

DESCRIPTION

Species Reactivity	Mouse
Specificity	Detects mouse CD30/TNFRSF8 in ELISAs and Western blots. In sandwich immunoassays, no cross-reactivity or interference with recombinant mouse (rm) TNF RI/TNFRSF1A, recombinant human (rh) TNF RI/TNFRSF1A, rhTNF RII/TNFRSF1B, rmTNF- α /TNFSF2, rhTNF- α /TNFSF2, rhTNF- β /TNFSF1, rhCD30/TNFRSF8, or rmCD30 Ligand/TNFSF8 is observed. In Western blots, no cross-reactivity with rhNGFR/TNFRSF16, rmOPG/TNFRSF11B, rmRANK/TNFRSF11A, rmTNF RI/TNFRSF1A, rmTNF RII/TNFRSF1B, rmEDAR, rmTAJ/TNFSF19, rmLTR β /TNFRSF3, rm4-1BB/TNFRSF9, rmCD27/TNFRSF7, rhCD30/TNFRSF8, rmCD40/TNFRSF5, rhDR3/TNFRSF12, rhDR6/TNFRSF21, rmFas/TNFRSF6, rmGITR/TNFRSF18, or rhHVEM/TNFRSF14 is observed.
Source	Monoclonal Rat IgG _{2A} Clone # 115705
Purification	Protein A or G purified from hybridoma culture supernatant
Immunogen	Mouse myeloma cell line NS0-derived recombinant mouse CD30/TNFRSF8 Phe19-Thr281 Accession # Q60846
Formulation	Lyophilized from a 0.2 μ m filtered solution in PBS with Trehalose. See Certificate of Analysis for details. *Small pack size (-SP) is supplied either lyophilized or as a 0.2 μ m filtered solution in PBS.

APPLICATIONS

Please Note: Optimal dilutions should be determined by each laboratory for each application. *General Protocols* are available in the *Technical Information* section on our website.

	Recommended Concentration	Sample
Western Blot	1 μ g/mL	Recombinant Mouse CD30/TNFRSF8 Fc Chimera (Catalog # 852-CD)
Mouse CD30/TNFRSF8 Sandwich Immunoassay		Reagent
ELISA Capture	2-8 μ g/mL	Mouse CD30/TNFRSF8 Antibody (Catalog # MAB8521)
ELISA Detection	0.1-0.4 μ g/mL	Mouse CD30/TNFRSF8 Biotinylated Antibody (Catalog # BAF852)
Standard		Recombinant Mouse CD30/TNFRSF8 Fc Chimera (Catalog # 852-CD)

PREPARATION AND STORAGE

Reconstitution	Reconstitute at 0.5 mg/mL in sterile PBS.
Shipping	The product is shipped at ambient temperature. Upon receipt, store it immediately at the temperature recommended below. *Small pack size (-SP) is shipped with polar packs. Upon receipt, store it immediately at -20 to -70 °C
Stability & Storage	Use a manual defrost freezer and avoid repeated freeze-thaw cycles. <ul style="list-style-type: none"> • 12 months from date of receipt, -20 to -70 °C as supplied. • 1 month, 2 to 8 °C under sterile conditions after reconstitution. • 6 months, -20 to -70 °C under sterile conditions after reconstitution.

BACKGROUND

CD30, also known as Ki-1 antigen and TNFRSF8, is a 120 kDa type I transmembrane glycoprotein belonging to the TNF receptor superfamily (1, 2). Mature mouse CD30 consists of a 264 amino acid (aa) extracellular domain (ECD) with three cysteine-rich repeats, a 27 aa transmembrane segment, and a 190 aa cytoplasmic domain (3). In contrast, human CD30 includes an additional 90 aa in the ECD and contains six cysteine-rich repeats. Within common regions of the ECD, mouse CD30 shares 53% and 80% aa sequence identity with human and rat CD30, respectively. CD30 is normally expressed on antigen-stimulated Th cells and B cells (4-6). However, it is upregulated in Hodgkin's disease (on Reed-Sternberg cells), other lymphomas, chronic inflammation, and autoimmunity (7). CD30 binds to CD30 Ligand/TNFSF8 which is expressed on activated Th cells, monocytes, granulocytes and medullary thymic epithelial cells (1, 5). CD30 signaling costimulates antigen-induced Th0 and Th2 proliferation and cytokine secretion but favors a Th2-biased immune response (8). In the absence of antigenic stimulation, it can still induce T cell expression of IL-13 (9). CD30 contributes to thymic negative selection by inducing the apoptotic cell death of CD4⁺CD8⁺ T cells (10, 11). In B cells, CD30 ligation promotes cellular proliferation and antibody production in addition to the expression of CXCR4, CCL3, and CCL5 (5, 12). An 85-90 kDa soluble form of CD30 is shed from the cell surface by TACE-mediated cleavage (13, 14). Soluble CD30 retains the ability to bind CD30 Ligand and functions as an inhibitor of normal CD30 signaling (15).

References:

1. Kennedy, M.K. *et al.* (2006) *Immunology* **118**:143.
2. Tarkowski, M. (2003) *Curr. Opin. Hematol.* **10**:267.
3. Bowen, M.A. *et al.* (1996) *J. Immunol.* **156**:442.
4. Hamann, D. *et al.* (1996) *J. Immunol.* **156**:1387.
5. Shanebeck, S.D. *et al.* (1995) *Eur. J. Immunol.* **25**:2147.
6. Gruss, H.-J. *et al.* (1994) *Blood* **83**:2045.
7. Ofizoglu E. *et al.* (2009) *Adv. Exp. Med. Biol.* **647**:174.
8. Del Prete, G. *et al.* (1995) *J. Exp. Med.* **182**:1655.
9. Harlin, H. *et al.* (2002) *J. Immunol.* **169**:2451.
10. Amakawa, R. *et al.* (1996) *Cell* **84**:551.
11. Chiarle, R. *et al.* (1999) *J. Immunol.* **163**:194.
12. Vinante, F. *et al.* (2002) *Blood* **99**:52.
13. Hansen, H.P. *et al.* (1995) *Int. J. Cancer* **63**:750.
14. Hansen, H.P. *et al.* (2000) *J. Immunol.* **165**:6703.
15. Hargreaves, P.G. and A. Al-Shamkhani (2002) *Eur. J. Immunol.* **32**:163.