

Product Datasheet

GAPDH Antibody (13H12) - BSA Free NBP2-27103

Unit Size: 0.1 mg

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

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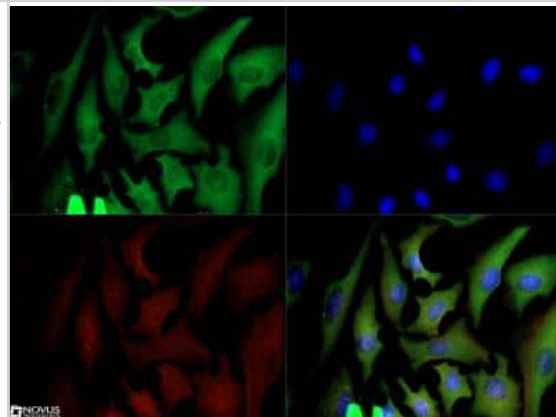
NBP2-27103

GAPDH Antibody (13H12) - BSA Free

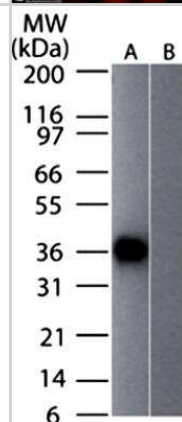
Product Information	
Unit Size	0.1 mg
Concentration	1.0 mg/ml
Storage	Store at 4C short term. Aliquot and store at -20C long term. Avoid freeze-thaw cycles.
Clonality	Monoclonal
Clone	13H12
Preservative	0.02% Sodium Azide
Isotype	IgG1 Kappa
Purity	Protein G purified
Buffer	PBS
Target Molecular Weight	36 kDa
Product Description	
Host	Mouse
Gene ID	2597
Gene Symbol	GAPDH
Species	Human, Mouse, Rat, Drosophila, Monkey, Primate, Sheep
Reactivity Notes	Based upon 91% sequence similarity with immunogen, this antibody is predicted to react with Guinea Pig, Sheep, Squirrel, Porcine/Pig, Ferret, Canine/Dog/Cat, Bovine, Reptile / Rattlesnake and several other species. Immunogen shows 82% similarity to Xenopus and Zebrafish. Rat, sheep, and monkey reactivity reported in scientific literature (PMID: 24796753, PMID: 27618403, and PMID: 24462973 respectively).
Marker	Cytosolic Marker
Immunogen	Amino acids between 275 and 325 of glyceraldehyde 3-phosphate dehydrogenase protein were used as the immunogen for this GAPDH antibody.
Product Application Details	
Applications	Western Blot, Simple Western, Immunocytochemistry/ Immunofluorescence, Immunohistochemistry, Immunohistochemistry-Paraffin
Recommended Dilutions	Western Blot 0.25 - 1 ug/ml, Simple Western 1:25, Immunohistochemistry 5 ug/ml, Immunocytochemistry/ Immunofluorescence 1:10, Immunohistochemistry-Paraffin 5 ug/ml
Application Notes	<p>GAPDH is a widely used loading control for quantitative Western blotting. In IHC-P, the staining of formalin-fixed tissues is enhanced by boiling tissue sections in 10 mM sodium citrate buffer, pH 6.0 for 10-20 min followed by cooling at RT for 20 min.</p> <p>In Simple Western only 10 - 15 uL of the recommended dilution is used per data point.</p> <p>See Simple Western Antibody Database for Simple Western validation: Tested in HeLa lysate 0.1 mg/mL, separated by Size, antibody dilution of 1:25, apparent MW was 44 kDa. Separated by Size-Wes, Sally Sue/Peggy Sue. WB reported in a verified customer review.</p>

Images

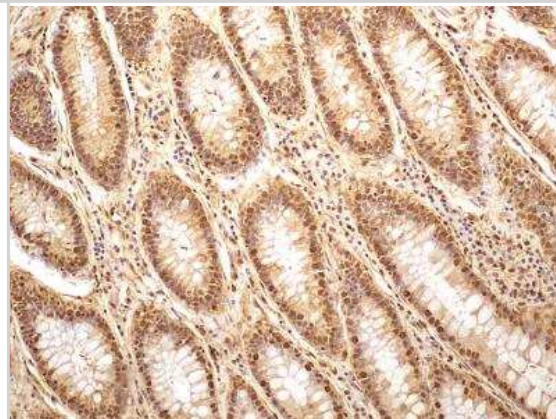
Immunocytochemistry/Immunofluorescence: GAPDH Antibody (13H12) [NBP2-27103] - GAPDH antibody was tested in HeLa cells with Dylight 488 (green). Nuclei and alpha-tubulin were counterstained with DAPI (blue) and Dylight 550 (red). A dilution of 1:10 was used. Image objective 40x.



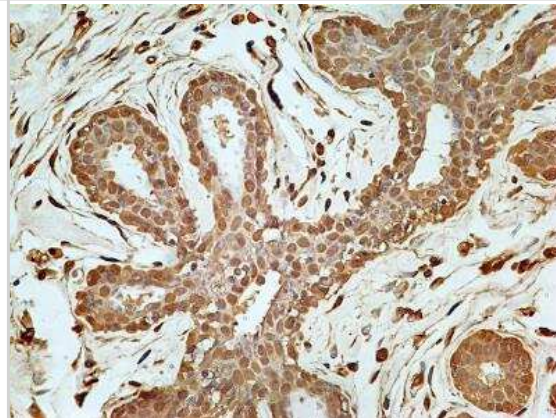
Western Blot: GAPDH Antibody (13H12) [NBP2-27103] - WB detection of GAPDH protein (theoretical molecular weight: 36 kDa) in HeLa cells lysate using GAPDH antibody (clone 13H12) in (A) the absence and (B) the presence of immunizing peptide.



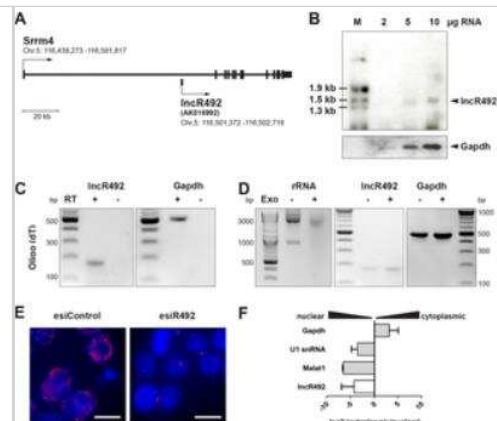
Immunohistochemistry-Paraffin: GAPDH Antibody (13H12) [NBP2-27103] - IHC-P detection GAPDH protein in a formalin-fixed paraffin-embedded section of human rectal carcinoma tissue using GAPDH antibody (clone 13H12) at 5 ug/ml concentration.



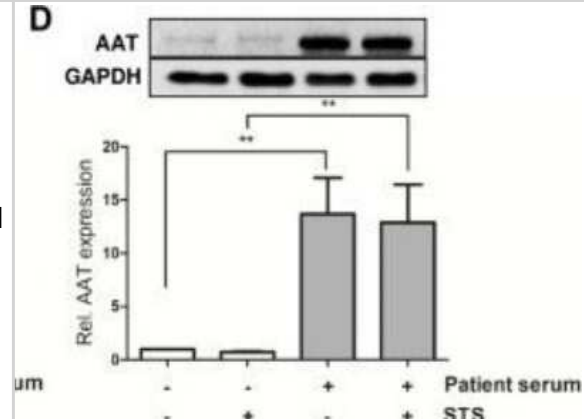
Immunohistochemistry-Paraffin: GAPDH Antibody (13H12) [NBP2-27103] - IHC-P detection GAPDH protein in a formalin-fixed paraffin-embedded section of normal human breast tissue using GAPDH antibody (clone 13H12) at 5 ug/ml concentration.



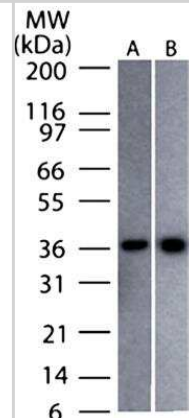
Western Blot: GAPDH Antibody (13H12) [NBP2-27103] - Characterization of IncR492. (B) Northern blot of IncR492 using increasing amounts of loaded total. Black arrow signifies IncR492-specific signal at approximately 1400 bp. A probe targeting Gapdh mRNA served as the loading control. (D) RT-PCR analysis of IncR492 and Gapdh expression. RNA extract was treated with a 5'-phosphate-dependent exonuclease, resulting in a degradation of f.ex. ribosomal RNA. Citation: Winzi M, Casas Vila N, Paszkowski-Rogacz M, Ding L, Noack S, Theis M, et al. (2018) The long noncoding RNA IncR492 inhibits neural differentiation of murine embryonic stem cells. PLoS ONE 13(1): e0191682. <https://doi.org/10.1371/journal.pone.0191682>



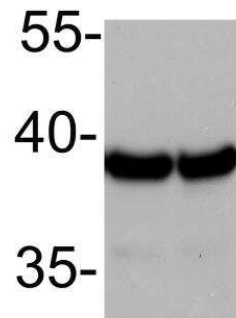
Western Blot: GAPDH Antibody (13H12) [NBP2-27103] - Serum from trauma patients induces intrinsic apoptosis resistance by up-regulating AAT expression in neutrophils. AAT protein expression was analyzed in neutrophils after 18 h of culture by western blot analysis. Relative expression was quantified vs. GAPDH expression. One representative blot of five independent experiments is depicted. ** $p < 0.01$ (one-way ANOVA with Newman keuls post-hoc test). Image collected and cropped by CiteAb from the following publication (<https://dx.plos.org/10.1371/journal.pone.0177450>), licensed under a CC-BY license.



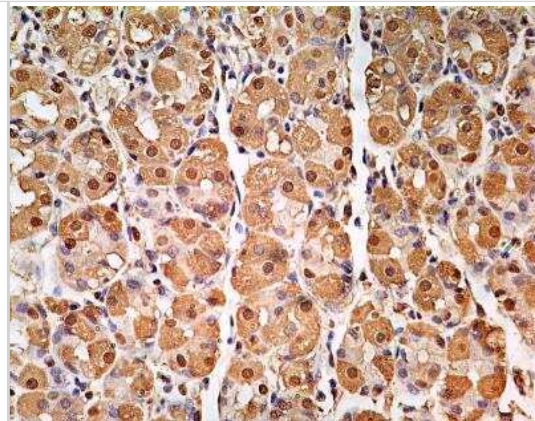
Western Blot: GAPDH Antibody (13H12) [NBP2-27103] - WB detection of GAPDH protein (theoretical molecular weight 36 kDa) in lysates of Mouse cell lines (A) NIH 3T3 (B) RAW 264.7 using GAPDH antibody (clone 13H12) at a concentration of 0.25 $\mu\text{g}/\text{ml}$.



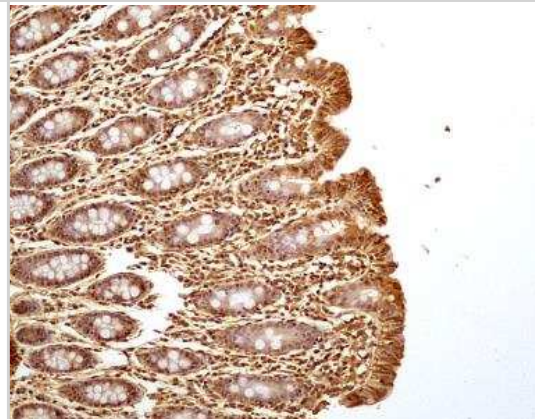
Western Blot: GAPDH Antibody (13H12) [NBP2-27103] - analysis of GAPDH in HeLa and HEK 293 cells (25 $\mu\text{g}/\text{lane}$) using anti-GAPDH antibody. Image from verified customer review.



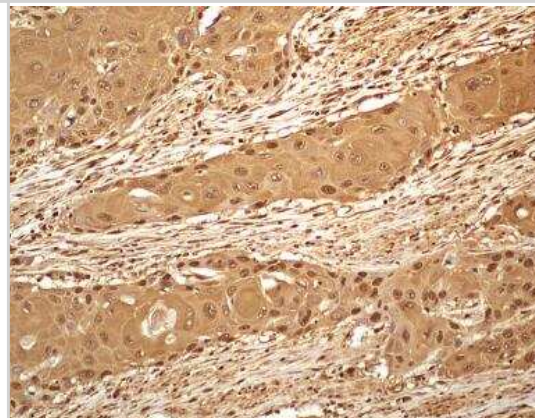
Immunohistochemistry-Paraffin: GAPDH Antibody (13H12) [NBP2-27103] - IHC-P detection GAPDH protein in a formalin-fixed paraffin-embedded section of normal human stomach tissue using GAPDH antibody (clone 13H12) at 5 ug/ml concentration.



Immunohistochemistry-Paraffin: GAPDH Antibody (13H12) [NBP2-27103] - IHC-P detection GAPDH protein in a formalin-fixed paraffin-embedded section of human colon tissue using GAPDH antibody (clone 13H12) at 5 ug/ml concentration.



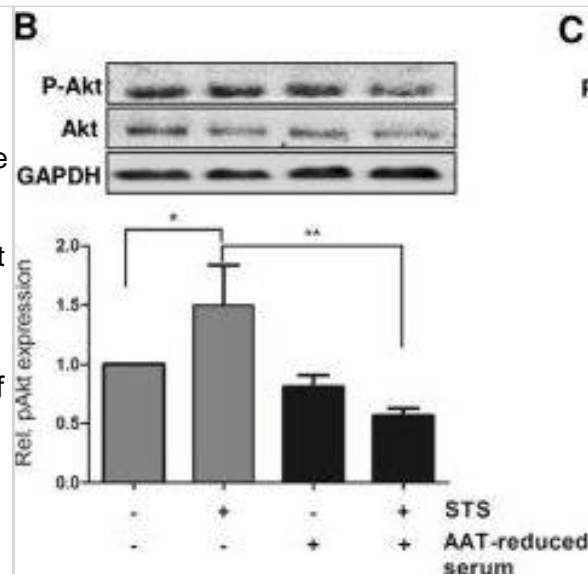
Immunohistochemistry-Paraffin: GAPDH Antibody (13H12) [NBP2-27103] - IHC-P detection GAPDH protein in a formalin-fixed paraffin-embedded tissue section of human esophageal squamous cell carcinoma (SCC) using GAPDH antibody (clone 13H12) at 5 ug/ml concentration.



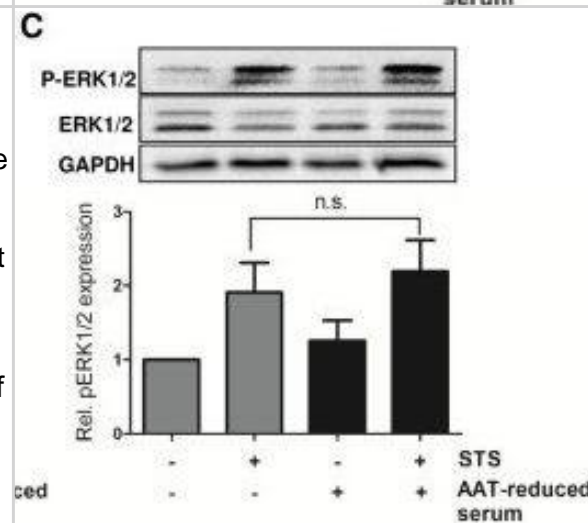
Simple Western: GAPDH Antibody (13H12) [NBP2-27103] - GAPDH/G3PDH Antibody (13H12) [NBP2-27103] - Simple Western lane view shows a specific band for GAPDH in 0.1 mg/ml of HeLa lysate. This experiment was performed under reducing conditions using the 12-230 kDa separation system.



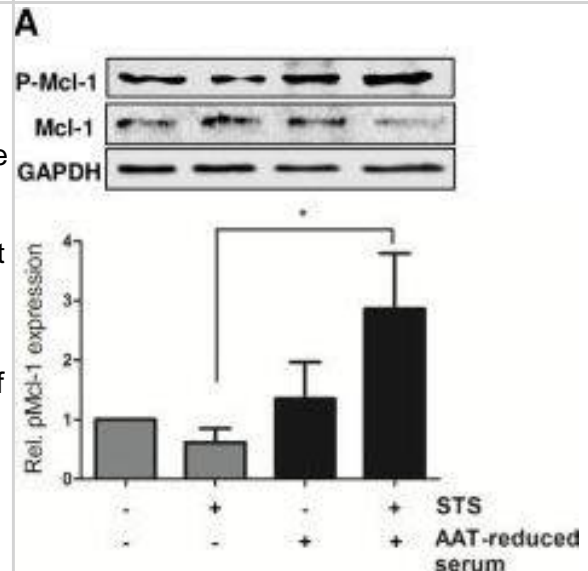
Western Blot: GAPDH Antibody (13H12) - BSA Free [NBP2-27103] - Effect of AAT on Mcl-1 phosphorylation, the activity of MAP kinases & caspases. Neutrophils from healthy volunteers ($2.5 \times 10^6/ml$) were cultured in medium supplemented with patient serum (3 mg protein/ml; 1%) & those containing low levels of AAT (AAT-reduce serum; 1%) in the presence of STS ($0.2 \mu M$). After 3 h, the expression of pMcl-1 (A, $n = 7$), pAkt (B, $n = 8$) & pERK1/2 (C, $n = 10$) were analyzed by western blot. Expression levels of the phosphorylated proteins were normalized to that of the unphosphorylated forms. GAPDH was used as loading control. One representative blot is displayed. * $p < 0.05$; ** $p < 0.01$; n.s. = not significant. D. After 4 h of incubation the activities of caspase-9 & caspase-3/-7 were quantified. Results are presented as means \pm SEM of eight independent experiments. No significant differences were found (one-way ANOVA with Newman keuls post-hoc test). Image collected & cropped by CiteAb from the following publication (<https://pubmed.ncbi.nlm.nih.gov/28493974>), licensed under a CC-BY license. Not internally tested by Novus Biologicals.



Western Blot: GAPDH Antibody (13H12) - BSA Free [NBP2-27103] - Effect of AAT on Mcl-1 phosphorylation, the activity of MAP kinases & caspases. Neutrophils from healthy volunteers ($2.5 \times 10^6/ml$) were cultured in medium supplemented with patient serum (3 mg protein/ml; 1%) & those containing low levels of AAT (AAT-reduce serum; 1%) in the presence of STS ($0.2 \mu M$). After 3 h, the expression of pMcl-1 (A, $n = 7$), pAkt (B, $n = 8$) & pERK1/2 (C, $n = 10$) were analyzed by western blot. Expression levels of the phosphorylated proteins were normalized to that of the unphosphorylated forms. GAPDH was used as loading control. One representative blot is displayed. * $p < 0.05$; ** $p < 0.01$; n.s. = not significant. D. After 4 h of incubation the activities of caspase-9 & caspase-3/-7 were quantified. Results are presented as means \pm SEM of eight independent experiments. No significant differences were found (one-way ANOVA with Newman keuls post-hoc test). Image collected & cropped by CiteAb from the following publication (<https://pubmed.ncbi.nlm.nih.gov/28493974>), licensed under a CC-BY license. Not internally tested by Novus Biologicals.



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Publications

Guo D, Liu S, Zhang J et Al. Prickle1-driven basement membrane deposition of the iPSC-derived embryoid bodies is separable from the establishment of apicobasal polarity Cell Prolif 2024-01-07 [PMID: 38185785]

Kate WD, Fanta M, Weinfeld M. et Al. Loss of the DNA repair protein, polynucleotide kinase/phosphatase, activates the type 1 interferon response independent of ionizing radiation Nucleic Acids Res 2024-09-09 [PMID: 39087523]

Vanda Balint, Mina Peric, Sanja Dacic, Danijela Stanisavljevic Ninkovic, Jelena Marjanovic, Jelena Popovic, Milena Stevanovic, Andrijana Lazic The Role of SOX2 and SOX9 Transcription Factors in the Reactivation-Related Functional Properties of NT2/D1-Derived Astrocytes. Biomedicines 2024-04-03 [PMID: 38672150]

Wang C, Terrigno M, Li J et al. Increased G3BP2-Tau interaction in tauopathies is a natural defense against Tau aggregation Neuron 2023-06-23 [PMID: 37385246]

Lin X, Fu B, Xiong Y et al. Unconventional secretion of unglycosylated ORF8 is critical for the cytokine storm during SARS-CoV-2 infection PLoS pathogens 2023-01-01 [PMID: 36689483] (WB)

Venkatramanan, S, Ibar, C Et al. TRIP6 is required for tension at adherens junctions. J Cell Sci 2021-03-11 [PMID: 33558314] (IF/IHC, Mouse)

Grotheer V, Skrynecki N, Oezel L et al. Osteogenic differentiation of human mesenchymal stromal cells and fibroblasts differs depending on tissue origin and replicative senescence Scientific reports 2021-06-07 [PMID: 34099837] (WB, Human)

Krassovka JM, Suschek CV, Prost M et al. The impact of non-toxic blue light (453 nm) on cellular antioxidative capacity, TGF-beta 1 signaling, and myofibrogenesis of human skin fibroblasts J. Photochem. Photobiol. B, Biol. 2020 -07-06 [PMID: 32659647] (WB, Human)

Srinivas C, Ramaiah MJ, Lavanya A et al Novel EPE Analogue Modulates Expression of Angiogenesis Associated microRNAs and Regulates Cell Proliferation by Targeting STAT3 in Breast Cancer PLoS ONE 2015-11-10 [PMID: 26551008] (WB, Human)

Modi, A;Singh, M;Gutti, G;Shanker, OR;Singh, VK;Singh, S;Singh, SK;Pradhan, S;Narayan, G; Benzothiazole derivative bearing amide moiety induces p53-mediated apoptosis in HPV16 positive cervical cancer cells Invest New Drugs 2019-08-20 [PMID: 31432292] (WB, Human)

Srivastava, S;Battu, MB;Khan, MZ;Nandicoori, VK;Mukhopadhyay, S; Mycobacterium tuberculosis PPE2 Protein Interacts with p67phox and Inhibits Reactive Oxygen Species Production J. Immunol. 2019-09-01 [PMID: 31375544] (WB, Mouse)

Eiraku, N;Chiba, N;Nakamura, T;Amir, MS;Seong, CH;Ohnishi, T;Kusuyama, J;Noguchi, K;Matsuguchi, T; BMP9 directly induces rapid GSK3- β phosphorylation in a Wnt-independent manner through class I PI3K-Akt axis in osteoblasts FASEB J. 2019-07-31 [PMID: 31365832]

More publications at <http://www.novusbio.com/NBP2-27103>



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

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NB720-B	Rabbit anti-Mouse IgG (H+L) Secondary Antibody [Biotin]
NBP1-43319-0.5mg	Mouse IgG1 Kappa Isotype Control (P3.6.2.8.1)

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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.

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