

## Mouse Monoclonal Antibody to

# GSK3beta (phospho-Ser 9)

## clone 2D3

0098-100/GSK3b-2D3 Order No.:

100 Size (µg) 0098S Lot No.:



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05/060418F

Isotype	Species Reactivity	Applications	Mol. Weight	Ref.Cell Line	Epitope	Immunogen
lgG1	human, mouse	WB, ELISA	46 kDa	A431	R T T pS F A E (phosphoserine 9)	phosphopeptide conjugated to KLH

### **Background and Specificity:**

Glycogen Synthase Kinase 3 beta (GSK3beta) is a serine/threonine kinase that is acting independently of most signal transduction pathways. The activity of GSK3beta can be modulated by phosphorylation of Ser 9 and Tyr 216 in the activation loop. GSK3beta plays a crucial role in phosphorylation of beta-catenin and thus is an important kinase involved in the regulation of beta-catenin/wnt-signaling.

Mab GSK3b-2D3 specifically recognizes GSK3beta phosphorylated at Ser 9 at 46 kDa in Western blot and ELISA applications. The antibody does not crossreact with GSK3alpha pSer21.

# **Related Products**

mab to GSK3alpha

#0211-100/GSK3a-10C1

mab to GSK3alpha (phospho-Ser21)

mab to GSK3beta #0135-100/GSK3b-11B9

mab to GSK3beta (phospho-Ser9)

#0185-100/GSK3b-3A8

mab to GSK3beta (phospho-Tyr215) mab to GSK3alpha (phospho-Tyr278) #0258-100/GSK3b-6D3

**Purification:** 

The antibody was purified from serum-free cell culture supernatant by subsequent thiophilic adsorption and size

exclusion chromatography.

liquid; 0.1mg/ml in in PBS/0.09% Na-Azide/PEG and Formulation:

Sucrose/50% Glycerol

Reconstitution:

Aliquote and store at -20°C up to 1 year. Stability:

Avoid repeated freeze / thaw cycles.

#0832: Cell lysate from pervanadate treated A431 cells. **Positive Control:** 

Immunoblotting: 0.5 µg/ml for HRPO/ECL detection

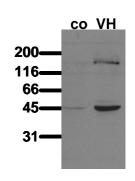
Recommended blocking buffer: Casein/Tween 20 based blocking and blot incubation buffer, e.g. nanoTools product

#3031-500/CPPT or #3031-3000/CPPT.

Immunoprecipitation: ND ND Immunocytochemistry:

use at 0.05 µg/ml ELISA:

> All products are supplied for research and investigational use only. Not for use in humans or laboratory animals.



#### **Phosphospecificity**

Whole cell extracts of control (co) and pervanadate (VH) treated A431 tumor cells were applied to SDS-PAGE (ca 20.000 cells per lane) and transferred to PVDF membranes. Immunoblots were probed with mab GSK3b-2D3 (0.5 µg/ ml) for 1h at RT and developed by ECL (exp. time: 30 sec).