

14-3-3 γ (C-16): sc-731

BACKGROUND

14-3-3 proteins regulate many cellular processes relevant to cancer biology, notably apoptosis, mitogenic signaling and cell-cycle checkpoints. Seven isoforms comprise this family of signaling intermediates, denoted 14-3-3 β , γ , ϵ , ζ , η , θ and σ . 14-3-3 proteins form dimers that present two binding sites for ligand proteins, thereby bringing together two proteins that may not otherwise associate. These ligands largely share a 14-3-3 consensus binding motif and exhibit serine/threonine phosphorylation. 14-3-3 proteins function in broad regulation of these ligand proteins, by cytoplasmic sequestration, occupation of interaction domains and import/export sequences, prevention of degradation, activation/repression of enzymatic activity and facilitation of protein modification, and thus loss of expression contributes to a vast array of pathogenic cellular activities.

CHROMOSOMAL LOCATION

Genetic locus: YWHAG (human) mapping to 7q11.23, YWHAH (human) mapping to 22q12.3; Ywhag (mouse) mapping to 5 G2, Ywhah (mouse) mapping to 5 B1.

SOURCE

14-3-3 γ (C-16) is an affinity purified rabbit polyclonal antibody raised against a peptide mapping at the C-terminus of 14-3-3 γ of human origin.

PRODUCT

Each vial contains 200 μ g IgG in 1.0 ml of PBS with < 0.1% sodium azide and 0.1% gelatin.

Blocking peptide available for competition studies, sc-731 P, (100 μ g peptide in 0.5 ml PBS containing < 0.1% sodium azide and 0.2% BSA).

APPLICATIONS

14-3-3 γ (C-16) is recommended for detection of 14-3-3 γ and, to a lesser extent, 14-3-3 η of mouse, rat and human origin by Western Blotting (starting dilution 1:200, dilution range 1:100-1:1000), immunoprecipitation [1-2 μ g per 100-500 μ 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).

14-3-3 γ (C-16) is also recommended for detection of 14-3-3 γ and, to a lesser extent, 14-3-3 η in additional species, including equine, canine, bovine, porcine and avian.

Molecular Weight of 14-3-3 γ : 33 kDa.

Positive Controls: NIH/3T3 whole cell lysate: sc-2210, K-562 whole cell lysate: sc-2203 or 14-3-3 η (m): 293T Lysate: sc-117813.

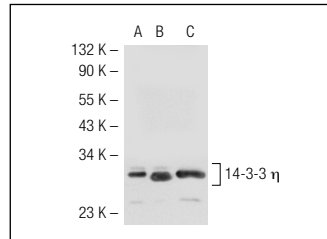
STORAGE

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

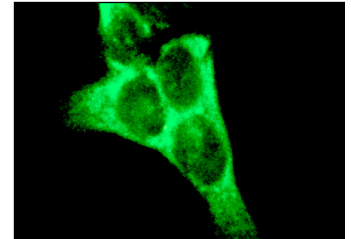
RESEARCH USE

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

DATA



14-3-3 γ (C-16): sc-731. Western blot analysis of 14-3-3 η expression in non-transfected 293T: sc-117752 (A), mouse 14-3-3 η transfected 293T: sc-117813 (B) and NIH/3T3 (C) whole cell lysates.



14-3-3 γ (C-16): sc-731. Immunofluorescence staining of methanol-fixed A-431 cells showing cytoplasmic localization.

SELECT PRODUCT CITATIONS

- Li, Y., et al. 2002. Regulation of TSC2 by 14-3-3 binding. *J. Biol. Chem.* 277: 44593-44596.
- Bolton, D.L., et al. 2008. 14-3-3 ζ binding to cell cycle regulatory factors is enhanced by HIV-1 Vpr. *Biol. Direct* 3: 17.
- Omi, K., et al. 2008. 14-3-3 ζ is indispensable for aggregate formation of polyglutamine-expanded huntingtin protein. *Neurosci. Lett.* 431: 45-50.
- Wang, J., et al. 2009. 14-3-3 ζ contributes to tyrosine hydroxylase activity in MN9D cells: Localization of dopamine regulatory proteins to mitochondria. *J. Biol. Chem.* 284: 14011-14019.
- Mangin, P.H., et al. 2009. Identification of five novel 14-3-3 isoforms interacting with the GPIIb-IX complex in platelets. *J. Thromb. Haemost.* 7: 1550-1555.
- Titus, M.A., et al. 2009. 14-3-3 η amplifies androgen receptor actions in prostate cancer. *Clin. Cancer Res.* 15: 7571-7581.
- Liang, X., et al. 2010. AS160 modulates aldosterone-stimulated epithelial sodium channel forward trafficking. *Mol. Biol. Cell* 21: 2024-2033.
- Zhong, J., et al. 2011. The interactome of a PTB domain-containing adapter protein, Odin, revealed by SILAC. *J. Proteomics* 74: 294-303.
- Lee, J.H. and Lu, H. 2011. 14-3-3 γ inhibition of MDMX-mediated p21 turnover independent of p53. *J. Biol. Chem.* 286: 5136-5142.
- Lopitz-Otsoa, F., et al. 2012. Integrative analysis of the ubiquitin proteome isolated using tandem ubiquitin binding entities (TUBEs). *J. Proteomics* 75: 2998-3014.


 MONOS
Satisfation
Guaranteed

Try **14-3-3 γ (D-6): sc-398423** or **14-3-3 γ (6A1): sc-69955**, our highly recommended monoclonal alternatives to 14-3-3 γ (C-16).