

## OX40 (ACT35) Mouse mAb



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

Applications: W, IP, IHC-Bond, HC-P, IF-IC, FC-FP, FC-L	Reactivity: H	<b>Sensitivity:</b> Endogenous	<b>MW (kDa):</b> 35-50	<b>Source/Isotype:</b> Mouse IgG2a	<b>UniProt ID:</b> #P43489	Entrez-Gene Id 7293
Product Usage Information		Application				Dilution
Imormation		Western Blotting				1:1000
		Immunoprecipitation IHC Leica Bond				1:50 1:400
		Immunohistochemist	ry (Paraffin)			1:400
		Immunofluorescence		uistry)		1:1000
		Flow Cytometry (Fixed		113ti y)		1:200
		Flow Cytometry (Live)	in crifficabilized)			1:200
Storage		Supplied in 10 mM sodium HEPES (pH 7.5), 150 mM NaCl, 100 $\mu$ g/ml BSA, 50% glycerol and less than 0.02% sodium azide. Store at –20°C. Do not aliquot the antibody.				
		For a carrier free (BSA and azide free) version of this product see product #99419.				
Specificity/Sensitivity		OX40 (ACT35) Mouse mAb recognizes endogenous levels of total OX40 protein.				
Source / Purification		Monoclonal antibody is produced by immunizing animals with HuT 102 cell line.				
Background		OX40 (TNFRSF4, CD134) is a member of the tumor necrosis factor (TNF) receptor superfamily that regulates T cell activity and immune responses. The OX40 protein contains four cysteine rich domains, a transmembrane domain, and a cytoplasmic tail containing a QEE motif (1,2). OX40 is primarily expressed on activated CD4+ and CD8+ T-cells, while the OX40 ligand (OX40L, TNFSF4, CD252) is predominantly expressed on activated antigen presenting cells (3-7). The engagement of OX40 with OX40L leads to the recruitment of TNF receptor-associated factors (TRAFs) and results in the formation of a TCR-independent signaling complex. One component of this complex, PKC0, activates the NF-kB pathway (2,8). OX40 signaling through Akt can also enhance TCR signaling directly (9). Research studies indicate that the OX40L-OX40 pathway is associated with inflammation and autoimmune diseases (10). Additional research studies show that OX40 agonists augment anti-tumor immunity in several cancer types (11,12).				
Background Re	1. Croft, M. (2010) Annu Rev Immunol 28, 57-78. 2. So, T. and Croft, M. (2012) Front Immunol 3, 133. 3. Paterson, D.J. et al. (1987) Mol Immunol 24, 1281-90. 4. Mallett, S. et al. (1990) EMBO J 9, 1063-8. 5. Dürkop, H. et al. (1995) Br J Haematol 91, 927-31. 6. Godfrey, W.R. et al. (1994) J Exp Med 180, 757-62. 7. Al-Shamkhani, A. et al. (1997) J Biol Chem 272, 5275-82. 8. So, T. et al. (2011) Proc Natl Acad Sci U S A 108, 2903-8. 9. So, T. and Croft, M. (2013) Front Immunol 4, 139. 10. Gough, M.J. and Weinberg, A.D. (2009) Adv Exp Med Biol 647, 94-107. 11. Weinberg, A.D. et al. (2011) Immunol Rev 244, 218-31. 12. Linch, S.N. et al. (2015) Front Oncol 5, 34.					

**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  $4^{\circ}$ C with gentle shaking, overnight.

**Applications Key** 

**W**: Western Blotting **IP**: Immunoprecipitation **IHC-Bond**: IHC Leica Bond **IHC-P**: Immunohistochemistry (Paraffin) **IF-IC**: Immunofluorescence (Immunocytochemistry) **FC-FP**: Flow Cytometry

(Fixed/Permeabilized) FC-L: Flow Cytometry (Live)

Cross-Reactivity Key H: Human

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