

PRODUCT INFORMATION

TGFB2 **Target**

LDS4; G-TSF; TGF-beta2 Synonyms

Recombinant human TGFB2(303-414) protein Description

with N-terminal human Fc tag

Delivery In Stock **Uniprot ID** P61812 **Expression Host HEK293**

Tag N-Human Fc tag

Molecular

Reconstitution

Background

Storage & Shipping

hFc(Glu99-Ala330) TGFB2(Ala303-Ser414) Characterization

The protein has a predicted molecular mass of **Molecular Weight**

38.9 kDa after removal of the signal peptide. The apparent molecular mass of hFc-TGFB2(303-414) is approximately 35-55 kDa due to glycosylation.

The purity of the protein is greater than 95% as determined by SDS-PAGE and Coomassie blue Purity

staining.

Lyophilized from sterile PBS, pH 7.4. Normally 5 % Formulation &

- 8% trehalose is added as protectants before lyophilization. Please see Certificate of Analysis for specific instructions of reconstitution.

Store at -20°C to -80°C for 12 months in lyophilized form. After reconstitution, if not intended for use within a month, aliquot and store

at -80°C (Avoid repeated freezing and thawing). Lyophilized proteins are shipped at ambient

temperature.

This gene encodes a secreted ligand of the TGF-

beta (transforming growth factor-beta)

superfamily of proteins. Ligands of this family bind various TGF-beta receptors leading to recruitment and activation of SMAD family transcription factors that regulate gene expression. The encoded preproprotein is proteolytically processed to generate a latencyassociated péptide (LAP) and a mature peptide, and is found in either a latent form composed of a

mature peptide homodimer, a LAP homodimer, and a latent TGF-beta binding protein, or in an active form consisting solely of the mature peptide homodimer. The mature peptide may also form heterodimers with other TGF-beta family members. Disruption of the TGF-beta/SMAD pathway has been implicated in a variety of human cancers. A chromosomal translocation that includes this gene is associated with Peters' anomaly, a congenital defect of the anterior

chamber of the eye. Mutations in this gene may be associated with Loeys-Dietz syndrome. This gene encodes multiple isoforms that may undergo similar proteolytic processing. [provided by RefSeq, Aug 2016]

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Usage Research use only

Unconjugated Conjugate

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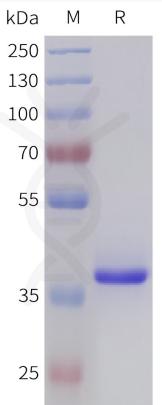


Figure 1. Human TGFB2(303-414) Protein, hFc Tag on SDS-PAGE under reducing condition.

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