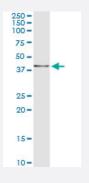


# NDN monoclonal antibody (M15), clone 2G8

Catalog # H00004692-M15 Size 100 ug

### Applications



### Immunoprecipitation

Immunoprecipitation of NDN transfected lysate using anti-NDN monoclonal antibody and Protein A Magnetic Bead, and immunoblotted with NDN MaxPab rabbit polyclonal antibody.

Specification	
Product Description	Mouse monoclonal antibody raised against a full length recombinant NDN.
Immunogen	NDN (NP_002478, 102 a.a. ~ 183 a.a) full length recombinant protein with GST tag. MW of the GST t ag alone is 26 KDa.
Sequence	AHELMWYVLVKDQKKMIWFPDMVKDVIGSYKKWCRSILRRTSLILARVFGLHLRLTSLHTMEFALV KALEPEELDRVALSN
Host	Mouse
Reactivity	Human
lsotype	lgG2b Kappa
Quality Control Testing	Antibody Reactive Against Recombinant Protein.
Storage Buffer	In 1x PBS, pH 7.4
Storage Instruction	Store at -20°C or lower. Aliquot to avoid repeated freezing and thawing.



## Applications

Immunoprecipitation

Immunoprecipitation of NDN transfected lysate using anti-NDN monoclonal antibody and Protein A Magnetic Bead, and immunoblotted with NDN MaxPab rabbit polyclonal antibody.

Protocol Download

ELISA

Gene Info — NDN	
Entrez GenelD	<u>4692</u>
GeneBank Accession#	<u>NM_002487</u>
Protein Accession#	<u>NP_002478</u>
Gene Name	NDN
Gene Alias	HsT16328, PWCR
Gene Description	necdin homolog (mouse)
Omim ID	<u>176270 602117</u>
Gene Ontology	Hyperlink
Gene Summary	This intronless gene is located in the Prader-Willi syndrome deletion region. It is an imprinted gen e and is expressed exclusively from the paternal allele. Studies in mouse suggest that the protein encoded by this gene may suppress growth in postmitotic neurons. [provided by RefSeq
Other Designations	OTTHUMP00000159437 necdin

#### Disease

- <u>Attention Deficit Disorder with Hyperactivity</u>
- <u>Autistic Disorder</u>
- Body Weight
- <u>NARP</u>

😵 Abnova

- <u>Obesity</u>
- Prader-Willi syndrome
- <u>Sleep Apnea</u>