

**GMP® IFN gamma (Interferon gamma), Human**

v. 250301

<b>Catalog number</b>	C01080-GMP-100 / C01080-GMP-1000
<b>Package</b>	100 µg / 1 mg
<b>Description</b>	Interferon gamma (IFN gamma) is an effective multifunctional cytokine which is released mainly by activated NK cells and T cells. IFN gamma is primarily characterized based on its anti-viral activities, and then been proved to have several functions such as anti-proliferative, immune-regulatory, and pro-inflammatory activities. IFN gamma can upregulate expression of MHC class I and II antigen by antigen-presenting cells.
<b>Expression System</b>	<i>Escherichia coli</i>
<b>Species of Origin</b>	Human
<b>Affinity Tag</b>	His Tag (C-term)
<b>Sequence</b>	Gln24-Gln166
<b>Endotoxin level</b>	<0.05 EU per 1 µg of the protein by the LAL method.
<b>Activity</b>	Measure by its ability to induce cytotoxicity in HT29 cells. The ED <sub>50</sub> for this effect is <1 ng/mL. The specific activity of recombinant human IFN gamma is approximately >2 x 10 <sup>6</sup> IU/mg, which is calibrated against the human IFN Gamma WHO Reference Material (NIBSC code: 87/586).
<b>Purity</b>	>95% as determined by SDS-PAGE analysis.
<b>Mycoplasma</b>	Not detected
<b>Form</b>	Lyophilized
<b>Storage Buffer</b>	Lyophilized from a 0.2 µm filtered solution of PBS, pH 8.0.
<b>Reconstitution</b>	It is recommended to reconstitute the lyophilized protein in sterile H <sub>2</sub> O to a concentration not less than 0.5 mg/mL and incubate the stock solution for at least 20 min to ensure sufficient re-dissolved.
<b>Stability &amp; Storage</b>	This product is stable after storage at: <ul style="list-style-type: none"><li>• -20°C for 12 months in lyophilized state from date of receipt.</li><li>• -20°C or -80°C for 1 month under sterile conditions after reconstitution.</li></ul> Avoid repeated freeze/thaw cycles.

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Croyez GMP® recombinant proteins are manufactured in ISO 13485:2016 and GMP-certified facility. The processes include:

**Specification**

- Animal-free reagent and laboratory
- Manufactured and tested under GMP guideline
- Testing and traceability of raw material
- Records of the maintenance and equipment calibration
- Personnel training records
- Batch-to-batch consistency
- Documentation of QA control and process changes
- Manufactured and tested under an ISO 13485:2016 certified quality management system
- Stability monitor of product shelf-life

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**Reference**

1. Schroder K. et al. (2004) *J Leukoc Biol.* 75,2: 163-89.
2. Schoenborn JR, Wilson CB. (2007) *Adv Immunol.* 96: 41-101.
3. Boehm U. et al. (1997) *Annu Rev Immunol.* 15: 749-95.
4. Ijzermans JN, Marquet RL. (1989) *Immunobiology.* 179,4-5: 456-73.
5. Razaghi A. et al. (2016) *J Biotechnol.* 240: 48-60.



SDS-PAGE analysis of GMP® IFN gamma, Human

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*For Research Use Only. Not for use in diagnostic or therapeutic procedures.*