

Datasheet

Recombinant mouse Leukemia Inhibitory Factor (LIF)

Catalog Number: PR16102 Product Type: Recombinant protein

Source: Recombinant mouse LIF is expressed in E. coli as a fusion protein with GST using the pGEX expression

system, cleaved from GST moeity with thrombin and purified by HPLC chromatography.

Purity: Greater than 95% by analytical HPLC and SDS-PAGE.

Endotoxin Levels: Endotoxin level is less than 0.1 ng per μg of LIF. Tested negative in both aseptic and microplasmic tests.

Activity: The activity of mouse LIF is determined by the ability to induce differentiation of murine M1 myeloid leukemic

cells. The minimum detectable concentration of mouse LIF in this assay is 0.5 ng/mL. The specific activity is >1 x 108 units/mg, where 50 units is defined as the amount of mouse LIF required to induce differentiation in

50% of the M1 colonies in 1 mL agar cultures.

Format: Liquid in PBS, pH 7.4 and 0.02% Tween 20. No preservatives added.

Storage: Maintain at 2-8°C until expiration date. Further dilutions should be made into buffer or

medium to which protein (e.g., 1% BSA) or Tween 20 has been added.

Leukemia Inhibitory Factor (LIF) is a lymphoid factor which promotes long-term maintenance of embryonic stem cells by suppressing spontaneous differentiation. LIF has a number of other activities including cholinergic neuron differentiation, control of stem cell pluripotency, bone and fat metabolism, mitogenesis of certain factor dependent cell lines and promotion of megakaryocyte production *in vivo*. Mouse LIF is a 20 kDa protein containing 181 amino acid residues.

REFERENCES:

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- 3. Vogiagis D. and Salamonsen L.A. (1999). The role of leukemia inhibitory factor in the establishment of pregnancy. *Journal of Endocrinology*. **160**: 181-190.
- 4. Kami K., Morikawa Y., Kawai Y. and Senba E. (1999). Leukemia Inhibitor Factor, Glial cell line derived neurotrophic factor, and their receptor expressions following muscle crush injury. *Muscle & Nerve* 22: 1576-1586.
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RECOMMENDED PROTOCOL

M1 Bioassay

- 1. The M1 bioassay is performed using *in vitro* semi-solid agar cultures, which contain approximately 100 cells in 1 mL volumes of DME containing 20 % FCS in 0.3% agar.
- 2. Add 100 µL of sample or mLIF (104 units/mL in 5% FCS in isotonic saline) in two-fold serial dilutions in duplicate to 35 mm petri dishes.
- 3. Add 100 µL of 5% FCS in isotonic saline to two control slides.
- 4. Incubate at 37°C in fully humidified atmosphere of 10% CO2 in air for 7 days.
- 5. Score the number of colonies that show differentiation (note: 50 units is defined as the amount of activity which results in 50% of the colonies being differentiated).