

## **Product datasheet for TP720562M**

# OriGene Technologies, Inc.

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## Desmin (DES) (NM\_001927) Human Recombinant Protein

#### **Product data:**

**Product Type:** Recombinant Proteins

**Description:** Recombinant protein of human desmin (DES)

Species: Human
Expression Host: E. coli

**Expression cDNA Clone** 

Val260-Leu470

or AA Sequence:

Tag: N-His

Predicted MW: 26.7 kDa

Concentration: lot specific

**Purity:** >95% as determined by SDS-PAGE and Coomassie blue staining

Buffer: Provided lyophilized from a 0.2 μm filtered solution of 20 mM Tris-HCl, 150 mM NaCl

**Endotoxin:** < 0.1 EU per μg protein as determined by LAL test

**Reconstitution Method:** Always centrifuge tubes before opening. Do not mix by vortex or pipetting. Dissolve the

lyophilized protein in ddH2O. It is not recommended to reconstitute a concentration less than 100 µg/ml. Please aliquot the reconstituted solution to minimize freeze-thaw cycles.

Storage: Store at -80°C.

Stability: Stable for at least 6 months from date of receipt under proper storage and handling

conditions.

**RefSeq:** NP 001918

**Locus ID:** 1674

**UniProt ID:** <u>P17661</u>, <u>Q53SB5</u>

**Cytogenetics:** 2q35

Synonyms: CDCD3; CSM1; CSM2; LGMD1D; LGMD1E; LGMD2R





### Desmin (DES) (NM\_001927) Human Recombinant Protein - TP720562M

Summary: This gene encodes a muscle-specific class III intermediate filament. Homopolymers of this

protein form a stable intracytoplasmic filamentous network connecting myofibrils to each other and to the plasma membrane. Mutations in this gene are associated with desminrelated myopathy, a familial cardiac and skeletal myopathy (CSM), and with distal myopathies.

[provided by RefSeq, Jul 2008]

**Protein Families:** Druggable Genome

Protein Pathways: Arrhythmogenic right ventricular cardiomyopathy (ARVC), Dilated cardiomyopathy,

Hypertrophic cardiomyopathy (HCM)

## **Product images:**

