

## Product datasheet for SC113748

### FANCL (NM\_018062) Human Untagged Clone

#### Product data:

Product Type:	Expression Plasmids
Product Name:	FANCL (NM_018062) Human Untagged Clone
Tag:	Tag Free
Symbol:	FANCL
Synonyms:	FAAP43; PHF9; POG
Mammalian Cell Selection:	None
Vector:	<u>pCMV6-XL5</u>
E. coli Selection:	Ampicillin (100 ug/mL)
Fully Sequenced ORF:	>OriGene ORF within SC113748 sequence for NM_018062 edited (data generated by NextGen Sequencing)

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ATGGCGGTGACGGAAGCGAGCCTGTTGCGCCAGTGCCCCCTGCTTCTGCCCCAGAACCGG
TCGAAAACCGTGTATGAGGGATTTCATCTCGGCTCAGGGAAGAGACTTCCACCTTAGGATA
GTGTTGCCTGAAGATTTACAACGAAGAATGCAAGATTATTATGTAGTTGGCAGCTGAGA
ACAATACTTAGTGATACCATCGAATAGTACAACAGAGAATGCAGCACTCTCCTGATCTA
ATGAGCTTTATGATGGAGTTGAAGATGCTTTTGGAAAGTTGCCTTAAAGAATAGACAAGAG
CTGTATGCACTACCTCCTCCTCCCGAGTCTACTCAAGCCTTATTGAAGAGATAGGAACT
CTTGTTGGGATAAACTTGTGTATGCGGATACCTGCTTCAGTACCATCAAGTAAAAGCA
GAAGATGCTTCTGGTAGAGAGCATTTAATCACTCTCAAGTTGAAGGCAAAGTATCCTGCA
GAATCACCAGATTATTTTGTGGATTTTCTGTTCCATTTTGTGCCTCCTGGACACCTCAG
AGCTCCTTAATAAGCATTATAGTCAGTTTTTGGCAGCAATAGAATCACTAAAGGCATTC
TGGGATGTTATGGATGAAATCGATGAGAAGACCTGGGTACTTGAGCCAGAAAAACCTCCA
CGGAGTGCAACAGCAGCAGAAATTGCATTAGGTAATAATGTTTCCATAAATATAGAGGTA
GACCCAGGCATCCTACTATGCTTCTGAGTGCTTCTTTTTTGGAGCTGACCATGTGGTA
AAACCCCTGGGAATTAAGCTGAGCAGGAACATACATTTGTGGGATCCAGAAAAATAGTGTG
TTACAAAATTTGAAAGATGTTTTAGAAAATTGATTTTCCAGCTCGTGCTATCCTGGAAAAA
TCTGATTTTACTATGGATTGTGGAATTTGTTATGCTTATCAACTGACGGTACCATTCT
GATCAAGTGTGTGATAATTCCCAGTGTGGACAACCTTTCCATCAAATATGCTTATATGAG
TGGCTGAGAGGACTACTAACTAGTAGACAGAGTTTTAACATCATATTTGGTGAATGTCCA
TATTGTAGTAAGCCAATTACCTTAAAAATGTCTGGAAGGAAACACTGA

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Clone variation with respect to NM\_018062.3  
760 c=>t;981 t=>c



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**5' Read Nucleotide Sequence:**

>OriGene 5' read for NM\_018062 unedited  
 NNGGGTGCACATTTGTATACGACTCACTATAGGCGGCCGCGATTCCGGCAGGAGGCAGGTC  
 TAGAGCTTTTTCTGTGTTTCTCCGGACTTCGAGCCATGGCGGTGACGGAAGCGAGCCTGTT  
 GCGCCAGTGCCCCCTGCTTCTGCCCCAGAACCGGTGAAAACCGTGTATGAGGGATTCAT  
 CTCGGCTCAGGGAAGAGACTTCCACCTTAGGATAGTGTTCCTGAAGATTACAACGAA  
 GAATGCAAGATTATTATGTAGTTGGCAGCTGAGAACAATACTTAGTGGATACCATCGAAT  
 AGTACAACAGAGAATGCAGCACTCCTGATCTAATGAGCTTTATGATGGAGTTGAAGAT  
 GCTTTTGGAAAGTTGCCTTAAAGAATAGACAAGAGCTGTATGCACTACCTCCTCCTCCCA  
 GTTCTACTCAAGCCTTATTGAAGAGATAGAACTCTTGGTTGGGATAAACTTGTGTATGC  
 GGATACCTGCTTCAGTACCATCAAGTAAAAGCAGAAGATGCTTCTGGTAGAGAGCATT  
 AATCACTCTCAAGTTGAAGGCAAAGTATCCTGCAGAATCACCAGATTATTTTGTGGATT  
 TCCTGTTCCATTTTGTGCTCCTGGACACCTCAGAGCTCCTTAATAAGCATTATAGTCA  
 GTTTTTGGCAGCAATAGAATCACTAAAGGCATTCTGGGATGTTATGGATGAAATCGATGA  
 GAAGACCTGNGTACTTGAGCCAGANAAACCTCCACGGAGTGCAACAGCACGCAGAATTGC  
 ATTAGGTAATAATGTTTCCATAAATATAGAGGTAGACCCAGGCATNCTACTATGCTTNC  
 TGAGTGCTCTTTTTTGGAGCTGACCATGTGGTAAAACCCCTGNGAATTAAGCTG

**3' Read Nucleotide Sequence:**

>OriGene 3' read for NM\_018062 unedited  
 ACGCGGCACGCAATCTAGTATCGAGTTTTTTTTTTTTTTTTTTTGGATCTTGATAACATTT  
 TATTTAGCATTCTTACACTACACAAAATAAATACTTGGATAACTCACGTCTAACAAAC  
 TAACTATACTATGTATTCTTTCTCCATAGGGAAAGCACAAGGAGAAGACAGAAATATCA  
 AGAGTCTCAAGAACCTTTGAATGAAGTAAACAGTTTCCCACAAAAATCAGCTATACACA  
 ATTCCCAAACCTATTTATGAGCCTCATCAAGATTTTACCAGTCCAGATATATTCAGAA  
 GTCAAGATCTCCATCTTGGTATAAATACACTTCCACAGTCAGCACGGGGATCACAGACTT  
 AGAAAGTTCAACTGGACTTTGGCCTACAATTTCCAGTTTACTCTTAGTGAAGAGACAAA  
 CGCAGATGTTTATTATTATCGCATCATCATACCTGTCCTTTTGGATGTTAGTATTTCTTGC  
 TTTATTTTTTCTCTGAAGATGATACAAAATTCCTTTTGGATAATTTTTTAAGTTTCCAGC  
 TCTTACCAGAAATGTTGTATTCTTATTTTCAAGTGTTCCTTCCAGACATTTTTAAGGTAAT  
 TGGCTTACTACAATATGGACATTCACCAAAATATGATGTTAAAACCTCTGTCTACTAGTTAG  
 TAGTCCTCTCAGCCACTCATATAAGCATATCTGATGAAAAGGCTGTCCACACTGGGAATT  
 ATCACACACTTGATCAGGAATGGTACCGTCAAGTTGATAAGCCTAACAAATTCACACTC  
 CTATAAAATCAGATTTTCCAGATAGACGAGCTGGAAAATAATTTCTAAACATCTTTCGAAT  
 TTGCACACACTTTTTTGAATCCACAAATGATGTGCTGCTCACTAATCCACAGGTTTACA  
 CATGGCACCTCCAAGAACCCAGGACCTGTAGAGGCCGGGTCTACTT

**Restriction Sites:**

ECoRI-NOT

**ACCN:**

NM\_018062

**Insert Size:**

1730 bp

**OTI Disclaimer:** Due to the inherent nature of this plasmid, standard methods to replicate additional amounts of DNA in E. coli are highly likely to result in mutations and/or rearrangements. Therefore, OriGene does not guarantee the capability to replicate this plasmid DNA. Additional amounts of DNA can be purchased from OriGene with batch-specific, full-sequence verification at a reduced cost. Please contact our customer care team at [custsupport@origene.com](mailto:custsupport@origene.com) or by calling 301.340.3188 option 3 for pricing and delivery.

The molecular sequence of this clone aligns with the gene accession number as a point of reference only. However, individual transcript sequences of the same gene can differ through naturally occurring variations (e.g. polymorphisms), each with its own valid existence. This clone is substantially in agreement with the reference, but a complete review of all prevailing variants is recommended prior to use. [More info](#)

**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:**

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\\_018062.2](#), [NP\\_060532.2](#)

**RefSeq Size:** 1750 bp

**RefSeq ORF:** 1128 bp

**Locus ID:** 55120

**UniProt ID:** [Q9NW38](#)

**Cytogenetics:** 2p16.1

**Protein Pathways:** Ubiquitin mediated proteolysis

**Gene Summary:** This gene encodes a ubiquitin ligase that is a member of the Fanconi anemia complementation group (FANC). Members of this group are related by their assembly into a common nuclear protein complex rather than by sequence similarity. This gene encodes the protein for complementation group L that mediates monoubiquitination of FANCD2 as well as FANCI. Fanconi anemia is a genetically heterogeneous recessive disorder characterized by cytogenetic instability, hypersensitivity to DNA crosslinking agents, increased chromosomal breakage, and defective DNA repair. Alternative splicing results in multiple transcript variants. [provided by RefSeq, May 2018]

Transcript Variant: This variant (2) has an alternate splice site in the CDS, as compared to variant 1. The resulting isoform (2) lacks an internal segment and has identical N- and C-termini, as compared to isoform 1.