500 µg/ml

## **Biorbyt Ltd.**

7 Signet Court, Swann Road Cambridge CB5 8LA United Kingdom

Email: <u>info@biorbyt.com</u>, <u>support@biorbyt.com</u> Phone: +44 (0)1223 859353 | Fax: +1 (415) 651-8558

## **Biorbyt LLC**

68 TW Alexander Drive Research Triangle Park

Durham NC 27713 United States

Email: <u>info@biorbyt.com</u>, <u>support@biorbyt.com</u> Phone: +1 (415) 906-5211 | Fax: +1 (415) 651-8558





**Cross Reactivity** No cross-reactivity with other proteins.

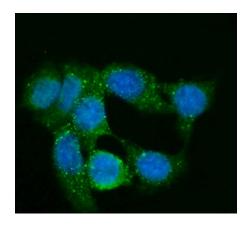
**Antibody Type** Primary Antibody

Clone Number 7C13

**Storage** Maintain refrigerated at 2-8°C for up to 2 weeks. For long term storage store at -

20°C in small aliquots to prevent freeze-thaw cycles.

**Note** For research use only



IF analysis of GNG2 using anti-GNG2 antibody. GNG2 was detected in an immunocytochemical section of Caco-2 cells. Enzyme antigen retrieval was performed using IHC enzyme antigen retrieval reagent for 15 mins. The cells were blocked with 10% goat serum. And then incubated with 5 μg/mL mouse anti-GNG2 Antibody overnight at 4°C. DyLight® 488 Conjugated Goat Anti-Mouse IgG was used as secondary antibody at 1:100 dilution and incubated for 30 minutes at 37°C. The section was counterstained with DAPI. Visualize using a fluorescence microscope and filter sets appropriate for the label used.

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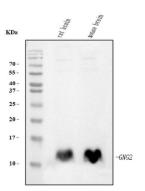
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Western blot analysis of GNG2 using anti-GNG2 antibody. Electrophoresis was performed on a 5-20% SDS-PAGE gel at 70V (Stacking gel) / 90V (Resolving gel) for 2-3 hours. The sample well of each lane was loaded with 30 ug of sample under reducing conditions. Lane 1: rat brain tissue lysates, Lane 2: mouse brain tissue lysates. After electrophoresis, proteins were transferred to a nitrocellulose membrane at 150 mA for 50-90 minutes. Blocked the membrane with 5% non-fat milk/TBS for 1.5 hour at RT. The membrane was incubated with mouse anti-GNG2 antigen affinity purified monoclonal antibody at 0.5 µg/mL overnight at 4°C, then washed with TBS-0.1% Tween 3 times with 5 minutes each and probed with a goat anti-mouse IgG-HRP secondary antibody at a dilution of 1:10000 for 1.5 hour at RT. The signal is developed using an Enhanced Chemiluminescent detection (ECL) kit with Tanon 5200 system. A specific band was detected for GNG2 at approximately 12 kDa. The expected band size for GNG2 is at 8 kDa.

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