

α PAK (N-20): sc-882



The Power to Question

BACKGROUND

Three isoforms of serine/threonine kinases, designated α PAK p68, β PAK p65 and γ PAK p62, have been shown to exhibit a high degree of sequence homology with the *S. cerevisiae* kinase Ste 20, involved in pheromone signaling. The α , β and γ PAK isoforms complex specifically with Rac1 and Cdc42 in their active GTP-bound state, inhibiting their intrinsic GTPase activity leading to their autophosphorylation. There are eight sites of autophosphorylation on γ PAK, including Ser 19, Ser 141 and Thr 402, and phosphorylation of Ser 141 and Thr 402 is correlated with γ PAK activation. Once phosphorylated and their affinity for Rac/Cdc42 reduced, the PAK isoforms disassociate from the complex to seek downstream substrates. One such putative substrate is Mek kinase, an upstream effector of Mek4 which is involved in the JNK signaling pathway. While the PAK isoforms interact in a GTP-dependent manner with Rac1 and Cdc42, they do not interact with Rho.

CHROMOSOMAL LOCATION

Genetic locus: PAK1 (human) mapping to 11q13.5; Pak1 (mouse) mapping to 7 E2.

SOURCE

α PAK (N-20) is an affinity purified rabbit polyclonal antibody raised against a peptide mapping at the N-terminus of α PAK of rat origin.

PRODUCT

Each vial contains 100 μ g IgG in 1.0 ml of PBS with < 0.1% sodium azide and 0.1% gelatin.

α PAK (N-20) is available conjugated to agarose (sc-882 AC), 500 μ g/0.25 ml agarose in 1 ml, for IP.

Blocking peptide available for competition studies, sc-882 P, (100 μ g peptide in 0.5 ml PBS containing < 0.1% sodium azide and 0.2% BSA).

APPLICATIONS

α PAK (N-20) is recommended for detection of α PAK p68 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) and solid phase ELISA (starting dilution 1:30, dilution range 1:30-1:3000). α PAK (N-20) is also recommended for detection of α PAK p68 in additional species, including equine, canine, bovine and porcine.

Suitable for use as control antibody for α PAK siRNA (h): sc-29700, α PAK siRNA (m): sc-29701, α PAK shRNA Plasmid (h): sc-29700-SH, α PAK shRNA Plasmid (m): sc-29701-SH, α PAK shRNA (h) Lentiviral Particles: sc-29700-V and α PAK shRNA (m) Lentiviral Particles: sc-29701-V.

Molecular Weight of α PAK: 65 kDa.

Positive Controls: NIH/3T3 whole cell lysate: sc-2210, F9 cell lysate: sc-2245 or 3T3-L1 cell lysate: sc-2243.

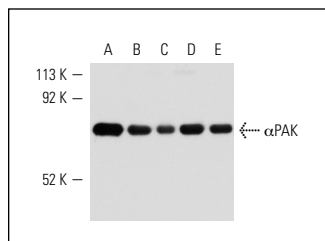
RESEARCH USE

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

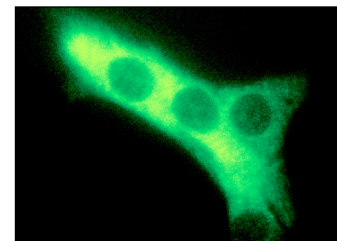
STORAGE

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

DATA



α PAK (N-20): sc-882. Western blot analysis of α PAK expression in EOC 20 (A), NIH/3T3 (B), F9 (C), 3T3-L1 (D) and C3H/10T1/2 (E) whole cell lysates.



α PAK (N-20): sc-882. Immunofluorescence staining of methanol-fixed NIH/3T3 cells showing cytoplasmic localization.

SELECT PRODUCT CITATIONS

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- Romero, A.M., et al. 2010. Chronic ethanol exposure alters the levels, assembly, and cellular organization of the actin cytoskeleton and microtubules in hippocampal neurons in primary culture. *Toxicol. Sci.* 118: 602-612.
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- Ming, J., et al. 2011. Interleukin-7 up-regulates cyclin D1 via activator protein-1 to promote proliferation of cell in lung cancer. *Cancer Immunol. Immunother.* 61: 79-88.
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- Itakura, A., et al. 2013. p21-Activated kinase (PAK) regulates cytoskeletal reorganization and directional migration in human neutrophils. *PLoS ONE* 8: e73063.
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