



PDHK1 (Phospho-Thr388) Antibody

#58026

Number: 58026

Amount: 100µg/100µl

Form of Antibody: Rabbit IgG in phosphate buffered saline (without Mg²⁺ and Ca²⁺), pH 7.4, 150mM NaCl, 0.02% sodium azide and 50% glycerol.

Storage/Stability: Store at -20°C/1 year

Immunogen: synthetic phosphopeptide corresponding to residues surrounding Thr388 of human PDHK1

Purification: The antibody was affinity-purified from rabbit antiserum by affinity-chromatography using epitope-specific phosphopeptide. The antibody against non-phosphopeptide was removed by chromatography using non-phosphopeptide corresponding to the phosphorylation site.

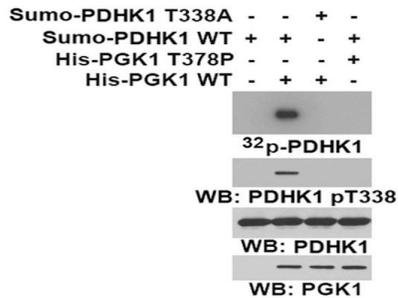
Specificity/Sensitivity: PDHK1 (Phospho-Thr388)antibody detects endogenous levels of PDHK1 only when phosphorylated at Threonine388 .

Reactivity: Human

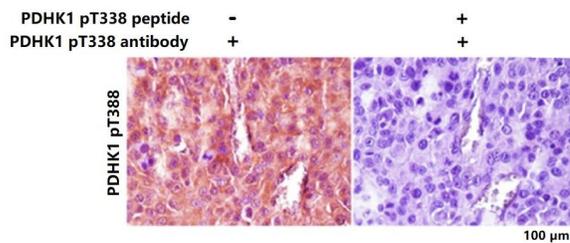
Applications:

Predicted MW: 43KD

WB :1:500~1:1000 IHC:1:50-200



In vitro phosphorylation analyses with autoradiography were performed by mixing purified WT PGK1 or PGK1 T378P with purified WT PDHK1 or PDHK1 T338A in the presence of [γ -32P] ATP. Immunoblotting analyses were performed with the indicated antibodies.



Validation of Antibody Specificities. IHC analyses of human GBM tissues were performed with the indicated antibodies in the presence or absence of specific blocking peptides. Scale bar, 100 µm.

Background :Mitochondrial PGK1 acts as a protein kinase to phosphorylate pyruvate dehydrogenase kinase 1 (PDHK1) at T338, which activates PDHK1 to phosphorylate and inhibit the pyruvate dehydrogenase (PDH) complex. This reduces mitochondrial pyruvate utilization, suppresses reactive oxygen species production, increases lactate production, and promotes brain tumorigenesis. PDHK1 T338 phosphorylation levels correlate with poor prognosis in glioblastoma patients[1] .

Reference:[1] Li X, Jiang Y, Meisenhelder J, Yang W, Hawke DH, Zheng Y, Xia Y, Aldape K, He J, Hunter T, Wang L, Lu Z. Mitochondria-Translocated PGK1 Functions as a Protein Kinase to Coordinate Glycolysis and the TCA Cycle in Tumorigenesis. *Mol Cell*. 2016 Mar 3;61(5):705-719. doi: 10.1016/j.molcel.2016.02.009. PMID: 26942675.