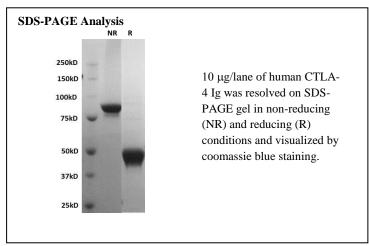


## Recombinant Human CTLA–4 Ig (Fc) Chimera

Catalog Number/Size:	P1001-100 100 μg	5		
	P1001-500 500 μg	5		
	P1001-B Bulk			
Source:	Human CTLA-4 (Accession# NP_005205.2) extracellular domain (Ala37-Phe162) fused with human IgG1 F produced from HEK293 cells.			
	Human CTLA-4 (Ala37-Phe162)	GSGGGG	Human IgG1 (Asp104-Lys330)	
	Accession# NP_005205.2	050000	Tiuman 1gOT (Asp104-Lys550)	
	N-terminal		C-terminal	
Structure:	Disulfide-linked homodimer			
Predicted N-terminal:	Ala 37			
Predicted Molecular Weight:	39.5 kDa (monomer)			
Apparent Molecular Weight on SDS-PAGE:	47.0 kDa, reducing conditions			
Formulation:	0.22 µm filtered protein solution in PBS			
Storage:	-20°C or below			
Estimated Purity:	>95% as determined by SDS-PAGE			
Protein Endotoxin level:	Not measured			
Protein Aggregation:	No obvious protein aggregates			





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## Recombinant Human CTLA-4 Ig (Fc) Chimera

Application:	Bioassay		
Product Description:	CTLA-4 (cytotoxic T lymphocyte-associated molecule-4) is a T cell surface glycoprotein and a member of the immunoglobulin superfamily. The 223-amino acid human protein contains a leader sequence, a single extracellular V domain, a transmembrane domain, and a cytoplasmic tail encoded by 4 exons, respectively (1). It is a disulfide-linked homodimer in the extracellular domain at cysteine residue 120 (2). CD28 is structurally related to CTLA-4, with CTLA-4 sharing 27% (murine) to 31% (human) amino acid identity with CD28 (1). Both CTLA-4 and CD28 bind to B7-1 (CD80) and B7-2 (CD86) on antigen presenting cells, with CTLA-4 binding at a 20- to 50-fold greater affinity than CD28 does (3). CD28 delivers an activation signal to T cells while CTLA-4 transmits an inhibitory signal to T cells (4). CD28 is found in considerable amounts on the cell surface of the majority of resting T cells whereas CTLA-4 presents significantly lower expression, predominantly appearing after T cell activation (5).		
Other Names:	CD152, ALPS5, CELIAC3, GRD4, GSE, IDDM12		
References:	<ol> <li>Harper, K. <i>et al.</i> (1991) J. Immunol. 147:1037.</li> <li>Linsley, P.S. <i>et al.</i> (1995) J. Biol. Chem. 270:15417.</li> <li>van der Merwe, P.A. <i>et al.</i> (1997) J. Exp. Med. 185:393.</li> <li>Krummel, M.F. and Allison, J.P. (1995) J. Exp. Med. 182:459.</li> </ol>		

5. Alegre, M.L. *et al.* (1996) J. Immunol. **157**:4762.

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