Ndn (36-V): sc-101224



The Power to Question

BACKGROUND

Prader-Willi syndrome (PWS) is a neurogenetic disorder resulting from the loss of paternal expression of genes localized in the 15q11-q12 region. Clinical manifestations of this disease include feeding problems in infancy, temper outbursts, perseveration, obsessive-compulsive symptoms and sleep disturbances. Necdin (Ndn) protein is generated from an intronless gene that is located in the Prader-Willi syndrome deletion region. Studies in mouse suggest that the protein encoded by this gene may suppress growth in post-mitotic neurons. Ndn expression in brain is restricted to post-mitotic neurons and parental alleles display a differential methylation profile in the coding region. Reduced expression of Ndn is responsible for at least a subset of the clinical manifestations of PWS, including skin picking and improved spatial memory.

REFERENCES

- 1. Watrin, F., et al. 1997. The mouse Necdin gene is expressed from the paternal allele only and lies in the 7C region of the mouse chromosome 7, a region of conserved synteny to the human Prader-Willi syndrome region. Eur. J. Hum. Genet. 5: 324-332.
- Jay, P., et al. 1997. The human Necdin gene, NDN, is maternally imprinted and located in the Prader-Willi syndrome chromosomal region. Nat. Genet. 17: 357-361.
- 3. Muscatelli, F., et al. 2000. Disruption of the mouse Necdin gene results in hypothalamic and behavioral alterations reminiscent of the human Prader-Willi syndrome. Hum. Mol. Genet. 9: 3101-3110.
- Oeffner, F., et al. 2001. Systematic screening for mutations in the human Necdin gene (NDN): identification of two naturally occurring polymorphisms and association analysis in body weight regulation. Int. J. Obes. Relat. Metab. Disord. 25: 767-779.
- 5. Online Mendelian Inheritance in Man, OMIM™. 2001. Johns Hopkins University, Baltimore, MD. MIM Number: 602117. World Wide Web URL: http://www.ncbi.nlm.nih.gov/omim/
- 6. LocusLink Report (LocusID: 4692). http://www.ncbi.nlm.nih.gov/LocusLink/

CHROMOSOMAL LOCATION

Genetic locus: NDN (human) mapping to 15q11.2.

SOURCE

Ndn (36-V) is a mouse monoclonal antibody raised against recombinant Ndn of human origin.

PRODUCT

Each vial contains 100 μg lgG_{2a} kappa light chain in 1.0 ml of PBS with < 0.1% sodium azide and 0.1% gelatin.

STORAGE

Store at 4° C, **DO NOT FREEZE**. Stable for one year from the date of shipment. Non-hazardous. No MSDS required.

APPLICATIONS

Ndn (36-V) is recommended for detection of Ndn of human origin by Western Blotting (starting dilution 1:200, dilution range 1:100-1:1000), immunoprecipitation [1-2 μ g per 100-500 μ g of total protein (1 ml of cell lysate)], immunofluorescence (starting dilution 1:50, dilution range 1:50-1:500), immunohistochemistry (including paraffin-embedded sections) (starting dilution 1:50, dilution range 1:50-1:500) and solid phase ELISA (starting dilution 1:30, dilution range 1:30-1:3000).

Suitable for use as control antibody for Ndn siRNA (h): sc-37318, Ndn shRNA Plasmid (h): sc-37318-SH and Ndn shRNA (h) Lentiviral Particles: sc-37318-V.

Molecular Weight (predicted) of Ndn: 36 kDa.

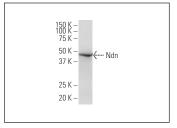
Molecular Weight (observed) of Ndn: 50 kDa.

Positive Controls: HL-60 whole cell lysate: sc-2209.

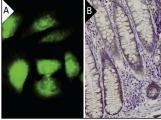
RECOMMENDED SUPPORT REAGENTS

To ensure optimal results, the following support reagents are recommended: 1) Western Blotting: use m-lgG κ BP-HRP: sc-516102 or m-lgG κ BP-HRP (Cruz Marker): sc-516102-CM (dilution range: 1:1000-1:10000), Cruz MarkerTM Molecular Weight Standards: sc-2035, UltraCruz[®] Blocking Reagent: sc-516214 and Western Blotting Luminol Reagent: sc-2048. 2) Immunoprecipitation: use Protein A/G PLUS-Agarose: sc-2003 (0.5 ml agarose/2.0 ml). 3) Immunofluorescence: use m-lgG κ BP-FITC: sc-516140 or m-lgG κ BP-PE: sc-516141 (dilution range: 1:50-1:200) with UltraCruz[®] Mounting Medium: sc-24941 or UltraCruz[®] Hard-set Mounting Medium: sc-359850. 4) Immunohistochemistry: use m-lgG κ BP-HRP: sc-516102 with DAB, 50X: sc-24982 and Immunohistomount: sc-45086, or Organo/Limonene Mount: sc-45087.

DATA



Ndn (36-V): sc-101224. Western blot analysis of Ndn expression in HL-60 whole cell lysate.



Ndn (36-V): sc-101224. Immunofluorescence staining of paraformaldehyde-fixed HeLa cells showing nuclear and cytoplasmic localization (A). Immunoperoxidase staining of formalin-fixed, paraffin-embedded human colon tissue showing membrane and cytoplasmic localization (B).

SELECT PRODUCT CITATIONS

 Sun, Y., et al. 2022. Noninvasive urinary protein signatures associated with colorectal cancer diagnosis and metastasis. Nat. Commun. 13: 2757.

RESEARCH USE

For research use only, not for use in diagnostic procedures