

## Human Interferon Gamma Recombinant Protein

<b>Catalog No.</b>	CR2025	<b>Quantity:</b>	50 µg
<b>Concentration:</b>	0.5 mg / ml		
<b>Description:</b>	Recombinant Human Interferon Gamma (hIFN-γ) has a MW = 16.775 kDa.		
<b>Source:</b>	cDNA obtained from human T lymphocyte mRNA expressed in <i>E. Coli</i> .		
<b>Formulation:</b>	Phosphate buffered saline (PBS) containing 0.1% bovine serum albumin (BSA).		
<b>Purity:</b>	>95%		
<b>Biological Activity:</b>	Interferon was titrated with the use of a cytopathic effect inhibition assay. In this antiviral assay for interferon about 1 unit/ml of interferon is the quantity necessary to produce a cytopathic effect of 50%. The units are determined with respect to the international reference standard for human interferon gamma (hIFN-γ) provided by the National Institutes of Health [see Pestka, S. (1986) "Interferon Standards and General Abbreviations," in Methods in Enzymology (S. Pestka, ed.), Academic Press, New York 119, 14-23]. Unit of activity measured on VERO cells with vesicular stomatitis virus (VSV).		
<b>Specific Activity:</b>	1.59 x 10 <sup>7</sup> units/mg		
<b>Storage &amp; Stability:</b>	After receipt, this product should be kept at -70°C or below for retention of full activity. When thawing, the contents of the tube should be apportioned in separate tubes in order to <b>avoid repeated freeze-thaw cycles</b> . Dilute only in buffers containing protein such as 0.1% bovine serum albumin (BSA).		
<b>Country of Origin:</b>	USA		

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