

For Research Use Only

# NIPSNAP1 Polyclonal antibody

Catalog Number:33020-1-AP



## Basic Information

|   |   |  |
|---|---|--|
| <b>Catalog Number:</b><br>33020-1-AP        | <b>GenBank Accession Number:</b><br>NM_003634.3     | <b>Purification Method:</b><br>Antigen affinity Purification |
| <b>Concentration:</b><br>350 µg/ml          | <b>GeneID (NCBI):</b><br>8508                       | <b>Recommended Dilutions:</b><br>WB 1:5000-1:50000           |
| <b>Source:</b><br>Rabbit                    | <b>UNIPROT ID:</b><br>Q9BPW8                        |  |
| <b>Isotype:</b><br>IgG                      | <b>Full Name:</b><br>nipsnap homolog 1 (C. elegans) |  |
| <b>Immunogen Catalog Number:</b><br>AG36474 | <b>Calculated MW:</b><br>33kDa, 284aa               |  |
|   | <b>Observed MW:</b><br>30 kDa                       |  |

## Applications

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| <b>Tested Applications:</b><br>WB, ELISA         | <b>Positive Controls:</b><br>WB : rat liver tissue, HepG2 cells, HuH-7 cells, mouse brain tissue, mouse liver tissue |
| <b>Species Specificity:</b><br>human, mouse, rat |  |

## Background Information

NIPSNAP1 has been shown to play a role in regulating synaptic activity in learning and memory but nonetheless, along with the brain, NIPSNAP1 is highly abundant in other organs such as the liver, kidney, and adrenals but its expression is less evident in other tissues such as skeletal muscle. Functional investigations revealed the knockdown of NIPSNAP1 in cancer cells induced cell cycle arrest and inhibited proliferation with the underlying phenotype resulting from senescence. Further analyses revealed that NIPSNAP1 engages in two separate binding interactions that prevent cellular senescence. First, NIPSNAP1 engages with the E3 ubiquitin ligase FBXL14 which otherwise serves to target c-Myc for proteasomal degradation. And moreover, this was shown to be part of a regulatory feedback loop since NIPSNAP1 is subject to transcriptional repression by c-Myc. The second interaction involves NIPSNAP1 binding to superoxide dismutase 2 (SOD2) which enhances its association with SIRT3, thereby deacetylating SOD2 and increasing its activity, in turn, alleviating elevated ROS levels. Consequently, silencing NIPSNAP1 expression contributes to P27-mediated senescence via both decreasing the levels of c-Myc together with increasing ROS levels. Together these findings reveal an important role for NIPSNAP1 as a negative regulator of cancer cell senescence, which functions to sustain the viability of cancer cells under growth factor deprivation stress (PMID: 37340421).

## Storage

**Storage:**  
Store at -20°C. Stable for one year after shipment.  
**Storage Buffer:**  
PBS with 0.02% sodium azide and 50% glycerol, pH7.3  
Aliquoting is unnecessary for -20°C storage

For technical support and original validation data for this product please contact:

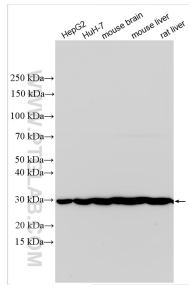
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## Selected Validation Data



Various lysates were subjected to SDS PAGE followed by western blot with 33020-1-AP (NIPSNAP1 antibody) at dilution of 1:15000 incubated at room temperature for 1.5 hours.