

Recombinant Equine Interferon-gamma

Information

100034181	
P42160	
Escherichia coli.	
Approximately 16.7 kDa, a single non-glycosylated polypeptide chain containin 143 amino acids.	
QAAFFKEIEN LKEYFNASNP DVGDGGPLFL DILKNWKEDS DKKIIQSQIV SFYFKLFENL KDNQVIQKSM DTIKEDLFVK F <mark>FN</mark> SSTSKLE DFQKLIQIPV NDLKVQRKAI SELIKVMNDL SPKANLRKRK R <mark>SQ</mark> NPFRGRR ALQ	
Sterile Filtered White lyophilized (freeze-dried) powder.	
Use a manual defrost freezer and avoid repeated freeze-thaw cycles - 12 months from date of receipt, -20 to -70 °C as supplied - 1 month, 2 to 8 °C under sterile conditions after reconstitution - 3 months, -20 to -70 °C under sterile conditions after reconstitution	
Lyophilized from a 0.2 μ m filtered concentrated solution in 2 \times PBS, pH 7.4, with 5 % trehalose.	
We recommend that this vial be briefly centrifuged prior to opening to bring the contents to the bottom. Reconstitute in sterile distilled water or aqueous buffer containing 0.1 % BSA to a concentration of 0.1-1.0 mg/mL. Stock solutions should be apportioned into working aliquots and stored at ≤ -20 °C. Further dilutions should be made in appropriate buffered solutions.	
Fully biologically active when compared to standard. The ED as determined by an anti-viral assay using human HeLa cells infected with encephalomyocarditis (EMC) virus is less than 10.0 ng/ml, corresponding to a specific activity of > 1.0 × 10 IU/mg.	
Gel pack.	
Centrifuge the vial prior to opening.	
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Components and Storage

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Components	100µg	500µg	
Recombinant Equine Interferon-gamma	100µg	500µg	

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- 12 months from date of receipt, -20 to -70 °C as supplied
- 1 month, 2 to 8 °C under sterile conditions after reconstitution
- 3 months, -20 to -70 °C under sterile conditions after reconstitution

Quality Control

Quality Control		SIO _{COT}
Purity	> 97 % by SDS-PAGE and HPLC analyses.	P. C. Land of the Co.
Endotoxin	Less than 0.1 EU/μg of rEqIFN-γ as determin	ned by LAL method.

Description

Reference









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