

PHGDH Antibody



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For Research Use Only. Not for Use in Diagnostic Procedures.

Applications: W	Reactivity: H M R	Sensitivity: Endogenous	MW (kDa): 57	Source/Isotype: Rabbit	UniProt ID: #043175	Entrez-Gene Id: 26227
Product Usage Information		Application Western Blotting			Dilution 1:1000	
Storage		Supplied in 10 mM so 20°C. Do not aliquot t		s), 150 mM NaCl, 100 μg	/ml BSA and 50% gl	ycerol. Store at –
Specificity/Sensitivity		PHGDH Antibody recognizes endogenous levels of total PHGDH protein.				
Source / Purification		Polyclonal antibodies are produced by immunizing animals with a synthetic peptide corresponding to residues surrounding Gln29 of human PHGDH protein. Antibodies are purified by protein A and peptide affinity chromatography.				
Background		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).				
		proliferation of cancer a subset of melanoma elevated PHGDH level	r cells (2-4), which is a and breast cancer s causes a strong d ce suggests that PH	crease in serine biosynth s supported by amplifica s (5,6). Suppression of P ecrease in cell proliferat GDH interacts with and y of glioma cells (7).	tion and overexpre HGDH expression in ion and inhibits tun	ssion of PHGDH in n cell lines with nor growth <i>in vivo</i>
Background References		 Locasale, J.W. (2013) Nat Rev Cancer 13, 572-83. Amelio, I. et al. (2013) Oncogene, [Epub ahead of print]. Ma, L. et al. (2013) Cell 152, 599-611. Maddocks, O.D. et al. (2013) Nature 493, 542-6. Possemato, R. et al. (2011) Nature 476, 346-50. Locasale, J.W. et al. (2011) Nat Genet 43, 869-74. Liu, J. et al. (2013) J Neurooncol 111, 245-55. 				
Species Reactivity		Species reactivity is de	etermined by testin	g in at least one approve	ed application (e.g.,	western blot).

Western Blot Buffer

Applications Key

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°C with gentle shaking, overnight.

W: Western Blotting

Cross-Reactivity Key

H: Human M: Mouse R: Rat

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