

## Phospho-Glycogen Synthase (Ser645) Ab

Cat.#: AF3165  
Size: 100ul, 200ul

Concn.: 1mg/ml  
Source: Rabbit

Mol.Wt.: 83,89kDa  
Clonality: Polyclonal

Application: WB 1:500-1:2000 IHC 1:50-1:200, 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: Phospho-Glycogen Synthase (Ser645) Ab detects endogenous levels of Glycogen Synthase only when phosphorylated at Serine 645.

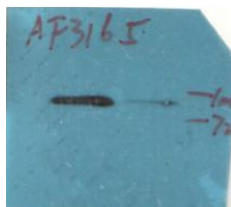
Immunogen: A synthesized peptide derived from human Glycogen Synthase around the phosphorylation site of Serine 645.

Uniprot: P13807

Description: GYS1 muscle glycogen synthase 1. Transfers glucosyl residue from UDP-glucose to glycogen. Regulated allosterically by glucose-6-phosphate, and by PKA-mediated phosphorylation.

Similarity: Belongs to the glycosyltransferase 3 family.

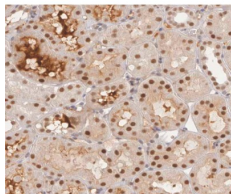
Storage Condition and Buffer: Rabbit IgG in phosphate buffered saline, pH 7.4, 150mM NaCl, 0.02% sodium azide and 50% glycerol. Store at -20 °C. Stable for 12 months from date of receipt.



Western blot analysis of Phospho-Glycogen Synthase (Ser645) Ab expression in PMA treated NIH-3T3 cells lysates. The lane on the right is treated with the antigen-specific peptide.



Western blot analysis of Glycogen Synthase phosphorylation expression in PMA treated NIH-3T3 whole cell lysates. The lane on the left is treated with the antigen-specific peptide.



AF3165 at 1/200 staining human kidney tissue sections by IHC-P. The tissue was formaldehyde fixed and a heat mediated antigen retrieval step in citrate buffer was performed. The tissue 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.



AF3165 staining NIH-3T3 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 primary Ab in 5% w/v milk, 1X TBS, 0.1% Tween@20 at 4°C with gentle shaking, overnight.

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