

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

STAT4 Ab

Cat.#: AF6441 Concn.: 1mg/ml Mol.Wt.: 86kDa Size: 100ul,200ul Source: Rabbit Clonality: Polyclonal

Application: WB 1:500-1:2000 IHC 1:50-1:200 IP, IF/ICC 1:100-1:500

Reactivity: Human, Mouse, Rat

Purification: The antiserum was purified by peptide affinity

chromatography using SulfoLink™ Coupling Resin (Thermo

Fisher Scientific).

Specificity: STAT4 Ab detects endogenous levels of total STAT4.

Immunogen: A synthesized peptide derived from human STAT4.

Uniprot: Q14765

Description: The protein encoded by this gene is a member of the STAT

family of transcription factors. In response to cytokines and growth factors, STAT family members are phosphorylated by the receptor associated kinases, and then form homo- or heterodimers that translocate to the cell nucleus where they

act as transcription activators.

Subcellular Location: Cytoplasm. Nucleus. Translocated into the nucleus in

response to phosphorylation.

Similarity: Belongs to the transcription factor STAT family.

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 extracts from various samples, using STAT4 Ab.

Lane 1: RAW264.7 cells, treated with blocking peptide;

Lane 2: RAW264.7 cells;

Lane 3: Hela cells.



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Western blot analysis of STAT4 expression in HeLa whole cell lysates, The lane on the left was treated with the antigenspecific peptide.



AF6441 at 1/100 staining Human breast cancer tissue by IHC-P. The sample was formaldehyde fixed and a heat mediated antigen retrieval step in citrate buffer was performed. The sample was then blocked and incubated with the Ab for 1.5 hours at 22°C. An HRP conjugated goat anti-rabbit Ab was used as the secondary.



AF6441 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 , 1X TBS, 0.1% Tween@20 at 4°C with gentle shaking, overnight.

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