

Recombinant Human Transforming Growth Factor Beta-1/TGFB1 Biotinylated

Catalog No. PKSH034059

Note: Centrifuge before opening to ensure complete recovery of vial contents.

Description

Synonyms Transforming Growth Factor Beta-1;TGF-Beta-1;Latency-Associated

Peptide;LAP;TGFB1;TGFB

Species Human

Expression Host HEK293 Cells **Sequence** Ala279-Ser390

AccessionP01137Calculated Molecular Weight14.6 kDaObserved molecular weight12-18 kDaTagN-Avi

Bioactivity Measured by its ability to inhibit TGF-beta 1 activity on TF-1 human

erythroleukemic cells. The ED₅₀ for this effect is 137.37pg/ml.

Properties

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

Endotoxin < 0.01 EU per μg of the protein as determined by the LAL method.

Storage Generally, lyophilized proteins are stable for up to 12 months when stored at -20 to

-80°C. Reconstituted protein solution can be stored at 4-8°C for 2-7 days. Aliquots

of reconstituted samples are stable at < -20°C for 3 months.

Shipping This product is provided as lyophilized powder which is shipped with ice packs.

Formulation Lyophilized from a 0.2 µm filtered solution of 50mM Glycine-HCl, 150mM NaCl,

pH2.5.

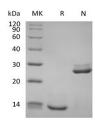
Normally 5% - 8% trehalose, mannitol and 0.01% Tween 80 are added as

protectants before lyophilization.

Please refer to the specific buffer information in the printed manual.

Reconstitution Please refer to the printed manual for detailed information.

Data



> 95 % as determined by reducing SDS-PAGE.

For Research Use Only

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Background

Transforming Growth Factor β -1 (TGF β -1) is a secreted protein which belongs to the TGF- β family. TGF β -1 is abundantly expressed in bone, articular cartilage and chondrocytes and is increased in osteoarthritis (OA). TGF β -1 performs many cellular functions, including the control of cell growth, cell proliferation, cell differentiation and apoptosis. The precursor is cleaved into a latency-associated peptide (LAP) and a mature TGFβ-1 peptide. TGFβ-1 may also form heterodimers with other TGFβ family members. It has been found that TGFβ-1 is frequently upregulated in tumor cells. Mutations in this gene results in Camurati-Engelmann disease.

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