



Catalog No:	97131
Lot No:	XXXXX
Source:	E. coli
Synonyms:	

## Description

Leptin antagonist triple mutant mouse recombinant is a single non-glycosilated polypeptide chain containing 146 amino acids and additional Ala at N-terminus and having a molecular mass of approx. 16 kDa. Mouse leptin antagonist was mutated, resulting in L39A/D40A/F41A mutant. Mouse leptin antagonist is bound to 20 kDa mono-PEG at N-terminus, resulting in 35. 6 kDa. The mouse leptin anatagonist runs as a 48 kDa. Leptin antagonist triple mutant was purified by proprietary chromatographic techniques.

#### **Physical Appearance**

White lyophilized (freeze-dried) powder.

## Formulation

Mouse leptin anatagonist triple mutant was lyophilized from a concentrated (0.65 mg/ml) solution with 0.003 mM NaHCO3.

#### Solubility

It is recommended to reconstitute the lyophilized leptin antagonist triple mutant in sterile water or sterile 0.4% NaHCO<sub>3</sub> adjusted to pH 8-9, not less than 100  $\mu$ g/ml, which can then be further diluted with other aqueous solutions.

### Stability

Lyophilized leptin antagonist triple mutant, although stable at room temperature for several weeks, should be stored desiccated below  $-18^{\circ}$ C. Upon reconstitution at > 0.1 mg/ml Leptin mutant and up to 2 mM and filter sterilization LEP mutant can be stored at 4°C or even room temperature for several weeks making it suitable for long term infusion studies using osmotic pumps. At lower concentration addition of a carrier protein (0.1% HSA or BSA) is suggested. Please prevent freeze-thaw cycles.

### Purity

Greater than 99.0% as determined by (a) Gel filtration analysis, (b) Analysis by SDS-PAGE.

## Activity

Leptin antagonist triple mutant mouse recombinant half-life in circulation after SC injection was over 20 hours. Leptin antagonist triple mutant mouse recombinant is capable of inhibiting leptin-induced proliferation of BAF/3 cells stably transfected with the long form of human leptin receptor. Leptin antagonist triple mutant mouse recombinant in vitro activity is 5 - 6 fold lower than the non-pegylated antagonist, though in vivo it has profound weight gain effect (as compared to the non-pegylated antagonist), resulting mainly from increased food intake.

## Usage

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