

TET2 (D6B9Y) Rabbit mAb



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		280	Source/Isotype: Rabbit IgG	UniProt ID: #Q6N021	Entrez-Gene Id: 54790
	For optimal ChIP results, use 20 μl of antibody and 10 μg of chromatin (approximately 4 x 10 ⁶ cells) per IP. This antibody has been validated using SimpleChIP [®] Enzymatic Chromatin IP Kits.				
	Application Dilution				
	Western Blotting			1:1000	
	• •			1:100	
	Chromatin IP			1:25	
	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 antibody.				rol and less than
tivity	TET2 (D6B9Y) Rabbit mAb recognizes endogenous levels of total TET2 protein.				
tion	Monoclonal antibody is produced by immunizing animals with recombinant protein specific to the carboxy terminus of human TET2 protein.				
	regulation of gene ex methylcytosine is a re DNMT3b, and is main depleted during DNA Translocation (TET) pr 5-hydroxymethylcytos formylcytosine (5-fC) a glycosylase (TDG), effo supporting active cyto dysplastic syndrome (which 30% progress to lymphoma (10). TET2	pression, genomic i pressive epigenetic tained by DNMT1 (3 replication. Howeve oteins TET1, TET2, a sine (5-hmC) (5). Add and 5-carboxylcytos ectively linking cyto beine demethylation MDS), a dysplasia o acute myeloid leu protein expression	mprinting, and mamma mark established de no 8, 4). 5-methylcytosine wer, subsequent studies had TET3 can catalyze the ditionally, TET proteins cine (5-caC), both of whice sine oxidation to the base (6,7). TET2 is the most of the femile (AML) (8, 9). It is a sis often reduced in solid	lian development (' vo by two enzymes as originally thougl ave shown that Ten e oxidation of meth an further oxidize 5 th are excised by the se excision repair po frequently mutated tic, and/or erythroitalso mutated in diffi	I,2). 5- , DNMT3a and nt to be passively -Eleven ylated cytosine to -hmC to form 5- ymine-DNA athway and gene in myeloid d cell lineages, of use large B-cell
erences	2. Turek-Plewa, J. and 3. Okano, M. et al. (19 4. Li, E. et al. (1992) <i>Ce</i> 5. Tahiliani, M. et al. (2 6. He, Y.F. et al. (2011) 7. Ito, S. et al. (2011) <i>S</i>	and Jagodziński, P.P. (2005) <i>Cell Mol Biol Lett</i> 10, 631-47. l. (1999) <i>Cell</i> 99, 247-57. 2) <i>Cell</i> 69, 915-26. al. (2009) <i>Science</i> 324, 930-5. 011) <i>Science</i> 333, 1303-7.			
	tivity tion erences	Western Blotting Immunoprecipitation Chromatin IP Supplied in 10 mM so 0.02% sodium azide. S tivity TET2 (D6B9Y) Rabbit r Monoclonal antibody carboxy terminus of h Methylation of DNA ar regulation of gene expended during DNA Translocation (TET) pr 5-hydroxymethylcytosine (5-fC) aglycosylase (TDG), effecting supporting active cytodysplastic syndrome (which 30% progress to lymphoma (10). TET2 melanoma, and oral s Perences 1. Hermann, A. et al. (194. Li, E. et al. (1992) C65. Tahiliani, M. et al. (2011)	Western Blotting Immunoprecipitation Chromatin IP Supplied in 10 mM sodium HEPES (pH 7.5 0.02% sodium azide. Store at -20°C. Do n tivity TET2 (D6B9Y) Rabbit mAb recognizes end Monoclonal antibody is produced by imm carboxy terminus of human TET2 protein Methylation of DNA at cytosine residues i regulation of gene expression, genomic i methylcytosine is a repressive epigenetic DNMT3b, and is maintained by DNMT1(3 depleted during DNA replication. Howeve Translocation (TET) proteins TET1, TET2, a 5-hydroxymethylcytosine (5-hmC) (5). Add formylcytosine (5-fC) and 5-carboxylcytos glycosylase (TDG), effectively linking cyto supporting active cytosine demethylation dysplastic syndrome (MDS), a dysplasia o which 30% progress to acute myeloid leu lymphoma (10). TET2 protein expression melanoma, and oral squamous cell carcin 1. Hermann, A. et al. (2004) Cell Mol Life 9 2. Turek-Plewa, J. and Jagodziński, P.P. (20 3. Okano, M. et al. (1999) Cell 99, 247-57. 4. Li, E. et al. (1992) Cell 69, 915-26. 5. Tahiliani, M. et al. (2009) Science 324, 9 6. He, Y.F. et al. (2011) Science 333, 1303-	Western Blotting Immunoprecipitation Chromatin IP Supplied in 10 mM sodium HEPES (pH 7.5), 150 mM NaCl, 100 µg, 0.02% sodium azide. Store at -20°C. Do not aliquot the antibody. TET2 (D6B9Y) Rabbit mAb recognizes endogenous levels of total of the carboxy terminus of human TET2 protein. Methylation of DNA at cytosine residues is a heritable, epigenetic regulation of gene expression, genomic imprinting, and mamma methylcytosine is a repressive epigenetic mark established de no DNMT3b, and is maintained by DNMT1 (3, 4). 5-methylcytosine we depleted during DNA replication. However, subsequent studies have a translocation (TET) proteins TET1, TET2, and TET3 can catalyze the 5-hydroxymethylcytosine (5-hmC) (5). Additionally, TET proteins c formylcytosine (5-fC) and 5-carboxylcytosine (5-caC), both of whice glycosylase (TDG), effectively linking cytosine oxidation to the bas supporting active cytosine demethylation (6,7). TET2 is the most of dysplastic syndrome (MDS), a dysplasia of myeloid, megakaryocy which 30% progress to acute myeloid leukemia (AML) (8, 9). It is a lymphoma (10). TET2 protein expression is often reduced in solid melanoma, and oral squamous cell carcinoma (11-13). Perences 1. Hermann, A. et al. (2004) Cell Mol Life Sci 61, 2571-87. 2. Turek-Plewa, J. and Jagodziński, P.P. (2005) Cell Mol Biol Lett 10, 3. Okano, M. et al. (1999) Cell 99, 247-57.	Western Blotting 1:1000 Immunoprecipitation 1:100 Chromatin IP 1:25 Supplied in 10 mM sodium HEPES (pH 7.5), 150 mM NaCl, 100 µg/ml BSA, 50% glycer 0.02% sodium azide. Store at -20°C. Do not aliquot the antibody. TET2 (D6B9Y) Rabbit mAb recognizes endogenous levels of total TET2 protein. Monoclonal antibody is produced by immunizing animals with recombinant protein carboxy terminus of human TET2 protein. Methylation of DNA at cytosine residues is a heritable, epigenetic modification that i regulation of gene expression, genomic imprinting, and mammalian development (*methylcytosine is a repressive epigenetic mark established de novo by two enzymes DNMT3b, and is maintained by DNMT1 (3, 4). 5-methylcytosine was originally though depleted during DNA replication. However, subsequent studies have shown that Ten Translocation (TET) proteins TET1, TET2, and TET3 can catalyze the oxidation of meth 5-hydroxymethylcytosine (5-hmC) (5). Additionally, TET proteins can further oxidize 5 formylcytosine (5-fC) and 5-carboxylcytosine (5-dC), both of which are excised by thy glycosylase (TDG), effectively linking cytosine oxidation to the base excision repair psupporting active cytosine demethylation (6,7). TET2 is the most frequently mutated dysplastic syndrome (MDS), a dysplasia of myeloid, megakaryocytic, and/or erythroid which 30% progress to acute myeloid leukemia (AML) (8, 9). It is also mutated in difful lymphoma (10). TET2 protein expression is often reduced in solid tumors such as promelanoma, and oral squamous cell carcinoma (11-13). Perences 1. Hermann, A. et al. (2004) Cell Mol Life Sci 61, 2571-87. 2. Turek-Plewa, J. and Jagodziński, P.P. (2005) Cell Mol Biol Lett 10, 631-47. 3. Okano, M. et al. (1999) Cell 99, 247-57. 4. Li, E. et al. (1992) Cell 69, 915-26. 5. Tahiliani, M. et al. (2001) Science 333, 1303-7.

Species Reactivity

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

Western Blot Buffer

IMPORTANT: 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

W: Western Blotting IP: Immunoprecipitation ChIP: Chromatin IP

Cross-Reactivity Key

H: Human

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