

## **Product datasheet for TP520185**

## OriGene Technologies, Inc.

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## Colec11 (NM\_027866) Mouse Recombinant Protein

**Product data:** 

**Product Type:** Recombinant Proteins

**Description:** Purified recombinant protein of Mouse collectin sub-family member 11 (Colec11), with C-

terminal MYC/DDK tag, expressed in HEK293T cells, 20ug

Species: Mouse

**Expression Host:** HEK293T

**Expression cDNA Clone** >MR220185 representing NM\_027866 or AA Sequence: Red=Cloning site Green=Tags(s)

MMMRDLALAGMLISLAFLSLLPSGCPQQTTEDACSVQILVPGLKGDAGEKGDKGAPGRPGRVGPTGEKGD MGDKGQKGTVGRHGKIGPIGAKGEKGDSGDIGPPGPSGEPGIPCECSQLRKAIGEMDNQVTQLTTELKFI KNAVAGVRETESKIYLLVKEEKRYADAQLSCQARGGTLSMPKDEAANGLMASYLAQAGLARVFIGINDLE KEGAFVYSDRSPMQTFNKWRSGEPNNAYDEEDCVEMVASGGWNDVACHITMYFMCEFDKENL

TRTRPLEQKLISEEDLAANDILDYKDDDDKV

Tag: C-MYC/DDK

**Predicted MW:** 29 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

**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 after receiving vials.

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 082142

**Locus ID:** 71693

UniProt ID: Q3SXB8, A0A0R4J0M6

RefSeq Size: 1383





## Colec11 (NM\_027866) Mouse Recombinant Protein – TP520185

Cytogenetics: 12 A2

RefSeq ORF: 816 Synonyms: CL-K1

Summary: This gene encodes a member of the collectin family of C-type lectins that possess collagen-like

sequences and carbohydrate recognition domains. Collectins are secreted proteins that play important roles in the innate immune system by binding to carbohydrate antigens on microorganisms, facilitating their recognition and removal. The encoded protein binds to multiple sugars with a preference for fucose and mannose. Mutations in the human gene are a cause of 3MC syndrome-2. Alternative splicing results in multiple transcript variants encoding

different isoforms. [provided by RefSeq, Sep 2015]