

## Phospho-Calnexin (Ser564) Ab

[Images\(3\)](#)

Cat.#: AF2438                                      Concn.: ~1mg/ml                                      Mol.Wt.: 90kDa  
 Size:    Source: Rabbit    Clonality: Polyclonal

Application:                                      WB 1:500-1:2000, IHC 1:50-1:200, IF/ICC 1:100-1:500  
 \*The optimal dilutions should be determined by the end user.

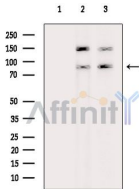
Reactivity:                                      Human,Mouse,Rat

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.

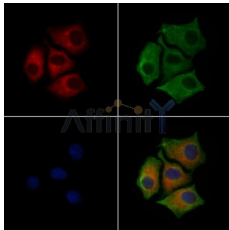
Purification:                                      The Ab is from purified rabbit serum by affinity purification via sequential chromatography on phospho-peptide and non-phospho-peptide affinity columns.

Immunogen:                                      A synthesized peptide derived from human Calnexin around the phosphorylation site of Ser564.

Uniprot:                                      P27824



Western blot analysis of extracts from HepG2 cells(heat-shock treatment) Hela cells(heat-shock treatment), using Phospho-Calnexin (Ser564) Ab. The lane on the left was treated with blocking peptide.



AF2438 staining HepG2 cells(30min of 4uM Forskolin 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(AF2438) 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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