

# Product Datasheet

## SREBP1 Antibody (2A4) - BSA Free NB600-582

Unit Size: 0.1 ml

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

[www.novusbio.com](http://www.novusbio.com)



[technical@novusbio.com](mailto:technical@novusbio.com)

**Publications: 73**

Protocols, Publications, Related Products, Reviews, Research Tools and Images at:  
[www.novusbio.com/NB600-582](http://www.novusbio.com/NB600-582)

Updated 9/9/2025 v.20.1

**Earn rewards for product  
reviews and publications.**

Submit a publication at [www.novusbio.com/publications](http://www.novusbio.com/publications)

Submit a review at [www.novusbio.com/reviews/destination/NB600-582](http://www.novusbio.com/reviews/destination/NB600-582)



**NB600-582**

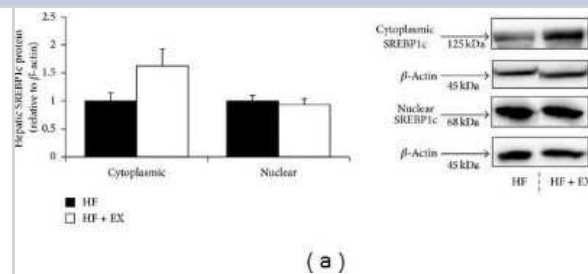
SREBP1 Antibody (2A4) - BSA Free

Product Information	
Unit Size	0.1 ml
Concentration	1 mg/ml
Storage	Store at 4C short term. Aliquot and store at -20C long term. Avoid freeze-thaw cycles.
Clonality	Monoclonal
Clone	2A4
Preservative	0.05% Sodium Azide
Isotype	IgG1 Kappa
Purity	Protein G purified
Buffer	PBS
Product Description	
Description	Novus Biologicals Knockout (KO) Validated Mouse SREBP1 Antibody (2A4) - BSA Free (NB600-582) is a monoclonal antibody validated for use in IHC, WB and Simple Western. Anti-SREBP1 Antibody: Cited in 69 publications. All Novus Biologicals antibodies are covered by our 100% guarantee.
Host	Mouse
Gene ID	6720
Gene Symbol	SREBF1
Species	Human, Mouse, Rat, Canine, Chicken, Hamster, Monkey, Golden Syrian Hamster
Reactivity Notes	Canine reactivity reported in scientific literature (PMID: 23720350). Hamster reactivity reported in scientific literature (PMID: 24393244). Chicken reactivity reported in multiple pieces scientific literature. Monkey reactivity reported in scientific literature (PMID: 26437365).
Immunogen	6 His-tag fusion protein of human SREBP1 corresponding to amino acids 301-407. [UniProt# P36956]
Product Application Details	
Applications	Western Blot, Simple Western, Immunohistochemistry-Frozen, Knockout Validated
Recommended Dilutions	Western Blot 1-2ug/ml, Simple Western 1:12.5, Immunohistochemistry-Frozen, Knockout Validated
Application Notes	<p>This SREBP1 (clone 2A4) antibody is useful for WB where a band can be seen at 125 kDa (precursor) and additional bands may be seen at 60-70 kDa (cleaved).</p> <p>In Simple Western only 10 - 15 uL of the recommended dilution is used per data point.</p> <p>See <a href="#">Simple Western Antibody Database</a> for Simple Western validation: Tested in HeLa lysate 1.0 mg/mL, separated by Size, antibody dilution of 1:12.5, apparent MW was 156 kDa. Separated by Size-Wes, Sally Sue/Peggy Sue. The use of this antibody in IHC paraffin embedded tissue has been questionable.</p>

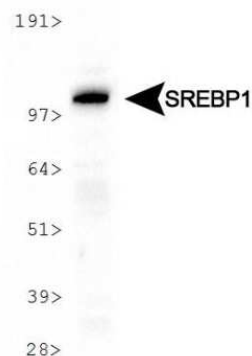


## Images

Western Blot: SREBP1 Antibody (2A4) [NB600-582] - Hepatic SREBP1c and SREBP2 protein abundance in C57BL/6 mice assigned to a high-fat (HF) or a high-fat/exercise (HF + EX) group for 8 weeks. Cytosolic and nuclear SREBP1c abundance. Image collected and cropped by CiteAb from the following publication (<https://www.hindawi.com/journals/jl/2013/908048/>), licensed under a CC-BY license.



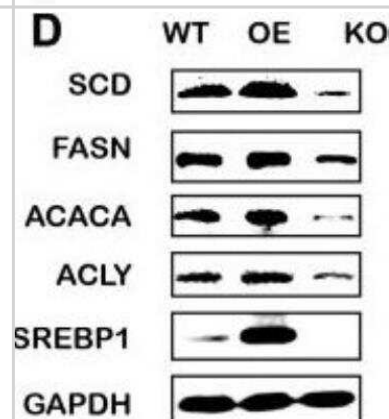
Western Blot: SREBP1 Antibody (2A4) [NB600-582] - Analysis of whole cell lysate from HeLa cells showing a single specific band for the expression of SREBP1 precursor protein (~120 kDa).



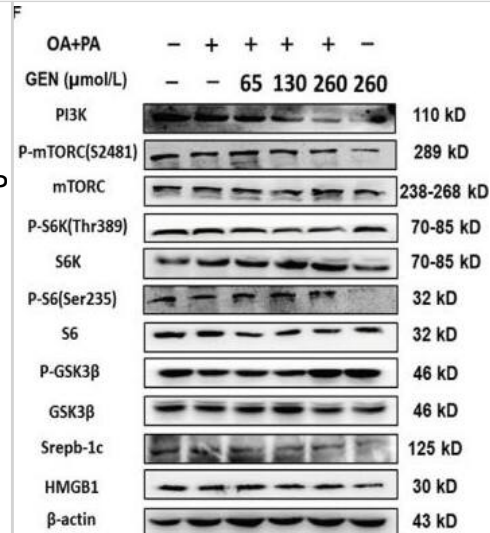
Simple Western: SREBP1 Antibody (2A4) [NB600-582] - Lane view shows a specific band for SREBP1 in 1.0 mg/ml of HeLa lysate. This experiment was performed under reducing conditions using the 12-230 kDa separation system.



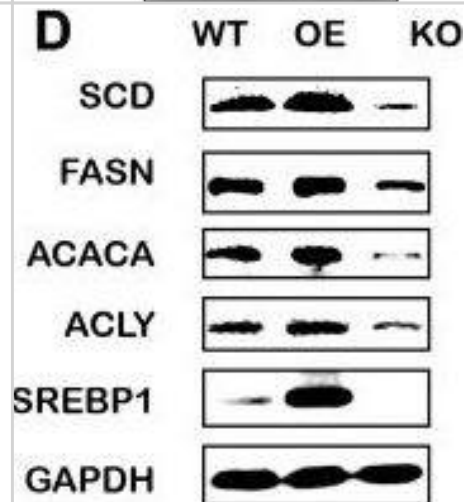
Western Blot: SREBP1 Antibody (2A4) [NB600-582] - KO and OE of SREBP1 influenced lipid metabolism in Bel-7402 cells. The protein level of lipid-associated genes in SREBP1-KO, SREBP1-OE, and WT Bel-7402 cells. Data were expressed as mean  $\pm$  standard error of the mean. \* $P < 0.05$ , \*\* $P < 0.01$ , and \*\*\* $P < 0.001$  vs. WT group, respectively.



Western Blot: SREBP1 Antibody (2A4) - BSA Free [NB600-582] - Effect of GEN on the expression of Nrf2-related signalling proteins in OA (660  $\mu\text{mol/L}$ ) & PA (330  $\mu\text{mol/L}$ ) induced HepG2 cells. GEN was added into HepG2 cells prior 1 h to stimulation of OA & PA for 18 h. A-C, Protein expression of Nrf2, PPAR $\alpha$ , PPAR $\gamma$  in nucleus & HO-1 in cytoplasm was detected by Western blot. (D & E) Protein expression of P-ACC, ACC, P-AKT, AKT, P-AMPK $\alpha$ , AMPK $\alpha$ , P-AMPK $\beta$  & AMPK $\beta$  was detected by Western blot. F-H, Protein expression of PI3K, P-mTORC, mTORC, P-S6K, S6K, P-S6, S6, SREBP-1c & HMGB1 was detected by Western blot. The similar results were collected from three dependent experiments. All data were expressed by mean  $\pm$  SEM (n = 5 in each group). ##P < .01 vs Control Group; \*P < .05 & \*\*P < .01 vs OA & PA Group Image collected & cropped by CiteAb from the following publication (<https://pubmed.ncbi.nlm.nih.gov/32293113>), licensed under a CC-BY license. Not internally tested by Novus Biologicals.



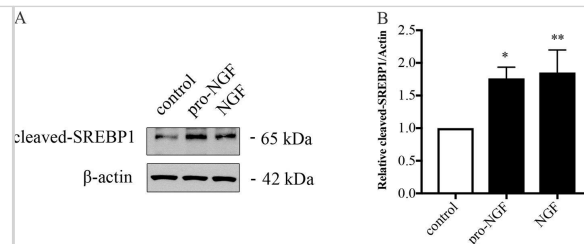
Western Blot: SREBP1 Antibody (2A4) - BSA Free [NB600-582] - KO & OE of SREBP1 influenced lipid metabolism in Bel-7402 cells. (A) Total cellular TG content of SREBP1-KO, SREBP1-OE, & WT Bel-7402 cells. (B) HPLC-MS determination of the SFAs to MUFAs ratios (palmitic acid to palmitoleic acid, 16:0 to 16:1) (stearic acid to oleic acid, 18:0 to 18:1) as well as the content of 16:0, 16:1, 18:0, & 18:1 in SREBP1-KO, SREBP1-OE, & WT Bel-7402 cells. (C) The transcript level of lipid-associated genes in SREBP1-KO, SREBP1-OE, & WT Bel-7402 cells. (D & E) The protein level of lipid-associated genes in SREBP1-KO, SREBP1-OE, & WT Bel-7402 cells. Data were expressed as mean  $\pm$  standard error of the mean. \*P < 0.05, \*\*P < 0.01, & \*\*\*P < 0.001 vs. WT group, respectively. Image collected & cropped by CiteAb from the following publication (<https://pubmed.ncbi.nlm.nih.gov/31297058>), licensed under a CC-BY license. Not internally tested by Novus Biologicals.



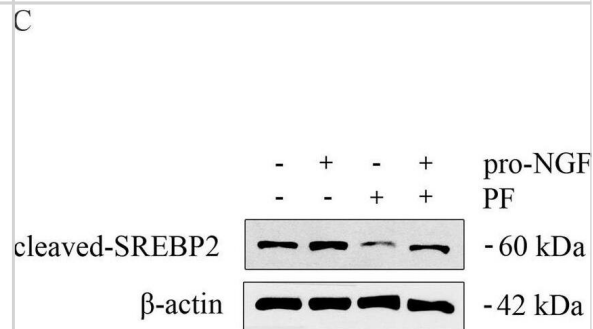
Western Blot: SREBP1 Antibody (2A4) - BSA Free [NB600-582] - Construction of knockout (KO) & overexpression (OE) cell lines. (A) The sequencing results of KO positive clone cells; (B) The messenger RNA (mRNA) level of SREBP1 in wild type (WT) & OE Bel-7402 cells. Data were expressed as mean  $\pm$  standard error of the mean. \*\*\*P < 0.001 vs. WT group; (C & D) The protein level of SREBP1 in WT, OE, & KO Bel-7402 cells. Data were expressed as mean  $\pm$  standard error of the mean. \*\*\*P < 0.001 vs. WT group. Image collected & cropped by CiteAb from the following publication (<https://pubmed.ncbi.nlm.nih.gov/31297058>), licensed under a CC-BY license. Not internally tested by Novus Biologicals.



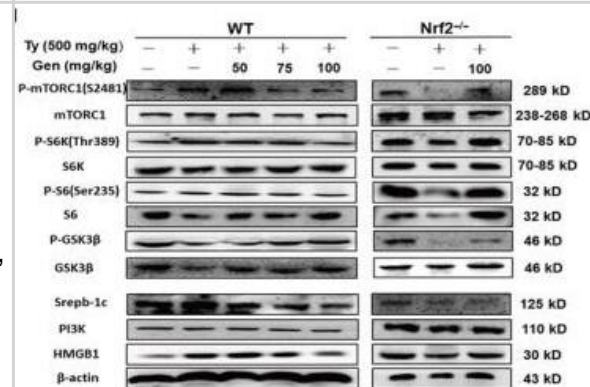
Western Blot: SREBP1 Antibody (2A4) - BSA Free [NB600-582] - p75NTR stimulation activated SREBP1 & SREBP2 in Huh7 hepatocyte cells. Human Huh7 cells were stimulated with 10 ng/ml pro-NGF or 50 ng/ml NGF for 16 h followed by immunoblotting using antibodies to detect the presence of cleaved/activated SREBP1 & SREBP2. a SREBP1, immunoblots & b quantification. Values are mean  $\pm$  SD, n = 3. \*\*P < 0.01 & \*p < 0.05 for treated vs. control cells. c, d SREBP2. Cells were treated with 10 ng/ml pro-NGF in the absence or presence of 0.5  $\mu$ M PF429242 (PF), which is a selective inhibitor of the Site 1 protease (S1P). c Immunoblots & d quantification. PF decreased the amount of processed SREBP2 in these cells; however, addition of pro-NGF increased the relative level of cleaved SREBP2. Values are mean  $\pm$  SD, n = 3. \*p < 0.05 for treated vs. the corresponding control Image collected & cropped by CiteAb from the following publication (<https://pubmed.ncbi.nlm.nih.gov/31296846>), licensed under a CC-BY license. Not internally tested by Novus Biologicals.



Western Blot: SREBP1 Antibody (2A4) - BSA Free [NB600-582] - p75NTR stimulation activated SREBP1 & SREBP2 in Huh7 hepatocyte cells. Human Huh7 cells were stimulated with 10 ng/ml pro-NGF or 50 ng/ml NGF for 16 h followed by immunoblotting using antibodies to detect the presence of cleaved/activated SREBP1 & SREBP2. a SREBP1, immunoblots & b quantification. Values are mean  $\pm$  SD, n = 3. \*\*P < 0.01 & \*p < 0.05 for treated vs. control cells. c, d SREBP2. Cells were treated with 10 ng/ml pro-NGF in the absence or presence of 0.5  $\mu$ M PF429242 (PF), which is a selective inhibitor of the Site 1 protease (S1P). c Immunoblots & d quantification. PF decreased the amount of processed SREBP2 in these cells; however, addition of pro-NGF increased the relative level of cleaved SREBP2. Values are mean  $\pm$  SD, n = 3. \*p < 0.05 for treated vs. the corresponding control Image collected & cropped by CiteAb from the following publication (<https://pubmed.ncbi.nlm.nih.gov/31296846>), licensed under a CC-BY license. Not internally tested by Novus Biologicals.



Western Blot: SREBP1 Antibody (2A4) - BSA Free [NB600-582] - Effect of GEN on the protein expression of Nrf2, ACC & mTORC signalling pathways in tyloxapol-induced mice. GEN (50, 75, 100 mg/kg) was administered to WT or Nrf2<sup>-/-</sup> mice prior 1 h to stimulation of tyloxapol (500 mg/kg) for 18 h. A-C, Protein expression of Nrf2, PPAR $\alpha$ , PPAR $\gamma$  in nucleus & HO-1 in cytoplasm of WT mice was detected by Western blot. (D & E) Protein expression of Nrf2, PPAR $\alpha$  & PPAR $\gamma$  in Nrf2<sup>-/-</sup> mice was detected by Western blot. F-H, Protein expression of P-ACC, ACC, P-AKT, AKT, P-AMPK $\alpha$ , AMPK $\alpha$ , P-AMPK $\beta$ , AMPK $\beta$  in WT & Nrf2<sup>-/-</sup> mice was detected by Western blot. I-M, Protein expression of PI3K, P-mTORC, mTORC, P-S6K, S6K, P-S6, S6, SREBP-1c & HMGB1 WT & Nrf2<sup>-/-</sup> mice was detected by Western blot. The similar results were collected from three dependent experiments. All data were expressed by mean  $\pm$  SEM (n = 5 in each group). ###P < .01 vs Control group; \*P < .05 & \*\*P < .01 vs Tyloxapol group Image collected & cropped by CiteAb from the following publication (<https://pubmed.ncbi.nlm.nih.gov/32293113>), licensed under a CC-BY license. Not internally tested by Novus Biologicals.



## Publications

Batayneh Z, Pan X, Gopaju R et al. Loss of hepatic carboxylesterase 3 (CES3) prevents the development of MASLD in mice. *Journal of lipid research* 2025-08-25 [PMID: 40865609]

Srinivasan M, Kota S, Bhopale K et al. Dysregulated hepatic alcohol metabolism: a key factor involved in the pathogenesis of alcohol-associated liver disease *American Journal of Physiology - Gastrointestinal and Liver Physiology* 2025-03-01 [PMID: 39907561]

Chiu CH, Chang CC, Lin JJ et al. Styrylpyrones from *Phellinus linteus* Mycelia Alleviate Non-Alcoholic Fatty Liver by Modulating Lipid and Glucose Metabolic Homeostasis in High-Fat and High-Fructose Diet-Fed Mice *Antioxidants (Basel)* 2022-04-30 [PMID: 35624762] (Western Blot, Hamster)

L Opazo-Ríos, M Soto-Catal, I Lázaro, A Sala-Vila, L Jiménez-Ca, M Orejudo, JA Moreno, J Egido, S Mas-Fontao Meta-Inflammation and De Novo Lipogenesis Markers Are Involved in Metabolic Associated Fatty Liver Disease Progression in BTBR ob/ob Mice *International Journal of Molecular Sciences*, 2022-04-02;23(7):. 2022-04-02 [PMID: 35409324] (Western Blot, Hamster)

Srinivasan MP, Bhopale KK, Amer SM et al. Linking Dysregulated AMPK Signaling and ER Stress in Ethanol-Induced Liver Injury in Hepatic Alcohol Dehydrogenase Deficient Deer Mice *Biomolecules* 2019-10-02 [PMID: 31581705] (Western Blot, Hamster)

Qiao P, Jia Y, Ma A et al. Dapagliflozin protects against nonalcoholic steatohepatitis in db/db mice *Frontiers in Pharmacology* 2022-08-19 [PMID: 36059948] (Western Blot, Hamster)

Low JY, Brennen WN, Meeker AK et al. Stromal CAVIN1 Controls Prostate Cancer Microenvironment and Metastasis by Modulating Lipid Distribution and Inflammatory Signaling *Molecular Cancer Research* 2020-09-01 [PMID: 32493699] (Western Blot, Hamster)

Thottakkattumana Parameswaran, V;Hild, C;Eichner, G;Ishaque, B;Rickert, M;Steinmeyer, J; Interleukin-1 Induces the Release of Lubricating Phospholipids from Human Osteoarthritic Fibroblast-like Synoviocytes *International journal of molecular sciences* [PMID: 35269552]

Balboni N, Babini G, Poeta E et Al. Transcriptional and metabolic effects of aspartate-glutamate carrier isoform 1 (AGC1) downregulation in mouse oligodendrocyte precursor cells (OPCs) *Cell Mol Biol Lett* 2024-03-29 [PMID: 38553684]

NL Cianciola, S Chung, D Manor, CR Carlin Adenovirus modulates Toll-like receptor 4 signaling by reprogramming ORP1L-VAP protein contacts for cholesterol transport from endosomes to the endoplasmic reticulum *J. Virol*, 2017-02-28;0(0):. 2017-02-28 [PMID: 28077646]

Cassim Bawa FN, Hu S, Gopaju R et Al. Adipocyte retinoic acid receptor ? prevents obesity and steatohepatitis by regulating energy expenditure and lipogenesis *Obesity (Silver Spring)* 2024-01-03 [PMID: 37873741]

Hu S, Cassim Bawa FN, Zhu Y et al. Loss of adipose ATF3 promotes adipose tissue lipolysis and the development of MASH *Communications Biology* 2024-10-10 [PMID: 39390075]

More publications at <http://www.novusbio.com/NB600-582>

## Procedures

### Western Blot Protocol Specific for NB600-582: SREBP1 Antibody (2A4)

#### Western Blot Protocol

1. Perform SDS-PAGE on samples to be analyzed, loading 20 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 overnight at 4C.
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%.

\*The above information is only intended as a guide. The researcher should determine what protocol best meets their needs. Please follow safe laboratory procedures.





### **Novus Biologicals USA**

10730 E. Briarwood Avenue  
Centennial, CO 80112  
USA  
Phone: 303.730.1950  
Toll Free: 1.888.506.6887  
Fax: 303.730.1966  
nb-customerservice@bio-techne.com

### **Bio-Techne Canada**

21 Canmotor Ave  
Toronto, ON M8Z 4E6  
Canada  
Phone: 905.827.6400  
Toll Free: 855.668.8722  
Fax: 905.827.6402  
canada.inquires@bio-techne.com

### **Bio-Techne Ltd**

19 Barton Lane  
Abingdon Science Park  
Abingdon, OX14 3NB, United Kingdom  
Phone: (44) (0) 1235 529449  
Free Phone: 0800 37 34 15  
Fax: (44) (0) 1235 533420  
info.EMEA@bio-techne.com

### **General Contact Information**

www.novusbio.com  
Technical Support: nb-technical@bio-techne.com  
Orders: nb-customerservice@bio-techne.com  
General: novus@novusbio.com

### **Products Related to NB600-582**

---

NB800-PC9	HeLa Nuclear Cell Lysate
NBP2-33376H	Blue Marker Antibody (6F4-F6) [HRP]
HAF007	Goat anti-Mouse IgG Secondary Antibody [HRP]
NB7539	Goat anti-Mouse IgG (H+L) Secondary Antibody [HRP]
NBP1-43319-0.5mg	Mouse IgG1 Kappa Isotype Control (P3.6.2.8.1)

---

### **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 100% guarantee, please visit [www.novusbio.com/guarantee](http://www.novusbio.com/guarantee)

Earn gift cards/discounts by submitting a review: [www.novusbio.com/reviews/submit/NB600-582](http://www.novusbio.com/reviews/submit/NB600-582)

Earn gift cards/discounts by submitting a publication using this product:  
[www.novusbio.com/publications](http://www.novusbio.com/publications)

