

## **CUTL1 Antibody**

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**Limited Uses** 



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

<b>Applications:</b> W, IP	Reactivity: H	<b>Sensitivity:</b> Endogenous	<b>MW (kDa):</b> 200	<b>Source/Isotype:</b> Rabbit	UniProt ID: #P39880	Entrez-Gene Id 1523
Product Usage Information		ApplicationDilutionWestern Blotting1:1000Immunoprecipitation1:200				
Storage		Supplied in 10 mM sodium HEPES (pH 7.5), 150 mM NaCl, 100 $\mu$ g/ml BSA and 50% glycerol. Store at – 20°C. Do not aliquot the antibody.				
Specificity/Sensitivity		CUTL1 Antibody recognizes endogenous levels of total CUTL1 protein. It recognizes the p200 isoform (the full-length) and is not expected to recognize the p110 and p75 isoforms.				
Source / Purification		Polyclonal antibodies are produced by immunizing animals with a synthetic peptide corresponding to residues surrounding Gln639 of human CUTL1 protein. Antibodies are purified by protein A and peptide affinity chromatography.				
Background		CUTL1 (Cut-like 1), also known as CUX1 (Cut homeobox 1) (CUX1), is a transcription factor that has been implicated in many cellular processes in different tissues, such as cell migration, neuronal differentiation, and DNA repair (1-5). CUTL1 expression and activities are altered in cancer. Research studies have shown the CUTL1 gene to be a frequent target of loss-of-heterozygocity in various cancers (6,7). On the other hand, CUTL1 expression is elevated in many cancers and is associated with shorter disease-free survival (8). These accumulating evidence suggest that decreased CUTL1 expression promote tumor initiation and increased CUTL1 expression facilitates tumor progression (9). While full-length CUTL1 is about 200 kDa (p200), short forms p110 and p75 can also be generated by proteolytic processing and alternative transcription initiation site, respectively (10, 11).				
Background References		<ol> <li>Rodríguez-Tornos, F.M. et al. (2016) Neuron 89, 494-506.</li> <li>Ramdzan, Z.M. et al. (2015) Oncotarget 6, 3613-26.</li> <li>Ramdzan, Z.M. et al. (2014) PLoS Biol 12, e1001807.</li> <li>Kedinger, V. et al. (2009) J Biol Chem 284, 27701-11.</li> <li>Vadnais, C. et al. (2012) Nucleic Acids Res 40, 4483-95.</li> <li>McNerney, M.E. et al. (2013) Blood 121, 975-83.</li> <li>Ramdzan, Z.M. and Nepveu, A. (2014) Nat Rev Cancer 14, 673-82.</li> <li>Michl, P. et al. (2005) Cancer Cell 7, 521-32.</li> <li>Hulea, L. and Nepveu, A. (2012) Gene 497, 18-26.</li> <li>Moon, N.S. et al. (2001) Mol Cell Biol 21, 6332-45.</li> <li>Goulet, B. et al. (2002) Cancer Res 62, 6625-33.</li> </ol>				
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 nonfat dry milk, 1X TBS, 0.1% Tween® 20 at 4°C with gentle shaking, overnight.				
Applications Key		W: Western Blotting IP: Immunoprecipitation				
Cross-Reactivity Key		H: Human				
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