## PKM2 (D78A4) XP<sup>®</sup> Rabbit mAb (PE Conjugate)



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

<b>Applications:</b> FC-FP	<b>Reactivity:</b> All	<b>Sensitivity:</b> Transfected Only	<b>Source/Isotype:</b> Rabbit IgG	UniProt ID: #P14618	Entrez-Gene Id: 5315
Product Usage Information		Application Flow Cytometry (Fixed/Permeabilized)			<b>Dilution</b> 1:50
Storage		Supplied in PBS (pH 7.2), less than 0.1% sodium azide and 2 mg/ml BSA. Store at $4^{\circ}$ C. Do not aliquot the antibody. Protect from light. Do not freeze.			
Specificity/Sensitivity		PKM2 (D78A4) $XP^{\otimes}$ Rabbit mAb detects endogenous levels of total PKM2 protein and does not cross-react with PKM1.			
Source / Purification		Monoclonal antibody is produced by immunizing animals with a synthetic peptide corresponding to residues surrounding Ser406 of human PKM2 protein.			
Description		This Cell Signaling Technology antibody is conjugated to phycoerythrin (PE) and tested in-house for direct flow cytometric analysis in human cells. This antibody is expected to exhibit the same species cross-reactivity as the unconjugated PKM2 (D78A4) XP <sup>®</sup> Rabbit mAb #4053.			
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).			
Background References		<ol> <li>Christofk, H.R. et al. (2008) Nature 452, 230-3.</li> <li>Mazurek, S. et al. (2005) Semin Cancer Biol 15, 300-8.</li> <li>Dombrauckas, J.D. et al. (2005) Biochemistry 44, 9417-29.</li> <li>Israelsen, W.J. et al. (2013) Cell 155, 397-409.</li> </ol>			

**Species Reactivity** 

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

**Applications Key** 

FC-FP: Flow Cytometry (Fixed/Permeabilized)

**Cross-Reactivity Key** 

**All:** All Species Expected

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