## SANTA CRUZ BIOTECHNOLOGY, INC.

# Enolase (A-5): sc-271384



#### BACKGROUND

Enolases have been characterized as highly conserved cytoplasmic glycolytic enzymes that may be involved in differentiation. Three isoenzymes have been identified:  $\alpha$  Enolase,  $\beta$  Enolase and  $\gamma$  Enolase.  $\alpha$  Enolase expression has been detected on most tissues, whereas  $\beta$  Enolase is expressed predominantly in muscle tissue and  $\gamma$  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. The 433 amino acid protein shows 67% homology to yeast Enolase and 94% homology to rat nonneural Enolase. Studies also indicate that  $\alpha$  Enolase is encoded by the same gene that encodes  $\tau$ -crystallin, a lens structural protein.

## REFERENCES

- Whitehead, M.C., et al. 1982. Synapse formation is related to the onset of neuron-specific Enolase immunoreactivity in the avian auditory and vestibular systems. Dev. Neurosci. 5: 298-307.
- 2. Giallongo, A., et al. 1986. Molecular cloning and nucleotide sequence of a full-length cDNA for human  $\alpha$  Enolase. Proc. Natl. Acad. Sci. USA 83: 6741-6745.

#### SOURCE

Enolase (A-5) is a mouse monoclonal antibody raised against amino acids 1-300 mapping at the N-terminus of  $\alpha$  Enolase of human origin.

#### PRODUCT

Each vial contains 200  $\mu g$  IgG\_1 kappa light chain in 1.0 ml of PBS with < 0.1% sodium azide and 0.1% gelatin.

Enolase (A-5) is available conjugated to agarose (sc-271384 AC), 500 µg/ 0.25 ml agarose in 1 ml, for IP; to HRP (sc-271384 HRP), 200 µg/ml, for WB, IHC(P) and ELISA; to either phycoerythrin (sc-271384 PE), fluorescein (sc-271384 FITC), Alexa Fluor<sup>®</sup> 488 (sc-271384 AF488), Alexa Fluor<sup>®</sup> 546 (sc-271384 AF546), Alexa Fluor<sup>®</sup> 594 (sc-271384 AF594) or Alexa Fluor<sup>®</sup> 647 (sc-271384 AF546), 200 µg/ml, for WB (RGB), IF, IHC(P) and FCM; and to either Alexa Fluor<sup>®</sup> 680 (sc-271384 AF680) or Alexa Fluor<sup>®</sup> 790 (sc-271384 AF790), 200 µg/ml, for Near-Infrared (NIR) WB, IF and FCM.

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## **APPLICATIONS**

Enolase (A-5) is recommended for detection of  $\alpha$  Enolase,  $\beta$  Enolase and  $\gamma$  Enolase of mouse, rat and human origin by Western Blotting (starting dilution 1:100, dilution range 1:100-1:1000), immunoprecipitation [1-2  $\mu$ g per 100-500  $\mu$ 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).

Molecular Weight of Enolase: 48 kDa.

Positive Controls: IMR-32 cell lysate: sc-2409, HeLa whole cell lysate: sc-2200 or KNRK whole cell lysate: sc-2214.

#### STORAGE

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

## DATA



Enolase (A-5): sc-271384. Western blot analysis of Enolase expression in HeLa (Å), IMR-32 (B), NIH/3T3 (C), KNRK (D) and SH-SYSY (E) whole cell lysates and rat brain tissue extract (F).



Enolase (A-5) Alexa Fluor<sup>®</sup> 488: sc-271384 AF488. Direct immunofluorescence staining of formalin-fixed SW480 cells showing cytoplasmic localization. Blocked with UltraCruz<sup>®</sup> Blocking Reagent: sc-516214 (**A**). Enolase (A-5) HRP: sc-271384 HRP. Direct immunoperoxidase staining of formalin fixed, paraffin-embedded human esophagus tissue showing cytoplasmic and nuclear staining of squamous epithelial cells. Blocked **B** with 0.25X UltraCruz<sup>®</sup> Blocking Reagent: sc-516214 (**B**).

#### **SELECT PRODUCT CITATIONS**

- Herrera-Martínez, M., et al. 2013. Actin, RhoA, and Rab11 participation during encystment in *Entamoeba invadens*. Biomed Res. Int. 2013: 919345.
- Hertz, R., et al. 2014. The *Entamoeba histolytica* Dnmt2 homolog (Ehmeth) confers resistance to nitrosative stress. Eukaryot. Cell 13: 494-503.
- Dutoit-Lefèvre, V., et al. 2015. An optimized fluorescence-based bidimensional immunoproteomic approach for accurate screening of autoantibodies. PLoS ONE 10: e0132142.
- Cho, H., et al. 2017. ENOblock, a unique small molecule inhibitor of the non-glycolytic functions of Enolase, alleviates the symptoms of type 2 diabetes. Sci. Rep. 7: 44186.
- Bakht, M.K., et al. 2018. Neuroendocrine differentiation of prostate cancer leads to PSMA suppression. Endocr. Relat. Cancer 26: 131-146.
- Jiang, S., et al. 2019. Acetylome profiling reveals extensive involvement of lysine acetylation in the conversion of muscle to meat. J. Proteomics 205: 103412.
- Tang, T., et al. 2020. Comparative proteomic and genomic analyses of Brucella abortus biofilm and planktonic cells. Mol. Med. Rep. 21: 731-743.
- Picciotto, S., et al. 2021. Isolation of extracellular vesicles from microalgae: towards the production of sustainable and natural nanocarriers of bioactive compounds. Biomater. Sci. 9: 2917-2930.
- Lim, J.S., et al. 2022. Mutual regulation between phosphofructokinase 1 platelet isoform and VEGF promotes glioblastoma tumor growth. Cell Death Dis. 13: 1002.

#### **RESEARCH USE**

For research use only, not for use in diagnostic procedures.