abcam

Product datasheet

Recombinant Human TGF beta 1 protein ab130946

描述

产品名称 重组人TGF beta 1蛋白

纯**度** > 98 % SDS-PAGE.

> 98% by HPLC

表达系统 HEK 293 cells

Accession P01137

蛋白长度 Full length protein

无动物成分 No

性质 Recombinant

种属 Human

序列 ALDTNYCFSSTEKNCCVRQLYIDFRKDLGWKWIHEPKGYHAN

FCLGPCPY

 ${\tt IWSLDTQYSKVLALYNQHNPGASAAPCCVPQALEPLPIVYYV}$

GRKPKVEQ LSNMIVRSCKCS

预测分子量 13 kDa

氨基酸 279 to 390

技术指标

Our Abpromise guarantee covers the use of ab130946 in the following tested applications.

The application notes include recommended starting dilutions; optimal dilutions/concentrations should be determined by the end user.

应**用** SDS-PAGE

HPLC

形式 Lyophilized

制备和贮存

稳定性和存储 Shipped at 4°C. After reconstitution store at -20°C. Avoid freeze / thaw cycles.

Constituent: 0.1% Trifluoroacetic acid

复溶 Resuspend in 10 mM citric acid pH 3.0 to a concentration of 0.1-1.0 mg/ml. Do not vortex. For

extended storage dilute further in a buffer containing a carrier protein (for example 0.1% BSA).

Store reconstituted protein at -20°C. Avoid repeated thawing and freezing.

常规信息

功能 Multifunctional protein that controls proliferation, differentiation and other functions in many cell

types. Many cells synthesize TGFB1 and have specific receptors for it. It positively and negatively regulates many other growth factors. It plays an important role in bone remodeling as it is a potent stimulator of osteoblastic bone formation, causing chemotaxis, proliferation and differentiation in

committed osteoblasts.

组织特异性 Highly expressed in bone. Abundantly expressed in articular cartilage and chondrocytes and is

increased in osteoarthritis (OA). Co-localizes with ASPN in chondrocytes within OA lesions of

articular cartilage.

疾病相关 Defects in TGFB1 are the cause of Camurati-Engelmann disease (CE) [MIM:131300]; also

known as progressive diaphyseal dysplasia 1 (DPD1). CE is an autosomal dominant disorder characterized by hyperostosis and sclerosis of the diaphyses of long bones. The disease typically presents in early childhood with pain, muscular weakness and waddling gait, and in some cases other features such as exophthalmos, facial paralysis, hearing difficulties and loss of vision.

序列相似性 Belongs to the TGF-beta family.

翻译后修饰 Glycosylated.

The precursor is cleaved into mature TGF-beta-1 and LAP, which remains non-covalently linked to

mature TGF-beta-1 rendering it inactive.

细胞定位 Secreted > extracellular space > extracellular matrix.

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