SANTA CRUZ BIOTECHNOLOGY, INC.

Omni-probe (M-21): sc-499

SANTA CRUZ BIOTECHNOLOGY

The Power to Question

BACKGROUND

Plasmid vectors for the expression of coding regions of eukaryotic genes in *E. coli*, insect and mammalian hosts are in common usage. Such expression vectors often encode hybrid fusion proteins containing part prokaryotic and part eukaryotic specified proteins. For instance, the prokaryotic pRSET A, B, C and pTrc His A, B, C; baculovirus pBlue Bac His A, B, C and mammalian pEBV His A, B, C expression vectors encode fusion proteins containing poly-histidine sequences fused to insertion cDNA sequences allowing for rapid purification on nickel-charged agarose resin. Omni-probe antibodies directed to specific vector sequences mapping between the polyhistidine tag and cDNA insert sequences provide a convenient and rapid means for identification of expressed fusion proteins in different hosts.

REFERENCES

1. Maniatis, T., et al. 1982. Molecular Cloning. Cold Spring Harbor, NY: Cold Spring Harbor Laboratory.
2. Nemenoff, R.A., et al. 1993. Phosphorylation and activation of a high molecular weight form of phospholipase A_2 by p42 microtubule-associated protein 2 kinase and protein kinase C. J. Biol. Chem. 268: 1960-1964.
3. Bowman, E.P., et al. 1993. Neutrophil phospholipase D is activated by a membrane-associated Rho family small molecular weight-GTP binding protein. J. Biol. Chem. 268: 21509-21512.

SOURCE

Omni-probe (M-21) is available as either rabbit (sc-499) or goat (sc-499-G) affinity purified polyclonal antibody raised against a peptide mapping between the (His)6 and polylinker sequence of Xpress series of prokaryotic and eukaryotic exspression vectors of origin.

PRODUCT

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

Blocking peptide available for competition studies, sc-499 P, (100 µg peptide in 0.5 ml PBS containing < 0.1% sodium azide and 0.2% BSA).

APPLICATIONS

Omni-probe (M-21) is recommended for detection of fusion proteins endoded by the prokaryotic pRSET A, B, C and pTrc His A, B, C; baculovirus pBlue Bac His A, B, C and mammalian pEBV His A, B, C expression vectors of N/A origin by Western Blotting (starting dilution 1:200, dilution range 1:100-1:1000) and immunoprecipitation [1-2 µg per 100-500 µg of total protein (1 ml of cell lysate)].

STORAGE

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

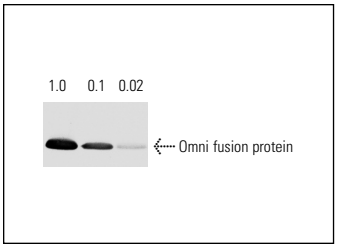
PROTOCOLS

See our web site at www.scbt.com or our catalog for detailed protocols and support products.

RESEARCH USE

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

DATA



Western blot analysis of an Omni fusion protein using Omni-probe (M-21): sc-499 antibody at IgG concentrations of 1.0 to 0.02 µg/ml.

SELECT PRODUCT CITATIONS

1. Nern, A., et al. 1998. A GTP-exchange factor required for cell orientation. Nature 391: 195-198.
2. Tanaka, T., et al. 2005. SLIM is a nuclear ubiquitin E3 ligase that negatively regulates Stat signaling. Immunity 22: 729-736.
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4. El-Shemerly, M., et al. 2008. ATR-dependent pathways control hExo1 stability in response to stalled forks. Nucleic Acids Res. 36: 511-519.
5. Dietschy, T., et al. 2009. p300-mediated acetylation of the Rothmund-Thomson-syndrome gene product RECQL4 regulates its subcellular localization. J. Cell Sci. 122: 1258-1267.
6. Nikolaienko, O., et al. 2009. Intersectin 1 forms a complex with adaptor protein Ruk/CIN85 *in vivo* independently of epidermal growth factor stimulation. Cell. Signal. 21: 753-759.
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