

## Phospho-5 Lipoxygenase (Ser664) Ab

[References\(1\)](#) [Images\(4\)](#)

Cat.#: AF8359  
 Size: 100ul,200ul

Concn.: ~1mg/ml  
 Source: Rabbit

Mol.Wt.: 77kDa  
 Clonality: Polyclonal

**Application:** WB 1:1000-3000, IF/ICC 1:100-1:500, ELISA(peptide) 1:20000-1:40000  
 \*The optimal dilutions should be determined by the end user.

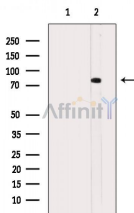
**Reactivity:** Human,Mouse,Rat

**Purification:** The Ab is from purified rabbit serum by affinity purification via sequential chromatography on phospho-peptide and non-phospho-peptide affinity columns.

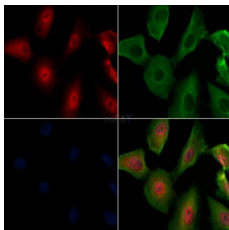
**Immunogen:** A synthesized peptide derived from human 5 Lipoxygenase around the phosphorylation site of Ser664.

**Uniprot:** P09917

**Storage:** 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 PC12 cells, using Phospho-5 Lipoxygenase (Ser664) Ab. The lane on the left was treated with blocking peptide.



AF8359 staining A549 cells(H2O2 treatment) by IF/ICC. The samples were fixed with PFA and permeabilized in 0.1% Triton X-100, then blocked in 10% serum for 45 minutes at 25°C. Samples were then incubated with primary Ab(#AF8359) and mouse anti-beta tubulin Ab(#T0023) for 1 hour at 37°C. An AlexaFluor594 conjugated goat anti-rabbit IgG Ab(Red) and an AlexaFluor488 conjugated goat anti-mouse IgG Ab(Green) were used as the secondary Ab.

The nuclear counter stain is DAPI (blue).

**IMPORTANT:** 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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