



POLYCLONAL ANTIBODY

For research use only. Not for clinical diagnosis.

Catalog No. 73-600

Anti-Necdin antibody, (rabbit polyclonal, NC243)

BACKGROUND

Necdin (neurally differentiated embryonal carcinoma-derived protein) is a 325-amino acid residue protein encoded by a cDNA clone isolated from neurally differentiated mouse embryonal carcinoma cells (ref.1). **Necdin** is a potent growth suppressor that is expressed predominantly in postmitotic cells such as neurons and muscle cells. **Necdin** has been implicated in the pathogenesis of Prader-Willi syndrome, a human neurodevelopmental disorder associated with genomic imprinting. Furthermore, **necdin** binds to major transcription factors E2F1 and p53, and also to NEFA and nucleobindin, both of which are calcium-binding proteins involved in intracellular calcium homeostasis. From these findings **necdin** is suggested to target various factors involved in the regulation of cell proliferation and survival, and plays a key role in development and differentiation of subsets of neurons in the brain. An antibody (named NC243) against mouse **necdin** was raised in rabbit (ref.2).

Product type	Primary antibodies
Host	Rabbit
Source	Serum
Form	Liquid Undiluted serum added with 0.05% sodium azide.
Volume	100 µl
Concentration	
Specificity	
Antigen	Recombinant GST-fused mouse necdin (aa 83-325).
Isotype	

Application notes WB, IHC, IC

Recommended use

Recommended dilutions

Western blotting: 1,000~3,000 fold dilution)
Immunohistochemistry: 500~1,000 fold dilution)
Immunocytochemistry: 500~1,000 fold dilution

Optimal dilutions/concentrations should be determined by the end user.

Data Link: Swiss-Prot [P25233](#) (mouse), [Q99608](#) (human)

Cross reactivity Mouse, Rat and Human Necdin

Storage Shipped and stored at 4°C for a few months. For longer period, aliquot and store at -20°C

- References**
- 1) Maruyama K *et al* (1991) "A novel brain-specific mRNA encoding nuclear protein (necdin) expressed in neurally differentiated embryonal carcinoma cells." *Biochem Biophys Res Commun* **178**: 291-296 PMID: [2069569](#)
 - 2) Niinobe M *et al.* (2000) "Cellular and subcellular localization of necdin in fetal and adult mouse brain." *Dev Neurosci* **22**: 310-319 PMID: [10965153](#)
 - 3) Kuwajima T *et al* (2004) "Necdin interacts with the Msx2 homeodomain protein via MAGE-D1 to promote myogenic differentiation of C2C12 cells." *J Biol Chem* **279**: 40484-40493 PMID: [15272023](#)
 - 4) Kurita M *et al* (2006) "Necdin downregulates Cdc2 expression to attenuate neuronal apoptosis." *J Neurosci* **26**: 12003-12013 PMID: [17108174](#)

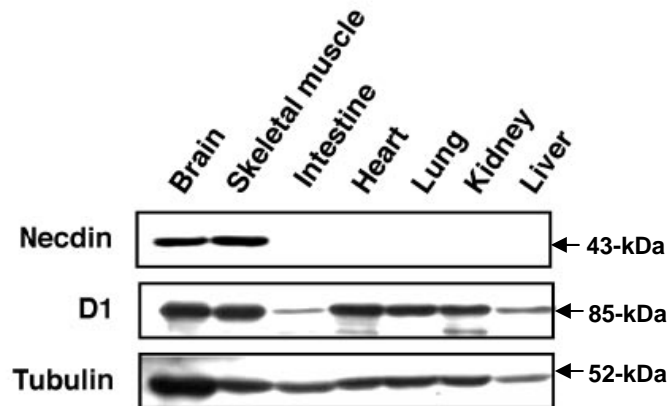


Fig.1 Western blotting using this antibody.

Western blot analysis of necdin and MAGE-D1 in mouse embryonic tissues (ref.3). Tissue lysates (20 ug) from E18.5 embryos were separated by SDS-PAGE and immunoblotted with antibodies against necdin, MAGE-D1 (D1) and tubulin. Endogenous ~43-kDa necdin protein was detected almost exclusively in the brain and skeletal muscle whereas endogenous ~85-kDa MAGE-D1 protein was expressed in a ubiquitous manner.

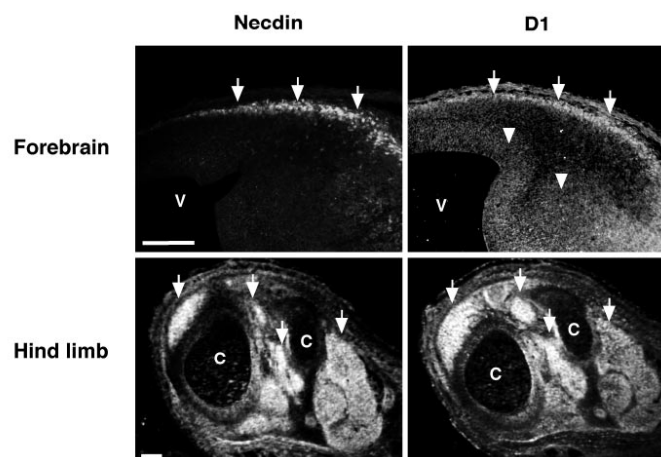


Fig.2 Immunohistochemistry using this antibody (ref.3).

Fluorescence immunohistochemistry for necdin and MAGE-D1 in mouse embryos. Adjacent sections of the forebrain at E12.5 embryo (upper panels) and the hind limb at E14.5 (lower panels) were stained with antibodies against necdin and MAGE-D1. The arrows point to the preplate (upper panels) and the skeletal muscles (lower panels). The arrowheads indicate the ventricular proliferative zone; V, ventricle; C, bone cavity. Necdin and MAGE-D1 were concentrated in the preplate of the forebrain at E12.5 and skeletal muscle tissues in the hind limb at E14.5. In developing neural tube, MAGE-D1 immunoreactivity was distributed in the ventricular zone as well as the marginal zone.



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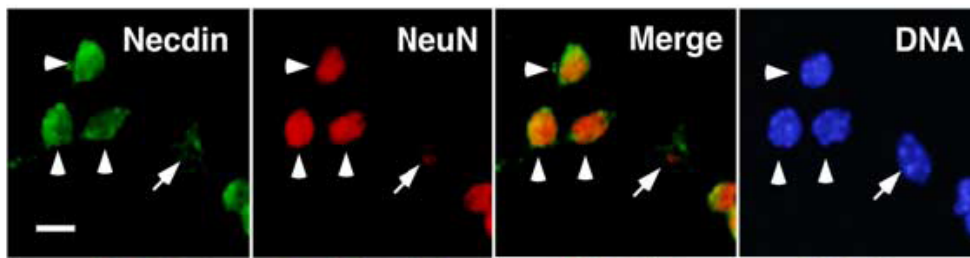


Fig.3 Immunocytochemistry using this antibody (ref.4).

Primary cerebellar granule neurons (CGNs) were fixed and double stained for nectdin and postmitotic neuron marker, NeuN. Chromosomal DNA was stained with Hoechst 33342. Arrowheads point to CGNs, and arrows point to CGN progenitors or non-neuronal cells.

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