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Phospho-PKM2 (Tyr105) Antibody



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Applications: WB	Reactivity: H M R Mk	Sensitivity: Endogenous	MW (kDa): 60	Source: Rabbit	UniProt ID: #P14618	Entrez-Gene Id: 5315	
							Product Usage Information
We	estern Blotting		1:1000				
Storage		Supplied in 10 mM sodium HEPES (pH 7.5), 150 mM NaCl, 100 μ g/ml BSA and 50% glycerol. Store at – 20°C. Do not aliquot the antibody.					
Specificity / Sensitiv	,	Phospho-PKM2 (Tyr105) Antibody detects endogenous levels of PKM2 protein only when phosphorylated at Tyr105. This antibody may slightly cross react with PKM1 phosphorylated at the equivalent site.					
Source / Purification	Pol	Polyclonal antibodies are produced by immunizing animals with a synthetic phosphopeptide corresponding					

Background

Pyruvate kinase is a glycolytic enzyme that catalyses the conversion of phosphoenolpyruvate to pyruvate. In mammals, the M1 isoform (PKM1) is expressed in most adult tissues (1). The M2 isoform (PKM2) is an alternatively spliced variant of M1 that is expressed during embryonic development (1). Research studies found that cancer cells exclusively express PKM2 (1-3). PKM2 is shown to be essential for aerobic glycolysis in tumors, known as the Warburg effect (1). When cancer cells switch from the M2 isoform to the M1 isoform, aerobic glycolysis is reduced and oxidative phosphorylation is increased (1). These cells also show decreased tumorigenicity in mouse xenografts (1). Recent studies showed that PKM2 is not essential for all tumor cells (4). In the tumor model studied, PKM2 was found to be active in the non-proliferative tumor cell population and inactive in the proliferative tumor cell population (4).

to the sequence surrounding Tyr105 of human PKM2 protein. Antibodies are purified by protein A and

Additional studies show that the oncogenic forms of FGFR1 directly phosphorylate Tyr105 of PKM2 and thereby inhibit the formation of active, tetrameric PKM2 (5). A PKM2 missense mutation found in cancer cells results in the replacement of Tyr105 by phenylalanine and leads to reduced cell proliferation during hypoxia and tumor growth in nude mice xenografts (5). These findings suggest that the phosphorylation at Tyr105 is a critical switch for the metabolism in cancer cells that promotes tumor growth (5).

Phosphorylation of PKM2 on Tyr105 was identified at Cell Signaling Technology (CST) using PhosphoScan $^{\otimes}$, CST's LC-MS/MS platform for phosphorylation site discovery.

Background References

1. Christofk, H.R. et al. (2008) Nature 452, 230-3.

peptide affinity chromatography.

- 2. Mazurek, S. et al. (2005) Semin Cancer Biol 15, 300-8.
- 3. Dombrauckas, J.D. et al. (2005) Biochemistry 44, 9417-29.
- 4. Israelsen, W.J. et al. (2013) *Cell* 155, 397-409.
- 5. Hitosugi, T. et al. (2009) Sci Signal 2, ra73.

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

Western Blot BufferIMPORTANT: For western blots, incubate membrane with diluted primary antibody in 5% w/v BSA, 1X TBS, 0.1% Tween® 20 at 4°C with gentle shaking, overnight.

Applications Key WB: Western Blotting

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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Limited Uses

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