PAK1 PBD Agarose Beads

CATALOG NUMBER: STA-411 STORAGE: -20°C

QUANTITY AND CONCENTRATION: 800 μL of 50% Agarose slurry, 400 μg PAK1-PBD in 1X PBS, 50% Glycerol

SHELF LIFE: 1 year from receipt under proper storage conditions; avoid multiple freeze thaw cycles

Background

Small GTP-binding proteins (or GTPases) are a family of proteins that serve as molecular regulators in signaling transduction pathways. Rac, a 21 kDa protein, belongs to the family of Rho GTPases regulating a variety of biological response pathways that include cell motility, cell division, gene transcription, and cell transformation. Like other small GTPases, Rac regulates molecular events by cycling between an inactive GDP-bound form and an active GTP-bound form. In its active (GTP-bound) state, Rac binds specifically to the p21-binding domain (PBD) of p21-activated protein kinase (PAK) to control downstream signaling cascades.

Presentation

PAK PBD Agarose beads, in color, are easy to visualize, minimizing potential loss during washes and aspirations of Rac-GTP pulldown (Figure 1).



Figure 1: PAK-PBD Beads in Color



Activity

Product specifically interacts and precipitaes GTP-bound Rac or Cdc 42 from cell lysate (Figures 2 & 3).

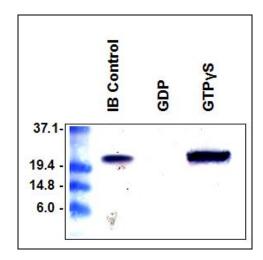


Figure 2: Rac Activation Assay. *Lane 1*, GTPase Immunoblot Positive Control. *Lane 2*, 293 cell lysate loaded with GDP and incubated with PAK PBD Agarose beads. *Lane 3*, 293 cell lysate loaded with GTPγS and incubated with PAK-1 PBD Agarose beads. Samples were immunoblotted with anti-Rac antibody.

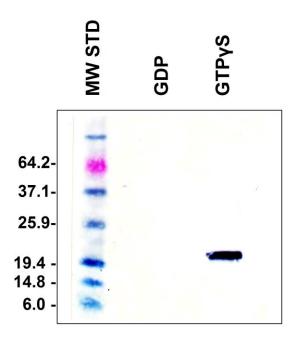


Figure 3: Cdc42 Activation Assay. *Lane 1*, MW Standard. *Lane 2*, 293 cell lysate loaded with GDP and incubated with PAK PBD Agarose beads. *Lane 3*, 293 cell lysate loaded with GTPγS and incubated with PAK-1 PBD Agarose beads. Samples were immunoblotted with anti-Cdc42 antibody.



References

- 1. Raftopoulou M., and Hall A. (2004) *Dev Biol.* **265**: 23-32.
- 2. Bar-Sagi D., and Hall A. (2000) Cell 103: 227-38.
- 3. Benard, V., Bohl, B. P., and Bokoch, G. M. (1999) J. Biol. Chem. 274, 13198-13204.

Recent Product Citations

- 1. Crespo, G.V. et al. (2023). The Rac inhibitor HV-107 as a potential therapeutic for metastatic breast cancer. *Mol Med.* **29**(1):75. doi: 10.1186/s10020-023-00678-7.
- 2. Bertrand-Chapel, A. et al. (2022). SMAD2/3 mediate oncogenic effects of TGF-β in the absence of SMAD4. *Commun Biol.* **5**(1):1068. doi: 10.1038/s42003-022-03994-6.
- 3. Bernadzki, K. M. et al. (2020). Arhgef5 Binds α-Dystrobrevin 1 and Regulates Neuromuscular Junction Integrity. *Front Mol Neurosci*. doi: 10.3389/fnmol.2020.00104.
- 4. Speranza, L. et al. (2017). Serotonin 5-HT7 receptor increases the density of dendritic spines and facilitates synaptogenesis in forebrain neurons. *J Neurochem*. doi: 10.1111/jnc.13962.
- 5. Bijata, M. et al. (2016). Dystroglycan controls dendritic morphogenesis of hippocampal neurons in vitro. *Front Cell Neurosci*. doi:10.3389/fncel.2015.00199.
- 6. Fusté, N. P. et al. (2016). Cytoplasmic cyclin D1 regulates cell invasion and metastasis through the phosphorylation of paxillin. *Nat Commun*. doi:10.1038/ncomms11581.
- 7. Alam, J. et al. (2014). N-acetylcysteine and the human serum components that inhibit bacterial invasion of gingival epithelial cells prevent experimental periodontitis in mice. *J Periodontal Implant Sci.* 44:266-273.
- 8. Morrison, A. R. et al. (2014). Chemokine-coupled β2 integrin-induced macrophage Rac2-myosin IIA interaction regulates VEGF-A mRNA stability and arteriogenesis. *J Exp Med.* **211**:1957-1968.
- 9. Pothula, S. et al. (2013). Regulation of Cdc42 expression and signaling is critical for promoting corneal epithelial wound healing. *Invest. Ophthalmol. Vis. Sci.* **54**: 5343-5352.
- 10. Cheng, J. et al.(2010).FSP-1 silencing in bone marrow cells suppresses neointima formation in vein graft. *Circ Res.* **110**:230-240.
- 11. Sabbatini, M. E. et al. (2010). CCK activates RhoA and Rac1 differentially through G-alpha-13 and G-alpha-q in mouse pancreatic acini. *Am. J. Physiol. Cell Physiol.* **298**:C592-C605.
- 12. Zhang, Q-G. et al. (2009). Estrogen attenuates ischemic oxidative damage via an estrogen receptor alpha-mediated inhibition of NADPH oxidase activation. *J. Neurosci.* **29**:13823-13836.

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Contact Information

Cell Biolabs, Inc. 7758 Arjons Drive San Diego, CA 92126

Worldwide: +1 858-271-6500 USA Toll-Free: 1-888-CBL-0505 E-mail: tech@cellbiolabs.com

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