

Product datasheet for UM800177CF

OriGene Technologies, Inc.

9620 Medical Center Drive, Ste 200 Rockville, MD 20850, US Phone: +1-888-267-4436 https://www.origene.com techsupport@origene.com EU: info-de@origene.com CN: techsupport@origene.cn

NSE (ENO2) Mouse Monoclonal Antibody [Clone ID: UMAB287]

Product data:

Product Type: Primary Antibodies

Clone Name: UMAB287
Applications: IHC, WB

Recommended Dilution: WB 1:1000, IHC 1:15000

Reactivity: Mouse, Rat **Host:** Mouse

Isotype: IgG2b

Clonality: Monoclonal

Immunogen: Human recombinant protein fragment corresponding to amino acids 188-293 of human NSE

(NP 001966) produced in E.coli.

Formulation: Lyophilized powder (original buffer 1X PBS, pH 7.3, 8% trehalose)

Reconstitution Method: For reconstitution, we recommend adding 100uL distilled water to a final antibody

concentration of about 1 mg/mL. To use this carrier-free antibody for conjugation experiment, we strongly recommend performing another round of desalting process. (OriGene recommends Zeba Spin Desalting Columns, 7KMWCO from Thermo Scientific)

Purification: Purified from mouse ascites fluids or tissue culture supernatant by affinity chromatography

(protein A/G)

Conjugation: Unconjugated

Predicted Protein Size: 47.3 kDa

Gene Name: Homo sapiens enolase 2 (ENO2), mRNA.

Database Link: NP 001966

Entrez Gene 13807 MouseEntrez Gene 24334 Rat

P09104

Background: This gene encodes one of the three enclase isoenzymes found in mammals. This isoenzyme,

a homodimer, is found in mature neurons and cells of neuronal origin. A switch from alpha enolase to gamma enolase occurs in neural tissue during development in rats and primates.

[provided by RefSeq, Jul 2008]

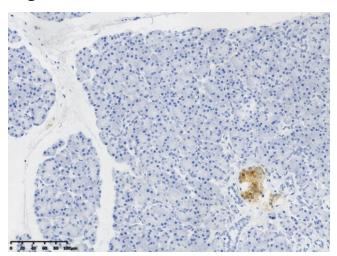
Synonyms: HEL-S-279; NSE



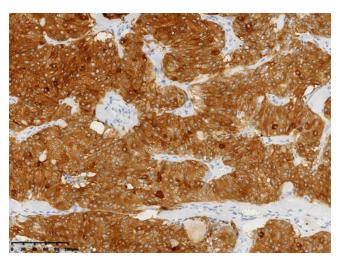


Protein Pathways: Glycolysis / Gluconeogenesis, Metabolic pathways, RNA degradation

Product images:



Immunohistochemical staining of paraffinembedded Human pancreas tissue within the normal limits using anti-NSE (ENO2) mouse monoclonal antibody. (Heat-induced epitope retrieval by 1mM EDTA in 10mM Tris buffer (pH8.0) at 120°C for 3 min, [UM800177]) (1:15000)

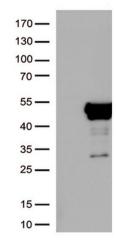


Immunohistochemical staining of paraffinembedded Carcinoma of Human thyroid tissue using anti-NSE (ENO2) mouse monoclonal antibody.(Heat-induced epitope retrieval by 1mM EDTA in 10mM Tris buffer (pH8.0) at 120°C for 3 min, [UM800177]) (1:15000)

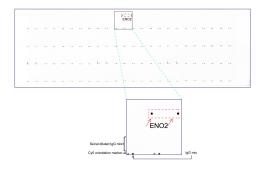




Immunohistochemical staining of paraffinembedded Human adult brain tissue within the normal limits using anti-NSE (ENO2) mouse monoclonal antibody.(Heat-induced epitope retrieval by 1mM EDTA in 10mM Tris buffer (pH8.0) at 120 oC for 3 min, [UM800177]) (1:15000)



HEK293T cells were transfected with the pCMV6-ENTRY control (Cat# [PS100001], Left lane) or pCMV6-ENTRY NSE (ENO2) (Cat# [RC201085], Right lane) cDNA for 48 hrs and lysed. Equivalent amounts of cell lysates (5 ug per lane) were separated by SDS-PAGE and immunoblotted with anti-NSE(ENO2) antibody (Cat# [UM800177]) (1:1000).



OriGene overexpression protein microarray chip was immunostained with UltraMAB anti-NSE (ENO2) mouse monoclonal antibody ([UM800177]). The positive reactive proteins are highlighted with two red arrows in the enlarged subarray. All the positive controls spotted in this subarray are also labeled for clarification. (1:100)