

Product datasheet

Recombinant Human EF-Ts protein (denatured) ab181937

[1 Image](#)

Description

| | |
|--|---|
| Product name | Recombinant Human EF-Ts protein (denatured) |
| Purity | > 85 % SDS-PAGE. |
| Expression system | Escherichia coli |
| Accession | <u>P43897-2</u> |
| Protein length | Full length protein |
| Animal free | No |
| Nature | Recombinant |
| Species | Human |
| Sequence | MSKELLMKLRRTGYSFVNCKKALETGGDLKQAEIWLH KEAQKEGWSKA AKLQGRKTKEGLIGLLQEGNTTVLVEVNCETDFVSRNLKF QLLVQQVALG TMMHCQTLKDQPSAYSQVQWLTPVNLALWEAEAGGSLE GFLNSSELSGLP AGPDREGSLKDQLALAIGKLGEMILKRAAWVKVPSGFYV GSYVHGAMQS PSLHKLVLGKYGALVICETSEQKTNLEDVGRRLGQHVVGM APLSVGSLEDD EPGGEAETKMLSQPYLLDPSITLGQYVQPQGVSVVDFVR FECGEGEEAAE TE |
| Predicted molecular weight | 33 kDa |
| Amino acids | 46 to 346 |
| Additional sequence information | NP_005717.2 |

Specifications

Our **Abpromise guarantee** covers the use of **ab181937** in the following tested applications.

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

Applications SDS-PAGE

Form Liquid

Preparation and Storage

Stability and Storage Shipped at 4°C. Store at +4°C short term (1-2 weeks). Upon delivery aliquot. Store at -20°C or -80°C. Avoid freeze / thaw cycle.

pH: 8.00
Constituents: 10% Glycerol (glycerin, glycerine), 2.4% Urea, 0.32% Tris HCl

General Info

Function Associates with the EF-Tu.GDP complex and induces the exchange of GDP to GTP. It remains bound to the aminoacyl-tRNA.EF-Tu.GTP complex up to the GTP hydrolysis stage on the ribosome.

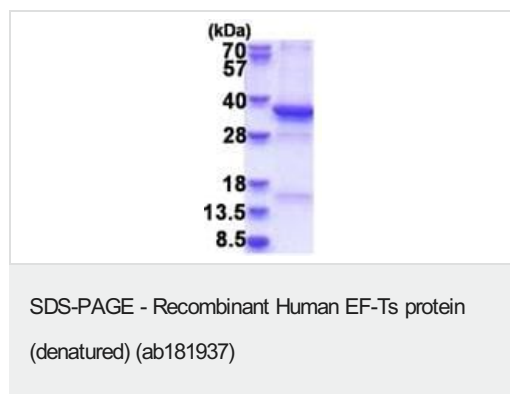
Tissue specificity Expressed in all tissues, with the highest levels of expression in skeletal muscle, liver and kidney.

Involvement in disease Defects in TFSM are the cause of combined oxidative phosphorylation deficiency type 3 (COXPD3) [MIM:610505]. Defects in the mitochondrial oxidative phosphorylation system result in devastating, mainly multisystem, diseases. COXPD3 symptoms include severe metabolic acidosis with encephalomyopathy or with hypertrophic cardiomyopathy. Patients show a severe defect in mitochondrial translation leading to a failure to assemble adequate amounts of three of the oxidative phosphorylation complexes.

Sequence similarities Belongs to the EF-Ts family.

Cellular localization Mitochondrion.

Images



15% SDS-PAGE analysis of 3 µg ab181937.

Please note: All products are "FOR RESEARCH USE ONLY. NOT FOR USE IN DIAGNOSTIC PROCEDURES"

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