## #66350 Store at -20C

## PHGDH (D8F3O) Rabbit mAb



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3 Trask Lane | Danvers | Massachusetts | 01923 | USA

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<b>Applications:</b> WB, IP	Reactivity: H	Sensitivity: Endogenous	<b>MW (kDa):</b> 57	Source/Isotype: Rabbit IgG	UniProt ID: #O43175	Entrez-Gene Id 26227	
Product Usage Information	Application			Dilution			
	We	estern Blotting			1:1000		
	Imi	munoprecipitation		1:50			
Storage		Supplied in 10 mM sodium HEPES (pH 7.5), 150 mM NaCl, 100 $\mu$ g/ml BSA, 50% glycerol and less than 0.02% sodium azide. Store at –20°C. Do not aliquot the antibody.					
Specificity / Sensitivity		PHGDH (D8F3O) Rabbit mAb recognizes endogenous levels of total PHGDH protein.					
Source / Purification	-	Monoclonal antibody is produced by immunizing animals with a synthetic peptide corresponding to residues surrounding Val304 of human PHGDH protein.					
Background	pho prov glut	Mammalian cells synthesize serine <i>de novo</i> by diverting a portion of the glycolytic intermediate 3-phosphoglycerate into the phosphorylated pathway of serine synthesis. This shift supports anabolism by providing precursors for the biosynthesis of proteins, nucleotides, creatine, porphyrins, phospholipids, and glutathione. Phosphoglycerate dehydrogenase (PHGDH) catalyzes the first step in the serine biosynthesis pathway by converting 3-phosphoglycerate into phosphohydroxy pyruvate (1).					
	can mel leve evic	Research studies demonstrate that an increase in serine biosynthesis supports growth and proliferation of cancer cells (2-4), which is supported by amplification and overexpression of PHGDH in a subset of melanoma and breast cancers (5,6). Suppression of PHGDH expression in cell lines with elevated PHGDH levels causes a strong decrease in cell proliferation and inhibits tumor growth <i>in vivo</i> (5). Additional evidence suggests that PHGDH interacts with and stabilizes FoxM1, which promotes the proliferation, invasion, and tumorigenicity of glioma cells (7).					
Background Referer	2. A 3. M 4. M 5. P 6. L	<ol> <li>Locasale, J.W. (2013) <i>Nat Rev Cancer</i> 13, 572-83.</li> <li>Amelio, I. et al. (2013) <i>Oncogene</i>, [Epub ahead of print].</li> <li>Ma, L. et al. (2013) <i>Cell</i> 152, 599-611.</li> <li>Maddocks, O.D. et al. (2013) <i>Nature</i> 493, 542-6.</li> <li>Possemato, R. et al. (2011) <i>Nature</i> 476, 346-50.</li> <li>Locasale, J.W. et al. (2011) <i>Nat Genet</i> 43, 869-74.</li> <li>Liu, J. et al. (2013) <i>J Neurooncol</i> 111, 245-55.</li> </ol>					

**Species Reactivity** 

Species reactivity is determined by testing in at least one approved application (e.g., western blot).

Western Blot Buffer

IMPORTANT: For western blots, incubate membrane with diluted primary antibody in 5% w/v nonfat dry milk, 1X TBS, 0.1% Tween® 20 at  $4^{\circ}$ C with gentle shaking, overnight.

**Applications Key** 

WB: Western Blotting IP: Immunoprecipitation

**Cross-Reactivity Key** 

H: human M: mouse R: rat Hm: hamster Mk: monkey Vir: virus Mi: mink C: chicken Dm: D. melanogaster

X: Xenopus Z: zebrafish B: bovine Dg: dog Pg: pig Sc: S. cerevisiae Ce: C. elegans Hr: horse

GP: Guinea Pig Rab: rabbit All: all species expected

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