Epo-R (Phospho-Tyr456) Ab

Cat.#: AF8436 Concn.: 1mg/ml Mol.Wt.: 55kDa Size: 50ul,100ul,200ul Source: Rabbit Clonality: Polyclonal

Application: WB 1:1000-3000, 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: Epo-R (Phospho-Tyr456) Ab detects endogenous levels of

Epo-R only when phosphorylated at Tyr456

Immunogen: A synthesized peptide derived from human Epo-R (Phospho-

Tyr456)

Uniprot: P19235

Subcellular Location: Plasma membrane; Extracellular region or secreted;

Tissue Specificity: Erythroid cells and erythroid progenitor cells. Isoform EPOR-F

is the most abundant form in EPO-dependent erythroleukemia cells and in late-stage erythroid

progenitors. Isoform EPOR-S and isoform EPOR-T are the predominant forms in bone marrow. Isoform EPOR-T is the most abundant from in early-stage erythroid progenitor

cells.

Similarity: The WSXWS motif appears to be necessary for proper

protein folding and thereby efficient intracellular transport and cell-surface receptor binding. The box 1 motif is required for JAK interaction and/or activation. Contains 1 copy of a cytoplasmic motif that is referred to as the immunoreceptor tyrosine-based inhibitor motif (ITIM). This motif is involved in modulation of cellular responses. The phosphorylated ITIM motif can bind the SH2 domain of several SH2-containing phosphatases. Belongs to the type I cytokine receptor family.

Type 1 subfamily.

Storage Condition and

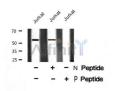
Buffer:

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



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Western blot analysis Epo-R (Phospho-Tyr456) using Jurkat whole cell lysates



AF8436 staining Hela 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.

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

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