

DESCRIPTION

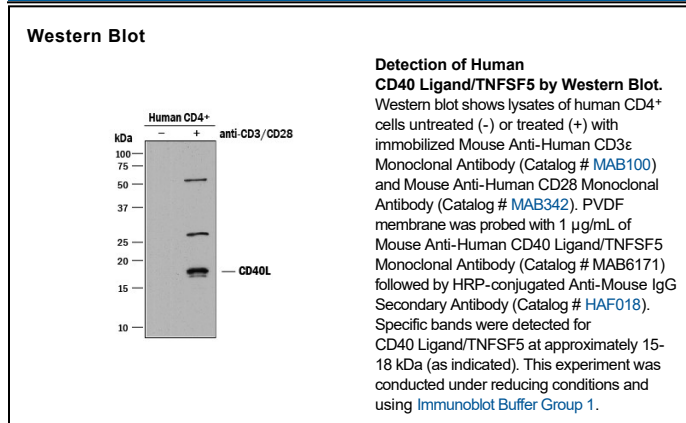
Species Reactivity	Human
Specificity	Detects human CD40 Ligand in direct ELISAs.
Source	Monoclonal Mouse IgG _{2B} Clone # 938601
Purification	Protein A or G purified from hybridoma culture supernatant
Immunogen	<i>E. coli</i> -derived recombinant human CD40 Ligand Glu108-Leu261 Accession # P29965
Formulation	Lyophilized from a 0.2 µm 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 µm 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 µg/mL	See Below

DATA



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

CD40 Ligand (CD40L), now renamed TNFSF5, but also known as CD154, TRAP and gp39, is a 34-39 kDa type II transmembrane glycoprotein that belongs to the TNF superfamily (1-3). Human CD40L is 261 amino acids (aa) in length and consists of a 22 aa cytoplasmic domain, a 24 aa transmembrane segment, and a 215 aa extracellular region that consists of multiple β -strands and one N-linked glycosylation site (4, 5). Although carbohydrates are present, they are not necessary for activity (6). As with other TNF superfamily members, CD40L will exist as a trimer, both as a membrane bound and soluble form (6-8). The soluble form is 18 kDa in size and about 150 aa in length, and arises from intracellular proteolytic processing. As a trimer, the soluble form is bioactive (7-9). Multiple mutations and alternate splice forms of CD40L exist, often associated with pathology and leading to truncated or nontrimerizable forms of CD40L (8). While CD40L is a ligand for CD40, the ligation of CD40L by CD40 initiates bidirectional signaling in both CD40 and CD40L expressing cells (10). The extracellular region of human CD40L is 99%, 88%, 86%, 82%, 75% and 75% aa identical to the extracellular regions of CD40L in rhesus monkey, bovine, porcine, canine, mouse and rat, respectively. CD40L binds to both CD40 and to integrin α IIb β 3 (CD41) (3, 11). In the cell membrane, it also associates with a unique splice variant of CD28 (CD28i) that may facilitate CD40L signal transduction (12). CD40L is expressed by monocytes, NK cells, mast cells, basophils, smooth muscle cells, endothelial cells, dendritic cells, activated and resting B cells, plus activated platelets and CD4+ T cells (13, 14). CD40L ligation of CD40 on dendritic cells (DC) initiates DC maturation and differentiation. CD40L signaling into naïve B cells promotes germinal center formation and isotype switching. With IL-21, CD40L generates IgA plus IgG3; with IL-4, CD40L generates IgG1 secretion (14, 15).

References:

1. Zhang, G. (2004) *Curr. Opin. Struct. Biol.* **14**:154.
2. Hehlgans, T. and K. Pfeffer (2005) *Immunology* **115**:1.
3. Quezada, S.A. *et al.* (2004) *Annu. Rev. Immunol.* **22**:307.
4. Graf, D. *et al.* (1992) *Eur. J. Immunol.* **22**:3191.
5. Hollenbaugh, D. *et al.* (1992) *EMBO J.* **11**:4313.
6. Khandekar, S.S. *et al.* (2001) *Prot. Exp. Purif.* **23**:301.
7. Pietravalle, F. *et al.* (1996) *J. Biol. Chem.* **271**:5965.
8. Garber, E. *et al.* (1999) *J. Biol. Chem.* **274**:33545.
9. Vakkalanka, R.K. *et al.* (1999) *Arthritis Rheum.* **42**:871.
10. Eissner, G. *et al.* (2004) *Cytokine Growth Factor Rev.* **15**:353.
11. Prasad, K.S. *et al.* (2003) *Proc. Natl. Acad. Sci. USA* **100**:12367.
12. Mikolajczak, S.A. *et al.* (2004) *J. Exp. Med.* **199**:1025.
13. Lievens, D. *et al.* (2009) *Thromb. Haemost.* **102**:206.
14. Elgueta, R. *et al.* (2009) *Immunol. Rev.* **229**:152.
15. Avery, D.T. *et al.* (2008) *J. Immunol.* **181**:1767.