

# **Product datasheet for TP301237**

#### OriGene Technologies, Inc.

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### FKBP12 (FKBP1A) (NM\_000801) Human Recombinant Protein

**Product data:** 

**Product Type:** Recombinant Proteins

**Description:** Recombinant protein of human FK506 binding protein 1A, 12kDa (FKBP1A), transcript variant

12B, 20 µg

Species: Human Expression Host: HEK293T

**Expression cDNA Clone** >RC201237 protein sequence or AA Sequence: Red=Cloning site Green=Tags(s)

MGVQVETISPGDGRTFPKRGQTCVVHYTGMLEDGKKFDSSRDRNKPFKFMLGKQEVIRGWEEGVAQMS

۷G

QRAKLTISPDYAYGATGHPGIIPPHATLVFDVELLKLE

**TRTRPLEQKLISEEDLAANDILDYKDDDDKV** 

Tag: C-Myc/DDK

Predicted MW: 11.8 kDa

**Concentration:** >0.05 μg/μL as determined by microplate BCA method

**Purity:** > 80% as determined by SDS-PAGE and Coomassie blue staining

**Buffer:** 25 mM Tris-HCl, 100 mM glycine, pH 7.3, 10% glycerol

**Preparation:** Recombinant protein was captured through anti-DDK affinity column followed by

conventional chromatography steps.

**Note:** For testing in cell culture applications, please filter before use. Note that you may experience

some loss of protein during the filtration process.

Storage: Store at -80°C.

Stability: Stable for 12 months from the date of receipt of the product under proper storage and

handling conditions. Avoid repeated freeze-thaw cycles.

**RefSeq:** NP 000792

**Locus ID:** 2280

UniProt ID: P62942



#### FKBP12 (FKBP1A) (NM\_000801) Human Recombinant Protein - TP301237

RefSeq Size: 1643

Cytogenetics: 20p13 RefSeq ORF: 324

Synonyms: FKBP-1A; FKBP-12; FKBP1; FKBP12; PKC12; PFIASE

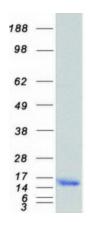
**Summary:** The protein encoded by this gene is a member of the immunophilin protein family, which play

a role in immunoregulation and basic cellular processes involving protein folding and trafficking. The protein is a cis-trans prolyl isomerase that binds the immunosuppressants FK506 and rapamycin. It interacts with several intracellular signal transduction proteins including type I TGF-beta receptor. It also interacts with multiple intracellular calcium release channels, and coordinates multi-protein complex formation of the tetrameric skeletal muscle ryanodine receptor. In mouse, deletion of this homologous gene causes congenital heart disorder known as noncompaction of left ventricular myocardium. Multiple alternatively spliced variants, encoding the same protein, have been identified. The human genome contains five pseudogenes related to this gene, at least one of which is transcribed. [provided

by RefSeq, Sep 2008]

**Protein Families:** Druggable Genome

## **Product images:**



Coomassie blue staining of purified FKBP1A protein (Cat# TP301237). The protein was produced from HEK293T cells transfected with FKBP1A cDNA clone (Cat# [RC201237]) using MegaTran 2.0 (Cat# [TT210002]).