For Research Use Only

Phospho-PAK1 (Ser144)/PAK2 (Ser141) Recombinant antibody, PBS Only

Catalog Number:85044-1-PBS



Purification Method:

Protein A purfication

CloneNo.:

242123D12

Basic Information

Catalog Number:

85044-1-PBS

Size: 1 mg/ml

Source:

55

Rabbit Isotype: IgG

p21 protein (Cdc42/Rac)-activated kinase 1

> Calculated MW: 553 aa, 62 kDa

BC109299

5058

Q13153 Full Name:

GeneID (NCBI):

UNIPROT ID:

GenBank Accession Number:

Observed MW: 61 kDa

Applications

Tested Applications: WB, Indirect ELISA Species Specificity: human, mouse, rat

Background Information

The human PAK family is divided into the group I (PAK1 to PAK3) and group II (PAK4 to PAK6). Group I PAK share some domains that are not present in the group II members. In particular, the autoinhibitory domain (AID) is important for regulation of the kinase activity of the group I family members. Autophosphorylation at PAK1 Ser144, or at the equivalent sites for the other PAK, stabilizes the open conformation and sustains high kinase activity. Mutation of tyrosines 131 or 429 is associated with reduced dimerization and enhanced kinase activity. (PMID: 31748572)

Storage

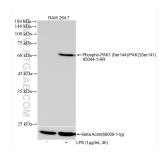
Storage:

Store at -80°C.

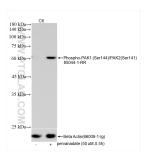
The product is shipped with ice packs. Upon receipt, store it immediately at -80°C Storage Buffer:

PBS Only

Selected Validation Data



Non-treated RAW 264.7 cells and LPS treated RAW 264.7 cells were subjected to SDS PAGE followed by western blot with 85044-1-RR (Phospho-PAK1 (Ser144)/PAK2 (Ser141) antibody) at dilution of 1:2000 incubated at room temperature for 1.5 hours. The membrane was stripped and re-blotted with Beta Actin (66009-1-ig) antibody as a loading control. This data was developed using the same antibody clone with 85044-1-PBS in a different storage buffer formulation.



Non-treated C6 cells and pervanadate treated C6 cells were subjected to SDS PAGE followed by western blot with 85044-1-RR (Phospho-PAK1 (Ser144)/PAK2 (Ser141) antibody) at dilution of 1:2000 incubated at room temperature for 1.5 hours. The membrane was stripped and re-blotted with Beta Actin (66009-1-lg) antibody as a loading control. This data was developed using the same antibody clone with 85044-1-PBS in a different storage buffer formulation.