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## TET2 (D6C7K) Rabbit mAb (PE Conjugate)

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

<b>Applications:</b> FC-FP	<b>Reactivity:</b> M	<b>Sensitivity:</b> Endogenous	<b>Source/Isotype:</b> Rabbit IgG	<b>UniProt ID:</b> #Q4JK59	<b>Entrez-Gene Id:</b> 214133
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Product Usage Information	Application	Dilution
<b>Storage</b>	Flow Cytometry (Fixed/Permeabilized)	1:50
<b>Specificity/Sensitivity</b>	Supplied in PBS (pH 7.2), less than 0.1% sodium azide and 2 mg/ml BSA. Store at 4°C. Do not aliquot the antibody. Protect from light. Do not freeze.	
<b>Source / Purification</b>	TET2 (D6C7K) Rabbit mAb (PE Conjugate) recognizes endogenous levels of total TET2 protein.	
<b>Description</b>	Monoclonal antibody is produced by immunizing animals with a synthetic peptide corresponding to residues surrounding Val1640 of Mouse TET2 protein.	
<b>Background</b>	This Cell Signaling Technology antibody is conjugated to phycoerythrin (PE) and tested in-house for direct flow cytometric analysis in mouse cells. This antibody is expected to exhibit the same species cross-reactivity as the unconjugated TET2 (D6C7K) Rabbit mAb #36449.	
<b>Background References</b>	Methylation of DNA at cytosine residues is a heritable, epigenetic modification that is critical for proper regulation of gene expression, genomic imprinting, and mammalian development (1,2). 5-methylcytosine is a repressive epigenetic mark established de novo by two enzymes, DNMT3a and DNMT3b, and is maintained by DNMT1 (3, 4). 5-methylcytosine was originally thought to be passively depleted during DNA replication. However, subsequent studies have shown that Ten-Eleven Translocation (TET) proteins TET1, TET2, and TET3 can catalyze the oxidation of methylated cytosine to 5-hydroxymethylcytosine (5-hmC) (5). Additionally, TET proteins can further oxidize 5-hmC to form 5-formylcytosine (5-fC) and 5-carboxylcytosine (5-caC), both of which are excised by thymine-DNA glycosylase (TDG), effectively linking cytosine oxidation to the base excision repair pathway and supporting active cytosine demethylation (6,7). TET2 is the most frequently mutated gene in myeloid dysplastic syndrome (MDS), a dysplasia of myeloid, megakaryocytic, and/or erythroid cell lineages, of which 30% progress to acute myeloid leukemia (AML) (8, 9). It is also mutated in diffuse large B-cell lymphoma (10). TET2 protein expression is often reduced in solid tumors such as prostate cancer, melanoma, and oral squamous cell carcinoma (11-13).	
	<ol style="list-style-type: none"> <li>Hermann, A. et al. (2004) <i>Cell Mol Life Sci</i> 61, 2571-87.</li> <li>Turek-Plewa, J. and Jagodziński, P.P. (2005) <i>Cell Mol Biol Lett</i> 10, 631-47.</li> <li>Okano, M. et al. (1999) <i>Cell</i> 99, 247-57.</li> <li>Li, E. et al. (1992) <i>Cell</i> 69, 915-26.</li> <li>Tahiliani, M. et al. (2009) <i>Science</i> 324, 930-5.</li> <li>He, Y.F. et al. (2011) <i>Science</i> 333, 1303-7.</li> <li>Ito, S. et al. (2011) <i>Science</i> 333, 1300-3.</li> <li>Langemeijer, S.M. et al. (2009) <i>Nat Genet</i> 41, 838-42.</li> <li>Yamazaki, J. et al. (2012) <i>Epigenetics</i> 7, 201-7.</li> <li>Asmar, F. et al. (2013) <i>Haematologica</i> 98, 1912-20.</li> <li>Nickerson, M.L. et al. (2013) <i>Hum Mutat</i> 34, 1231-41.</li> <li>Lian, C.G. et al. (2012) <i>Cell</i> 150, 1135-46.</li> <li>Jäwert, F. et al. (2013) <i>Anticancer Res</i> 33, 4325-8.</li> </ol>	

<b>Species Reactivity</b>	Species reactivity is determined by testing in at least one approved application (e.g., western blot).
<b>Applications Key</b>	<b>FC-FP:</b> Flow Cytometry (Fixed/Permeabilized)
<b>Cross-Reactivity Key</b>	<b>M:</b> Mouse
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