

IL-36α, Human, Recombinant

货号: PCK194

产品信息

别名 Interleukin-36 Alpha; FIL1 Epsilon; Interleukin-1 Epsilon; IL-1 Epsilon;

Interleukin-1 Family Member 6; IL-1F6; IL36A; FIL1E; IL1E; IL1F6

物种 Human

表达宿主 E.coli

序列信息 Lys6-Phe158

检索号 Q9UHA7

分子量 17.1 kDa

生物活性 Measured by its ability to induce IL-8 secretion in A431 human epithelial

carcinoma cells. The ED50 for this effect is 1.83 ng/ml.

产品特性

纯度 >95% as determined by reducing SDS-PAGE.

内毒素 <1.0 EU per µg as determined by LAL test.

保存 Lyophilized protein should be stored at -5~-20°C, stable for one year after receipt.

Reconstituted protein solution can be stored at 2-8°C for 2-7 days. Aliquots of

reconstituted samples are stable at -5~-20°C for 3 months.

运输 Ambient temperature or ice pack.

制剂 Lyophilized from a 0.2 µm filtered solution of 20 mM PB, 6% Trehalose, 5mM

EDTA, 0.05% Tween 80, pH7.4.

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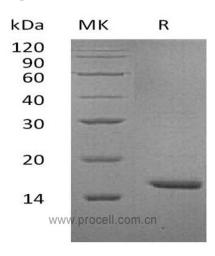
复融

Always centrifuge tubes before opening. Do not mix by vortex or pipetting. It is not recommended to reconstitute to a concentration less than 100 µg/ml.Dissolve the lyophilized protein in distilled water. Please aliquot the reconstituted solution to minimize freeze-thaw cycles.

背景介绍

Human Interleukin-36α (IL-36α) is a secreted Cytokine that belongs to the Interleukin 1 Cytokine family. IL-36α is expressed in the immune system and the fetal brain, but not in other tissues or multiple hematopoietic cell lines. IL-36α is the only IL-1 family member found to be expressed on Tcells. IL-36 α and IL-1F8 are involved in the regulation of adipose tissue gene expression. Importantly, IL-36α inhibits PPARγ expression, which may lead to a reduction in adipocyte differentiation suggesting metabolic effects of this Cytokine. IL-36α, along with IL-1F8 and IL-1F9, has been shown to act as an agonist by activating the pathway involving NFkB and MAPK in an IL-1Rrp2 dependent manner. This suggest that IL-36α may signal in a similar fashion to IL-1 and IL-18 in having a binding Receptor which upon ligation, recruits a second Receptor as a signaling component, forming an active heterodimeric Receptor complex.

SDS-PAGE



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