

## **HTATIP2/TIP30 Antibody**



Orders: 877-616-CELL (2355)

orders@cellsignal.com

Support: 877-678-TECH (8324)

Web: info@cellsignal.com

cellsignal.com

3 Trask Lane | Danvers | Massachusetts | 01923 | USA

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

Applications: W	Reactivity: H	<b>Sensitivity:</b> Endogenous	<b>MW (kDa):</b> 28, 32	<b>Source/Isotype:</b> Rabbit	UniProt ID: #Q9BUP3	Entrez-Gene Id: 10553
Product Usage Information		<b>Application</b> Western Blotting			<b>Dilution</b> 1:1000	
Storage		Supplied in 10 mM sodium HEPES (pH 7.5), 150 mM NaCl, 100 μg/ml BSA and 50% glycerol. Store at – 20°C. Do not aliquot the antibody.				
Specificity/Sensitivity		HTATIP2/TIP30 Antibody recognizes endogenous levels of total HTATIP2 protein.				
Source / Purification		Polyclonal antibodies are produced by immunizing animals with a synthetic peptide corresponding to residues surrounding His231 of human HTATIP2 protein. Antibodies are purified by protein A and peptide affinity chromatography.				
Background		The HIV-1 Tat interactive protein 2 (HTATIP2, TIP30, CC3) is an oxidoreductase that was originally identified as a metastatic tumor suppressor and Tat-mediated proapoptotic gene transcription cofactor (1,2). HTATIP2 protein contains a short-chain dehydrogenase (SDR) domain and a NADPH binding motif important for HTATIP2 interaction with importins and inhibition of nucleocytoplasmic transport (3,4). Research studies demonstrate that induced overexpression of HTATIP2 predisposes cells to apoptosis by inhibiting the nuclear transport of important signaling proteins (e.g. p53, activated notch1) and several key targets of the DNA repair process (5-7). HTATIP2 is part of a protein complex, with Rab5a, endophilin B1, and ACSL4, that may regulate EGFR receptor endosomal trafficking, degradation, and cytoplasmic/nuclear signaling (8,9). Silencing of HTATIP2 promotes tumor cell survival under low glucose conditions by inducing increased expression of mitochondrial respiratory proteins and glucose metabolic enzymes (10).				
Background References		1. Shtivelman, E. (1997) <i>Oncogene</i> 14, 2167-73. 2. Xiao, H. et al. (2000) <i>EMBO J</i> 19, 956-63. 3. El Omari, K. et al. (2005) <i>J Biol Chem</i> 280, 18229-36. 4. King, F.W. and Shtivelman, E. (2004) <i>Mol Cell Biol</i> 24, 7091-101. 5. Zhao, J. et al. (2008) <i>Cancer Res</i> 68, 4133-41. 6. Nakahara, J. et al. (2009) <i>J Clin Invest</i> 119, 169-81. 7. Fong, S. et al. (2010) <i>BMC Cell Biol</i> 11, 23. 8. Zhang, C. et al. (2011) <i>J Biol Chem</i> 286, 9373-81. 9. Li, A. et al. (2013) <i>Oncogene</i> 32, 2273-81, 2281e.1-12. 10. Chen, V. and Shtivelman, E. (2010) <i>Cell Cycle</i> 9, 4941-53.				
Species Reacti	vity	Species reactivity is d	etermined by testin	g in at least one approve	ed 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

**Cross-Reactivity Key** H: Human

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