Product Datasheet

DGAT2 Antibody NBP1-71700SS

Unit Size: 0.025 ml

Store at 4C short term. Aliquot and store at -20C long term. Avoid freeze-thaw cycles.

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Publications: 2

Protocols, Publications, Related Products, Reviews, Research Tools and Images at: www.novusbio.com/NBP1-71700

Updated 6/15/2014 v.20.1

NBP1-71700SS

0.025 ml
1 mg/ml
Store at 4C short term. Aliquot and store at -20C long term. Avoid freeze-thaw cycles.
Polyclonal
0.05% Sodium Azide
Immunogen affinity purified
PBS, 30% glycerol
Rabbit
84649
DGAT2
Human, Mouse, Yeast
Human, yeast and mouse. Immunogen has 100% identity to rat, bovine and porcine, and 86% identity to Xenopus.
A synthetic peptide made to an internal region of the human DGAT2 protein (within residues 100-200). [Swiss-Prot Q96PD7]
Western Blot
Western Blot 1 ug/ml
This DGAT2 antibody is useful for Western blot, where a band is seen ~43 kDa.
NBP1-71700] - Analysis of DGAT2 in 250> 150> 100> 75>



15> 10>

Publications

Zeng T, Zhang CL, Song FY et al. CMZ Reversed Chronic Ethanol-Induced Disturbance of PPAR-a Possibly by Suppressing Oxidative Stress and PGC-1a Acetylation, and Activating the MAPK and GSK3b Pathway PLoS ONE. 2014 Jun 04 [PMID: 24892905] (WB, Mouse)

Details:

DGAT2 antibody used for WB in specific pathogen-free/SPF male Kun-Ming mice treated or not with chronic ethanol and/or chlormethiazole (CMZ). DGAT2 was studied as rate-limiting enzyme of G synthesis and its levels were found increased in ethanol and CMZ/ethanol group mice liver. (Figure 9A)

Xu N, Zhang SO, Cole RA, McKinney SA, Guo F, Haas JT, Bobba S, Farese RV Jr, Mak HY. The FATP1-DGAT2 complex facilitates lipid droplet expansion at the ER-lipid droplet interface J Cell Biol 2012 Aug 27 [PMID: 22927462] (WB, Yeast)



Procedures

Western Blot protocolspecific for DGAT2 antibody (NBP1-71700)

Western Blot Protocol

- 1. Perform SDS-PAGE on samples to be analyzed, loading 40 ug of total protein per lane.
- 2. Transfer proteins to membrane according to the instructions provided by the manufacturer of the membrane and transfer apparatus.
- 3. Stain according to standard Ponceau S procedure (or similar product) to assess transfer success, and mark molecular weight standards where appropriate.
- 4. Rinse the blot.
- 5. Block the membrane using standard blocking buffer for at least 1 hour.
- 6. Wash the membrane in wash buffer three times for 10 minutes each.
- 7. Dilute primary antibody in blocking buffer and incubate 1 hour at room temperature.
- 8. Wash the membrane in wash buffer three times for 10 minutes each.
- 9. Apply the diluted HRP conjugated secondary antibody in blocking buffer (as per manufacturers instructions) and incubate 1 hour at room temperature.
- 10. Wash the blot in wash buffer three times for 10 minutes each (this step can be repeated as required to reduce background).
- 11. Apply the detection reagent of choice in accordance with the manufacturers instructions.

*Note: Tween-20 can be added to the blocking or antibody dilution buffer at a final concentration of 0.05-0.2%.





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Limitations

This product is for research use only and is not approved for use in humans or in clinical diagnosis. Primary Antibodies are guaranteed for 1 year from date of receipt.

For more information on our guarantee, please visit www.novusbio.com/guarantee.

