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**R**Dsystems

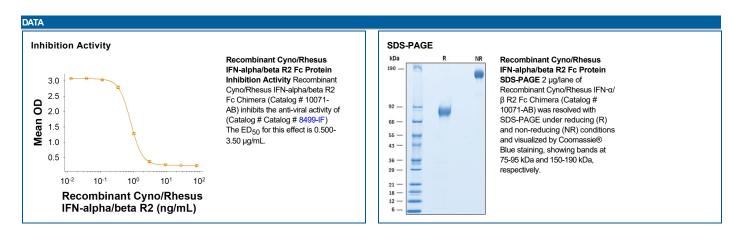
## Recombinant Cynomolgus Monkey/Rhesus Macaque IFN-α/β R2 Fc Chimera

Catalog Number: 10071-AB

Source	Chinese Hamster Ovary cell line, CHO-derived IFN-alpha/beta R2 protein			
	Cyno/Rhesus IFN-alpha R2 (Ile 27-Lys243) Accession # XP_005548871.1	IEGRMD	Human IgG <sub>1</sub> (Pro100-Lys330)	
	N-terminus		C-terminus	
N-terminal Sequence Analysis	lle 27			
Structure / Form	Disulfide-linked homodimer			
Predicted Molecular Mass	51.4 kDa			

SPECIFICATIONS		
SDS-PAGE	75 - 95 kDa, reducing conditions	
Activity	Measured by its ability to inhibit Type-I IFN-mediated anti-viral activity. The ED <sub>50</sub> for this effect is 0.500-3.50 μg/mL in the presence of 30 pg/mL of Recombinant Human IFN-β (Catalog # 8499-IF).	
Endotoxin Level	<0.10 EU per 1 $\mu$ g of the protein by the LAL method.	
Purity	>95%, by SDS-PAGE visualized with Silver Staining and quantitative densitometry by Coomassie® Blue Staining.	
Formulation	Lyophilized from a 0.2 µm filtered solution in PBS. See Certificate of Analysis for details.	

PREPARATION AND STORAGE		
Reconstitution	Reconstitute at 500 μg/mL in PBS.	
Shipping	The product is shipped at ambient temperature. Upon receipt, store it immediately at the temperature recommended below.	
Stability & Storage	<ul> <li>Use a manual defrost freezer and avoid repeated freeze-thaw cycles.</li> <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>3 months, -20 to -70 °C under sterile conditions after reconstitution.</li> </ul>	



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### BACKGROUND

**R**DSYSTEMS

IFN-alpha/beta R2, also known as IFNAR2, is a 100 kDa glycoprotein in the class II cytokine receptor family. These proteins form heterodimeric receptor complexes that transduce signals from the interferon, IL-10, and IL-28 families of cytokines (1, 2). Mature IFN-alpha / beta R2 consists of an extracellular domain (ECD), containing two fibronectin type III repeats, a transmembrane segment, and a cytoplasmic domain. Alternative splicing generates a secreted isoform that corresponds to the ECD as well as a 50 kDa transmembrane isoform with a substituted and truncated cytoplasmic region (3, 4). The short isoform is impaired in its ability to activate signaling molecules and functions as a dominant negative receptor subunit (5-7). The mature ECD of cynomolgus IFN-alpha/ beta R2 shares 93% and 100% amino acid (aa) sequence identity with that of human and rhesus, respectively. IFN-alpha/beta R2, in association with IFN-alpha/beta R1, is required for mediating the antiviral, antiproliferative, and apoptotic effects of the type I interferons IFN-alpha and IFN-beta. IFN-alpha / beta R2 is the principal ligand binding subunit of the receptor. Ligand binding is stabilized by the subsequent association with IFN-alpha / beta R1 is equired for complex (8, 9). IFN-alpha / beta R2 is also subject to presenilin-dependent intramembrane proteolysis, resulting in the liberation of nearly the entire ECD as well as the cytoplasmic domain which migrates to the nucleus and can inhibit gene transcription (10). High concentrations of soluble IFN-alpha / beta R2 bin and neutralize IFN-alpha / beta R2 is also subject to presenilin-dependent intramembrane proteolysis, resulting in the liberation of nearly the entire ECD as well as the cytoplasmic domain which migrates to the nucleus and can inhibit gene transcription (10). High concentrations of soluble IFN-alpha / beta R2 bin and neutr

#### References:

- 1. Langer, J.A. et al. (2004) Cytokine Growth Factor Rev. 15:33.
- 2. Pestka, S. et al. (2004) Immunol. Rev. 202:8.
- 3. Lutfalla, G. et al. (1995) EMBO J. 14:5100.
- 4. Novick, D. et al. (1995) J. Leukocyte Biol. 57:712.
- 5. Pfeffer, L.M. et al. (1997) J. Biol. Chem. 272:11002.
- 6. Gazziola, C. *et al.* (2005) Int. J. Oncol. **26**:129.
- 7. Kotenko, S.V. and S. Pestka (2000) Oncogene 19:2557.
- 8. Lamken, P. *et al.* (2004) J. Mol. Biol. **341**:303.
- 9. Arduini, R.M. *et al.* (1999) Prot. Sci. **8**:1867.
- 10. Saleh, A.Z.M. *et al.* (2004) Oncogene **23**:7076.
- 11. McKenna, S.D. *et al.* (2004) J. Interferon Cytokine Res. **24**:119.
- 12. Tyler, D.R. et al. (2007) Mol. Immunol. 44:1864.

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