

Product datasheet for **RC221796**

Filamin B (FLNB) (NM_001457) Human Tagged ORF Clone

Product data:

Product Type:	Expression Plasmids
Product Name:	Filamin B (FLNB) (NM_001457) Human Tagged ORF Clone
Tag:	Myc-DDK
Symbol:	FLNB
Synonyms:	ABP-278; ABP-280; AOI; FH1; FLN-B; FLN1L; LRS1; SCT; TAPP; TAP
Mammalian Cell Selection:	Neomycin
Vector:	pCMV6-Entry (PS100001)
E. coli Selection:	Kanamycin (25 ug/mL)
ORF Nucleotide Sequence:	>RC221796 representing NM_001457 Red=Cloning site Blue=ORF Green=Tags(s)

TTTTGTAATACGACTCACTATAGGGCGCCGGGAATTCGTCGACTGGATCCGGTACCGAGGAGATCTGCC
GCC**CGATCGCC**

ATGCCGTTAACCGAGAAGGATCTAGCTGAGGACGCGCCTTGAAGAAGATCCAGCAGAACACGTTACAC
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CACATCCCTGGCAGCCCTTTTCATGTACAGTGCCT

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Protein Sequence:

>RC221796 representing NM_001457
 Red=Cloning site Green=Tags(s)

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TRTRPLEQKLI SEEDLAANDILDYKDDDDKV

Restriction Sites:

Sgfl-Mlul

Components:	The ORF clone is ion-exchange column purified and shipped in a 2D barcoded Matrix tube containing 10ug of transfection-ready, dried plasmid DNA (reconstitute with 100 ul of water).
Reconstitution Method:	<ol style="list-style-type: none">1. Centrifuge at 5,000xg for 5min.2. Carefully open the tube and add 100ul of sterile water to dissolve the DNA.3. Close the tube and incubate for 10 minutes at room temperature.4. Briefly vortex the tube and then do a quick spin (less than 5000xg) to concentrate the liquid at the bottom.5. Store the suspended plasmid at -20°C. The DNA is stable for at least one year from date of shipping when stored at -20°C.
RefSeq:	NM_001457.4
RefSeq Size:	9467 bp
RefSeq ORF:	7809 bp
Locus ID:	2317
UniProt ID:	O75369
Cytogenetics:	3p14.3
Protein Pathways:	Focal adhesion, MAPK signaling pathway
MW:	278.2 kDa
Gene Summary:	This gene encodes a member of the filamin family. The encoded protein interacts with glycoprotein Ib alpha as part of the process to repair vascular injuries. The platelet glycoprotein Ib complex includes glycoprotein Ib alpha, and it binds the actin cytoskeleton. Mutations in this gene have been found in several conditions: atelosteogenesis type 1 and type 3; boomerang dysplasia; autosomal dominant Larsen syndrome; and spondylocarpotarsal synostosis syndrome. Multiple alternatively spliced transcript variants that encode different protein isoforms have been described for this gene. [provided by RefSeq, Nov 2009]