

## Phospho-EZH2 (Ser311) Ab

[Images\(3\)](#)

Cat.#: AF3583	Concn.: ~1mg/ml	Mol.Wt.: 85kD, 100kD
Size:	Source: Rabbit	Clonality: Polyclonal

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

\*The optimal dilutions should be determined by the end user.

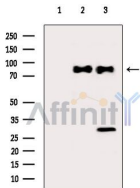
Reactivity: Human,Mouse,Rat

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.

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 EZH2 around the phosphorylation site of Ser311.

Uniprot: Q15910

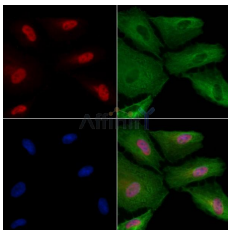


Western blot analysis of extracts from various samples, using Phospho-EZH2 (Ser311) Ab.

Lane 1: Hepg2 cells (serum starvation treatment), blocked with antigen-specific peptides.

Lane 2: Hepg2 cells (serum starvation treatment).

Lane 3: Raw264.7 cells (heat-shock treatment).



AF3583 staining A549 cells (4h of LPS 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 (AF3583) and mouse anti-beta tubulin Ab (T0023) for 1 hour at 37°C. An AlexaFluor594 conjugated goat anti-rabbit IgG(H+L) Ab (Red) and an AlexaFluor488 conjugated goat anti-mouse IgG(H+L) 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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