NSE Rabbit mAb

Catalog No: #49015

Package Size: #49015-1 50ul #49015-2 100ul



Orders: order@signalwayantibody.com Support: tech@signalwayantibody.com

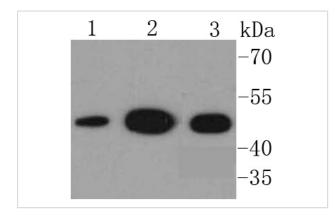
Description

Product Name	NSE Rabbit mAb
Host Species	Recombinant Rabbit
Clonality	Monoclonal
Clone No.	SC06-28
Purification	ProA affinity purified
Applications	WB, ICC, IHC
Species Reactivity	Hu, Ms, Rt, zebrafish
Immunogen Description	recombinant protein
Conjugates	Unconjugated
Other Names	2 phospho D glycerate hydrolyase antibody 2-phospho-D-glycerate hydro-lyase antibody Eno 2 antibody
	ENO2 antibody ENOG antibody ENOG_HUMAN antibody Enolase 2 (gamma, neuronal) antibody Enolase
	2 antibody Enolase 2 gamma neuronal antibody Enolase2 antibody Epididymis secretory protein Li 279
	antibody Gamma enolase antibody Gamma-enolase antibody HEL S 279 antibody Neural enolase antibody
	Neuron specific enolase antibody Neuron specific gamma enolase antibody Neuron-specific enolase
	antibody Neurone specific enolase antibody NSE antibody
Accession No.	Swiss-Prot#:P09104
Calculated MW	47 kDa
Formulation	1*TBS (pH7.4), 1%BSA, 40%Glycerol. Preservative: 0.05% Sodium Azide.
Storage	Store at -20°C

Application Details

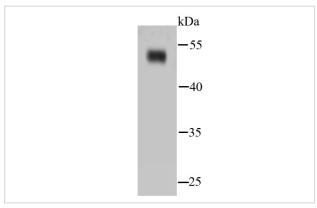
WB: 1:1,000-1:2,000 IHC: 1:50-1:200 ICC: 1:50-1:200

Images

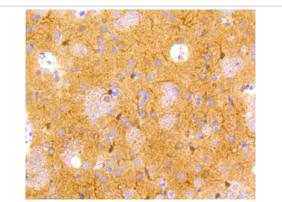


Western blot analysis of NSE on different lysates using anti-NSE antibody at 1/1,000 dilution. Positive control:

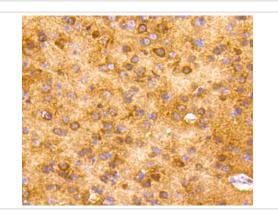
Lane 1: HepG2 Lane 2: Hela Lane 3: 293



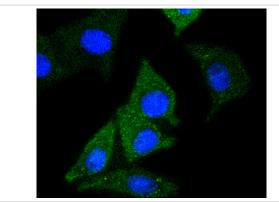
Western blot analysis of NSE on hybrid fish (crucian-carp) brain tissue lysates using anti-NSE antibody at 1/500 dilution.



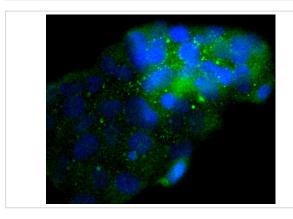
Immunohistochemical analysis of paraffin-embedded rat brain tissue using anti-NSE antibody. Counter stained with hematoxylin.



Immunohistochemical analysis of paraffin-embedded human tonsil tissue using anti-NSE antibody. Counter stained with hematoxylin.



ICC staining NSE in SH-SY-5Y cells (green). The nuclear counter stain is DAPI (blue). Cells were fixed in paraformaldehyde, permeabilised with 0.25% Triton X100/PBS.



ICC staining NSE in 293 cells (green). The nuclear counter stain is DAPI (blue). Cells were fixed in paraformaldehyde, permeabilised with 0.25% Triton X100/PBS.

Background

Enolases have been characterized as highly conserved cytoplasmic glycolytic enzymes that may be involved in differentiation. Three isoenzymes have been identified, α Enolase, β Enolase and γ Enolase. α Enolase expression has been detected on most tissues, whereas β Enolase is expressed predominantly in muscle tissue and γ Enolase is detected only in nervous tissue. These isoforms exist as both homodimers and heterodimers, and they play a role in converting phosphoglyceric acid to phosphenolpyruvic acid in the glycolytic pathway.

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Note: This product is for in vitro research use only and is not intended for use in humans or animals.