Store at -20°C

Gluconeogenesis Antibody Sampler Kit

#72624

Cell Signaling TECHNOLOGY®

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For Research Use Only. Not For Use In Diagnostic Procedures.

Product Includes	Product #	Quantity	Mol. Wt.	Isotype/Source
Enolase-1 Antibody	3810	20 µL	47 kDa	Rabbit
Enolase-2 (E2H9X) XP® Rabbit mAb	24330	20 µL	47 kDa	Rabbit IgG
FBP1/FBPase 1 (D2T7F) Rabbit mAb	59172	20 µL	39 kDa	Rabbit IgG
GPI (E2Q8J) XP® Rabbit mAb	94068	20 µL	60 kDa	Rabbit IgG
PCK1 (D12F5) Rabbit mAb	12940	20 µL	63 kDa	Rabbit IgG
PCK2 (D3E11) Rabbit mAb	8565	20 µL	71 kDa	Rabbit IgG
PGAM1 (D3J9T) Rabbit mAb	12098	20 µL	28 kDa	Rabbit IgG
PGK1 Antibody	68540	20 µL	43 kDa	Rabbit
Pyruvate Carboxylase Antibody	66470	20 µL	130 kDa	Rabbit
Anti-rabbit IgG, HRP-linked Antibody	7074	100 µL		Goat

See www.cellsignal.com for individual component applications, species cross-reactivity, dilutions, and additional application protocols.

Description: The Gluconeogenesis Antibody Sampler Kit provides an economical means of detecting select components involved in the gluconeogenesis metabolism pathway. The kit includes enough antibodies to perform two western blot experiments with each primary antibody.

Background: Enolase is an important glycolytic enzyme involved in the interconversion of 2-phosphoglycerate to phosphoenolpyruvate. Mammalian enolase exists as three subunits: enolase-1 (α-enolase), enolase-2 (γ-enolase), and enolase-3 (β-enolase). Expression of the enolase isoforms differs in a tissue specific manner (1). Enolase-1 plays a key role in anaerobic metabolism under hypoxic conditions and may act as a cell surface plasminogen receptor during tissue invasion (2,3). Abnormal expression of enolase-1 is associated with tumor progression in some cases of breast and lung cancer (4-7). Alternatively, an enolase-1 splice variant (MBP-1) binds the c-myc promoter p2 and may function as a tumor suppressor. For this reason, enolase-1 is considered as a potential therapeutic target in the treatment of some forms of cancer (8). Research studies have shown elevated levels of neuron-specific enolase-2 in neuroblastoma (1) and small-cell lung cancer (9,10). Fructose-1,6-bisphosphatase 1 (FBP1 or FBPase 1), a rate-limiting enzyme in gluconeogenesis, catalyzes the conversion of fructose-1,6-bisphosphate to fructose-6-phosphate (11). Inhibition of *FBP1* expression in basal-like breast cancer (BLBC) cells leads to metabolic reprogramming, including enhanced glycolysis, which leads to increased glucose uptake, biosynthesis of macromolecules, and activation of PKM2 (11). This metabolic reprogramming endows tumor cells with cancer stem cell (CSC)-like properties, thereby increasing their tumorigenicity (11). Depletion of FBP1 was also reported in more than 600 clear cell renal cell carcinoma (ccRCC) tumors, suggesting that FBP1 may inhibit ccRCC tumor progression (12). Glucose-6-phosphate isomerase (GPI) is a multi-functional protein belonging to the glucose phosphate isomerase family (13,14). As an intracellular metabolic enzyme, GPI plays a pivotal role in glycolysis and gluconeogenesis by catalyzing the interconversion of D-glucose-6-phosphate and D-fructose-6-phosphate (15). GPI is also secreted, where it functions as a cytokine (referred to as Autocrine Motility Factor, AMF), acting via the E3-ubiquitin-protein ligase AMFR/gp78 (16). In normal tissues, GPI/AMF has been shown to promote both immune cell maturation and neuronal cell survival (17,18). It is also secreted in abundance by some tumor cells (19), where it has been shown to promote tumor cell migration and metastasis (20,21). Phosphoenolpyruvate carboxykinase 1 (PCK1, PEPCK1, or PEPCK-C) is a cytosolic enzyme responsible for the conversion of oxaloacetate to phosphoenolpyruvate (22). PCK1 and PCK2 are involved in controlling the rate-limiting step of gluconeogenesis in the liver, which generates glucose from non-carbohydrate substrates, such as lactate and glycerol (23,24). PCK2 (PEPCK2 or PEPCK-M) encodes an isoform of phosphoenolpyruvate carboxykinase (PEPCK) that is found in the mitochondria of renal and hepatic tissues (22). Phosphoglycerate mutase (PGAM1) catalyzes the conversion of 3-phosphoglycerate to 2-phosphoglycerate during glycolysis (25-29). Research studies have shown increased PGAM1 expression in cancer (25-28) and mental disease (29). PGK1 (phosphoglycerate kinase) is an essential enzyme in the glycolysis pathway (30). It catalyzes the reversible phospho-transfer reaction from 1,3-diphosphoglycerate to ADP to form ATP and 3-phosphoglycerate. The expression of PGK1 is upregulated in many cancer types and plays an important role in cancer cell proliferation and metastasis (31-34). Pyruvate carboxylase (PC) catalyzes the carboxylation of pyruvate to oxaloacetate to replenish TCA cycle intermediates. It is also critical in regulating gluconeogenesis in the liver (35).

Storage: Supplied in 10 mM sodium HEPES (pH 7.5), 150 mM NaCl, 100 µg/ml BSA, 50% glycerol and less than 0.02% sodium azide. Store at –20°C. *Do not aliquot the antibodies.*

Please visit www.cellsignal.com for validation data and a complete listing of recommended companion products.

Background References:

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Applications: **W**—Western **IP**—Immunoprecipitation **IHC**—Immunohistochemistry **ChIP**—Chromatin Immunoprecipitation **IF**—Immunofluorescence **F**—Flow cytometry **E-P**—ELISA-Peptide **Species Cross-Reactivity:** **H**—human **M**—mouse **R**—rat **Hm**—hamster **Mk**—monkey **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 **All**—all species expected Species enclosed in parentheses are predicted to react based on 100% homology.

Specificity/Sensitivity: Each antibody in the Gluconeogenesis Antibody Sampler Kit detects endogenous levels of its target protein. Enolase-1 Antibody does not cross-react with enolase-2. Enolase-2 (E2H9X) XP® Rabbit mAb does not cross-react with human enolase-1 protein. PGAM1 (D3J9T) Rabbit mAb may cross-react with overexpressed PGAM2 protein.

Source/Purification: Monoclonal antibodies are produced by immunizing animals with synthetic peptides corresponding to residues near the carboxy terminus of human enolase-2 protein, human PGAM1 protein, and human PCK1 protein, near the amino terminus of human PCK2 protein, and surrounding Val106 of human FBP1/FBPase 1 protein and Pro8 of human GPI protein.

Polyclonal antibodies are produced by immunizing animals with synthetic peptides corresponding to residues surrounding Thr298 of human PGK1 protein, near the carboxy terminus of human pyruvate carboxylase protein, and corresponding to residues surrounding Ala76 of human enolase-1 protein. Antibodies are purified by protein A and peptide affinity chromatography. Pyruvate Carboxylase Antibody is purified by peptide affinity chromatography.