

p- α PAK (Thr 423)-R: sc-21903-R

BACKGROUND

p21-activated kinases (PAK) are serine/threonine kinases that link Rho GTPases to cytoskeletal reorganization and nuclear signaling. Three common isoforms are PAK1, PAK2, and PAK3 (also known as α PAK p68, β PAK p65 and γ PAK p62, respectively). α , β and γ PAK isoforms associate with Rac 1 and Cdc42 in their active GTP-bound state, inhibiting their intrinsic GTPase activity and mediating their autophosphorylation. γ PAK can undergo phosphorylation on Ser 19, Ser 141 and Thr 402, and phosphorylation of Ser 141 and Thr 402 correlates with γ PAK activation. Autophosphorylation of α PAK Thr 423 (Thr 402 for β PAK and Thr 421 for γ PAK) is catalyzed by Cdc42 and is required for kinase activation of PAK. Once phosphorylated and their affinity for Rac/Cdc42 reduced, PAK isoforms disassociate from the complex to seek downstream substrates. One such substrate is MEK kinase, an upstream effector of MEK4 which is involved in the JNK signaling pathway.

REFERENCES

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- Boguski, M.S. and McCormick, F. 1993. Proteins regulating Ras and its relatives. *Nature* 366: 643-654.
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- Minden, A., Lin, A., McMahon, M., Lange-Carter, C., Derijard, B., Davis, R.J., Johnson, G.L. and Karin, M. 1994. Differential activation of ERK and JNK mitogen-activated protein kinases by Raf-1 and MEK. *Science* 266: 1719-1723.

CHROMOSOMAL LOCATION

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

SOURCE

p- α PAK (Thr 423)-R is a rabbit polyclonal antibody raised against a short amino acid sequence containing Thr 423 phosphorylated α PAK of mouse origin.

STORAGE

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

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-21903 P, (100 μ g peptide in 0.5 ml PBS containing < 0.1% sodium azide and 0.2% BSA).

APPLICATIONS

p- α PAK (Thr 423)-R is recommended for detection of Thr 423 phosphorylated α PAK of mouse, rat and human origin by Western Blotting (starting dilution 1:200, dilution range 1:100-1:1000), 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).

p- α PAK (Thr 423)-R is also recommended for detection of correspondingly phosphorylated α PAK in additional species, including equine, canine, bovine, porcine and avian.

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 p- α PAK: 65 kDa.

RECOMMENDED SECONDARY REAGENTS

To ensure optimal results, the following support (secondary) reagents are recommended: 1) Western blotting: use goat anti-rabbit IgG-HRP: sc-2004 (dilution range: 1:2000-1:100,000) or Cruz Marker™ compatible goat anti-rabbit IgG-HRP: sc-2030 (dilution range: 1:2000-1:5000), Cruz Marker™ Molecular Weight Standards: sc-2035, TBS Blotto B Blocking Reagent: sc-2335 (use 50 mM NaF, sc-24988, as diluent), Western Blotting Luminol Reagent: sc-2048 and Lambda Phosphatase: sc-200312A. 2) Immunofluorescence: use goat anti-rabbit IgG-FITC: sc-2012 (dilution range: 1:100-1:400) or goat anti-rabbit IgG-TR: sc-2780 (dilution range: 1:100-1:400) with UltraCruz™ Mounting Medium: sc-24941.

SELECT PRODUCT CITATIONS

- Gustafson, B. and Smith, U. 2006. Cytokines promote Wnt signaling and inflammation and impair the normal differentiation and lipid accumulation in 3T3-L1 preadipocytes. *J. Biol. Chem.* 281: 9507-9516.

RESEARCH USE

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



Try p- α PAK (17.Thr 423): sc-135755 or p- α PAK (66.Thr 423): sc-135754, our highly recommended monoclonal alternatives to p- α PAK (Thr 423).