

Phospho-JAK2 (Tyr570) Ab

Cat.#: AF3025 Concn.: 1mg/ml Mol.Wt.: 125kDa Size: 100ul,200ul Source: Rabbit Clonality: Polyclonal

Application: WB 1:500-1:2000, IF/ICC 1:100-1:500

Reactivity: Human, Mouse, Rat

Purification: The Ab is from purified rabbit serum by affinity purification

via sequential chromatography on phospho- and non-

phospho-peptide affinity columns.

Specificity: Phospho-JAK2 (Tyr570) Ab detects endogenous levels of JAK2

only when phosphorylated at Tyrosine 570.

Immunogen: A synthesized peptide derived from human JAK2 around the

phosphorylation site of Tyrosine 570.

Uniprot: 060674

Description: This gene product is a protein tyrosine kinase involved in a

specific subset of cytokine receptor signaling pathways. It has been found to be constituitively associated with the prolactin receptor and is required for responses to gamma

interferon.

Subcellular Location: Endomembrane system. Nucleus.

Tissue Specificity: Ubiquitously expressed throughout most tissues.

Similarity: Possesses 2 protein kinase domains. The second one

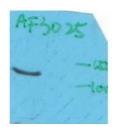
probably contains the catalytic domain, while the presence of slight differences suggest a different role for protein kinase 1 (By similarity). Belongs to the protein kinase superfamily. Tyr protein kinase family. JAK subfamily.

Storage Condition and

Buffer:

Rabbit IgG in phosphate buffered saline, pH 7.4, 150mM NaCl, 0.02% sodium azide and 50% glycerol. Store at -20

°C.Stable for 12 months from date of receipt.



Western blot analysis of Phospho-JAK2 (Tyr570) Ab expression in etoposide treated 293 cells lysates. The lane on the right is treated with the antigen-specific peptide.



Affinity Biosciences website: www.affbiotech.com order: order@affbiotech.com



Western blot analysis of JAK2 phosphorylation expression in etoposide treated 293 whole cell lysates, The lane on the left is treated with the antigen-specific peptide.



AF3025 staining K562 by IF/ICC. The sample were fixed with PFA and permeabilized in 0.1% Triton X-100,then blocked in 10% serum for 45 minutes at 25°C. The primary Ab was diluted at 1/200 and incubated with the sample for 1 hour at 37°C. An Alexa Fluor 594 conjugated goat anti-rabbit IgG (H+L) Ab, diluted at 1/600, was used as the secondary Ab.

<code>IMPORTANT:</code> For western blot, incubate membrane with diluted primary Ab in 5% w/v milk , 1% TBS, 0.1% Tween@20 at 4°C with gentle shaking, overnight.

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