

**DESCRIPTION**

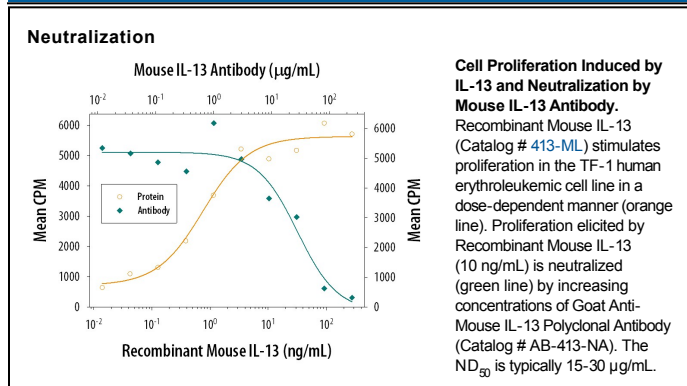
<b>Species Reactivity</b>	Mouse
<b>Specificity</b>	Detects mouse IL-13 in direct ELISAs and Western blots. In direct ELISAs and Westerns, less than 1% cross-reactivity with recombinant human IL-13 is observed. Neutralizes the biological activity of recombinant mouse IL-13, but will not neutralize the biological activity of recombinant human IL-13.
<b>Source</b>	Polyclonal Goat IgG
<b>Purification</b>	Protein A or G purified
<b>Immunogen</b>	<i>E. coli</i> -derived recombinant mouse IL-13 Ser26-Phe131 Accession # P20109
<b>Endotoxin Level</b>	<0.10 EU per 1 µg of the antibody by the LAL method.
<b>Formulation</b>	Lyophilized from a 0.2 µm filtered solution in PBS with Trehalose. See Certificate of Analysis for details.

**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.

	<b>Recommended Concentration</b>	<b>Sample</b>
<b>Western Blot</b>	1 µg/mL	Recombinant Mouse IL-13 (Catalog # 413-ML)
<b>Neutralization</b>		Measured by its ability to neutralize IL-13-induced proliferation in the TF-1 human erythroleukemic cell line. Kitamura, T. <i>et al.</i> (1989) <i>J. Cell Physiol.</i> <b>140</b> :323. The Neutralization Dose (ND <sub>50</sub> ) is typically 15-30 µg/mL in the presence of 10 ng/mL Recombinant Mouse IL-13.

**DATA**



**PREPARATION AND STORAGE**

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

**BACKGROUND**

IL-13 is a 17 kDa immunoregulatory cytokine that plays a key role in the pathogenesis of allergic asthma and atopy. It is secreted by Th1 and Th2 CD4<sup>+</sup> T cells, NK cells, visceral smooth muscle cells, eosinophils, mast cells, and basophils (1-3). IL-13 circulates as a monomer with two internal disulfide bonds that contribute to a bundled four  $\alpha$ -helix configuration (4, 5). Mature mouse IL-13 shares 57%, 75%, and 58% amino acid sequence identity with human, rat, and rhesus IL-13, respectively. Despite the low homology, it exhibits cross-species activity between human, mouse, and rat (6, 7). IL-13 has diverse activities on numerous cell types (8). On macrophages, IL-13 suppresses the production of proinflammatory cytokines and other cytotoxic substances. On B cells, IL-13 induces immunoglobulin class switching to IgE, upregulates the expression of MHC class II, CD71, CD72, and CD23, and costimulates proliferation. IL-13 upregulates IL-6 while downregulating IL-1 and TNF- $\alpha$  production by fibroblasts and endothelial cells. IL-13 binds with low affinity to IL-13 R $\alpha$ 1, triggering IL-13 R $\alpha$ 1 association with IL-4 R $\alpha$ . This high affinity receptor complex also functions as the type 2 IL-4 receptor complex (9, 10). Additionally, IL-13 binds with high affinity to IL-13 R $\alpha$ 2 which is expressed intracellularly, on the cell surface, and as a soluble molecule (11-14). IL-13 R $\alpha$ 2 regulates the bioavailability of both IL-13 and IL-4 and is overexpressed in glioma and several bronchial pathologies (10, 15, 16). Compared to wild type IL-13, the atopy-associated R110Q variant of IL-13 elicits increased responsiveness from eosinophils that express low levels of IL-13 R $\alpha$ 2 (17).

**References:**

1. Wills-Karp, M. (2004) *Immunol. Rev.* **202**:175.
2. Nakajima, H. and K. Takatsu (2007) *Int. Arch. Allergy Immunol.* **142**:265.
3. Brown, K.D. *et al.* (1989) *J. Immunol.* **142**:679.
4. Moy, F.J. *et al.* (2001) *J. Mol. Biol.* **310**:219.
5. Eisenmesser, E.Z. *et al.* (2001) *J. Mol. Biol.* **310**:231.
6. Ruetten, H. and C. Thiemermann (1997) *Shock* **8**:409.
7. Lakkis, F.G. *et al.* (1997) *Biochem. Biophys. Res. Commun.* **235**:529.
8. Wynn, T.A. (2003) *Annu. Rev. Immunol.* **21**:425.
9. Andrews, A.L. *et al.* (2002) *J. Biol. Chem.* **277**:46073.
10. Tabata, Y. *et al.* (2007) *Curr. Allergy Asthma Rep.* **7**:338.
11. Chiamonte, M.G. *et al.* (2003) *J. Exp. Med.* **197**:687.
12. Daines, M.O. and G.K. Hershey (2002) *J. Biol. Chem.* **277**:10387.
13. Matsumura, M. *et al.* (2007) *Biochem. Biophys. Res. Commun.* **360**:464.
14. Tabata, Y. *et al.* (2007) *J. Immunol.* **177**:7905.
15. Andrews, A.L. *et al.* (2006) *J. Allergy Clin. Immunol.* **118**:858.
16. Joshi, B.H. *et al.* (2006) *Vitam. Horm.* **74**:479.
17. Andrews, A-L. *et al.* (2007) *J. Allergy Clin. Immunol.* **120**:91.