

## TACE (D22H4) Rabbit mAb



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

Applications: W	Reactivity: H	<b>Sensitivity:</b> Endogenous	<b>MW (kDa):</b> 135	<b>Source/Isotype:</b> Rabbit IgG	UniProt ID: #P78536	Entrez-Gene Id: 6868
Product Usage Information	2	<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, 50% glycerol and less than 0.02% sodium azide. Store at –20°C. Do not aliquot the antibody.				
Specificity/Sensitivity		TACE (D22H4) Rabbit mAb recognizes endogenous levels of total TACE protein.				
Source / Purification		Monoclonal antibody is produced by immunizing animals with recombinant protein specific to the amino terminus of human TACE protein.				
Background		TACE (TNF-α converting enzyme), also known as ADAM17, is a transmembrane metalloprotease that plays a key role in the cleavage of a number of cell surface molecules in a process known as "shedding". TACE is abundantly expressed in many adult tissues, but in fetal development, expression is differentially regulated (1). An important substrate of TACE is pro-TNF-α (1). Increased expression of TACE is associated with several pathological conditions, including osteoarthritis and rheumatoid arthritis, where the pro-inflammatory effects of increased TNF-α contribute to disease pathogenesis (2,3). Regulation of other important molecules by TACE, such as EGFR and Notch, has recently been documented. TACE is responsible for the shedding of EGFR ligands such as amphiregulin and TNF-α. Some tumors have hyperactivated EGFR due to upregulated TNF-α production and upregulated TACE, making TACE a potential target for drug development (4). TACE activates Notch in a ligand-independent manner and has been shown to play a role in the development of the <i>Drosophila</i> nervous system (5). TACE has also been proposed to act as an α-secretase for amyloid precursor protein (APP) (6), and to be involved in the renewal and proliferation of neural stem cells (7).				
Background References		<ol> <li>Black, R.A. et al. (1997) Nature 385, 729-33.</li> <li>Amin, A.R. (1999) Osteoarthritis Cartilage 7, 392-4.</li> <li>Patel, I.R. et al. (1998) J Immunol 160, 4570-9.</li> <li>Kenny, P.A. (2007) Differentiation 75, 800-8.</li> <li>Delwig, A. and Rand, M.D. (2008) Cell Mol Life Sci 65, 2232-43.</li> <li>Deuss, M. et al. (2008) Curr Alzheimer Res 5, 187-201.</li> <li>Rubio-Araiz, A. et al. (2008) Mol Cell Neurosci 38, 374-80.</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 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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