

## Product datasheet for **SC120068**

### FANCA (NM\_000135) Human Untagged Clone

#### Product data:

Product Type:	Expression Plasmids
Product Name:	FANCA (NM_000135) Human Untagged Clone
Tag:	Tag Free
Symbol:	FANCA
Synonyms:	FA; FA-H; FA1; FAA; FACA; FAH; FANCH
Mammalian Cell Selection:	None
Vector:	<u>pCMV6-XL4</u>
E. coli Selection:	Ampicillin (100 ug/mL)
Fully Sequenced ORF:	>NCBI ORF sequence for NM_000135, the custom clone sequence may differ by one or more nucleotides

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ATGTCGACTCGTGGGTCCCGAACTCCGCCTCGGGCCAGGACCCAGGGGGCCCGGAGGGCCTGGGCCG
AGCTGCTGGCGGAAGGGTCAAGAGGGAAAAATAATCCTGAAAGGGCACAGAAATTAAGGAATCAGC
TGTGCGCCTCCTGCGAAGCCATCAGGACCTGAATGCCCTTTTGCTTGAGGTAGAAGGTCCACTGTGTAA
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TAGGCTCTGCTTTCAGGATCAAGCCTCAAGGCTGGGGTTCCCGTGGGTATTCTCTCAGCCGGGATGGT
TGCCCTAGCGTGGGACAGATCTGCACGGCTCCAGCGGAGACCAGTCACCCTGTGCTGCTGACTGTGGAG
CAGAGAAAGAAGCTGTCTCCCTGTTAGAGTTTGCTCAGTATTTATTGGCACACAGTATGTTCTCCCGTC
TTTCCTTCTGTCAAGAATTATGGAAAATACAGAGTTCCTTGTGCTTGAAGCGGTGTGGCATCTTACGT
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TTCAGGAATCTGTGCTGCCTTTGTGAACAGATGGAAGCATCCTGCCAGCATGCTGACGTCGCCAGGGCCA
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TGGGGACATTACTGAGCCCCACAGCCAAGCTCTTCAGGATGTTGAAAAGGCCATCATGGTGTGGAGCAT  
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TCCCCGCCTGCTCACACCTCGAGTGCTCCCAAGTCCCTGACTCCCCTGTGGCGTTTATAGAGTCTCT  
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TTATTA AAAAGTTTCAGTTCCTCATGTTGAGATTGTTCTCAGAGGCCCGACAGCCTTTTCTGAGGAGGA  
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ACCTGGAGCTGGAAATCAACCTGAAGCTGATGCTCTTTCAGATACTGAACGGCAGGACTTCCACCAGTG  
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CTGATTTGGTCTTTGGTGGCCGACAGGAAATGAGGATATTATTTCCAGATTGCAGGAGATGGTAGCTGA  
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ATTTTCCGACAGCGCTCCAGGCTCTGACAAGCGGGTGGAGCTGGCTGCCAGCCTTACAGACAGAGGG  
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AGAACAGCCATCACTGCCAGATGCGAGCAGTTCTTCCACTTGGTCAACTCTGAGATGAGAACTTCTGC  
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TGCATGTTGCTGTGGACATGACTTGAAGCTGGTCCAGCTCTTCTGGCTGGGGATACAAGCACAGTTTC  
ACCTCCAGCTGGCAGGAGCCTGGAGCTCAAGGGTCAGGGCAACCCCGTGGAACTGATAACAAAAGCTCGT  
CTTTTTCTGCTGCAGTTAATACCTCGGTGCCGAAAAAGAGCTTCTCACACGTGGCAGAGCTGCTGGCTG  
ATCGTGGGACTGCGACCCAGAGGTGAGCGCCGCCCTCCAGAGCAGACAGCAGGCTGCCCTGACGCTGA  
CCTGTCCCAGGAGCCTCATCTCTTCTGA

**5' Read Nucleotide Sequence:**

>OriGene 5' read for NM\_000135 unedited  
 GGTGCACAATTTGTATACGACTCACTATAGGCGGCCGCGNAATTCGCACGAGGGGCTGTA  
 GGCGCCAAGGCCATGTCCGACTCGTGGTCCCCTCCGCTCGGGCCAGGACCCAGGG  
 GGCCGCCGAGGGGCTGGGCCGAGCTGTGGCGGGAAGGGTCAAGAGGGAAAAATATAAT  
 CCTGAAGGGCACAGAAATTAAGGAATCAGCTGTGCGCCTCTCGAAGCCATCAGGACC  
 TGAATGCCCTTTTGCTTGAGAAGTCCACTGTGTAATAAATTTGCTCTCAGCAAAGTGAT  
 TGACTGTGACAGTTCTGAGGCCTATGCTAATCATTCTAGTTCATTTATAGGCTCGCTTT  
 GCAGGATCAAGCCTCAAGGCTGGGGTTCCCGTGGGTATTCTCTCAGCCGGGATGGTTGC  
 CTCTAGCGTGGGACAGATCTGCACGGCTCCAGCGGAGACCAGTCAACCCTGTGCTGTGAC  
 TGTGGAGCAGAGAAAGAAGCTGTCTCCCTGTTAGAGTTTGCTCAGTATTTATTGGCACA  
 CAGTATGTTCTCCCGTCTTCTCTGTCAAGAATTATGGAAAATACAGAGTTCTTTGTT  
 GCTTGAAGCGGTGTGGCATCTTCACGTACAAGCATTGTGAGCCTGCAAGAGCTGCTGGA  
 AAGCCATCCCACATGCATGCTGTGGGATCGTGGCTCTCAGGATCTGTGCTGCCTTTGT  
 GAACAGATGGGAGCATCCTGCCAGCATGCTGACGTGCCAGGGCCATGCTTTCTGAATTT  
 GTTCAAAGTTTGTGTTGAGGGGATTTAGAAAACTAGTATCTGAGAACAACGTGGAGCC  
 CGAAAAAGCCCGCAGTCCACGTTGAAGTACTGCAGAGAATGCTTAATTTGCACTTGA

**3' Read Nucleotide Sequence:**

>OriGene 3' read for NM\_000135 unedited  
 TGGACCGGGCAGCAATCTAGTGTGCGAGTTTTTTTTTTTTTTTTTTTCCCTAAANCAGT  
 CGAGGAGTATTTGTAATCCACTTTTTAGTGCAACAAGAGCTCCATGTTATGCTTGTAAATA  
 AATTTTACACGGGAGCTGGGCTGGTGTGCAAGTGGCAGGTCCTCCGTCAGAAGATATGAGG  
 CTCTGGGACAGGTCAGCGTCAGGGGCAGCCTGTGTCTGCTCTGGAGGGCGGCGCTCAC  
 CTCTGGGTCGAGTCCCCACGATCAGCCAGCAGCTCTGCCACGTGTGAGAAGCTCTTTTT  
 CGGGCACCGAGGTATTAAGTGCAGCAGAAAAAGACGAGCTTTTGTATCAGTTCACGGG  
 GTTGCCCTGACCTTTGAGCTCCAGGCTCCTGCCAGCTGGAGGTGAAACTGTGCTTGATC  
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 CTCTCCCTGATGGCCCGTCTTCATGGAAGTAGGAGAGAAAGACTGTAAAAAGCGAAAGG  
 CAGCAGCCTGGTGTGCTGATCCGGGGCCACCGGAGGAGAGCCGCCCCAGCCTGAGGTC  
 ACTCTCTGTCAACTGAAAGAGTGCCAGCCAGGATATCTTCTCTCTCTAAACACTCGAG  
 GATTGCTGCACAAACGTGAAAGCCTTTGGCAGGCTGTGGTGTATTTGAGGTCAGATG  
 TGACGACAGCAGGCCATCAAGGAGAAGAAGATAAGGATACCAATAGCTCCTCTCTCTCG  
 CAGTCCAGCTTTTAGCTGCTTCTGATGTTNTCTTCCCTGACTTTGTTGATNCANAGT  
 GCAGTGCANCAGTGAAATCCCTNCCGGTNTTGGGNTGCTGGGATGCANCCTCAGGGGAT  
 AAGAANTCGTGGGCAACTGCCGCCTTTCTGTACTTCTGCATTCGGGGGCAGGGCTTT  
 TGGCAGTGCTCTCCACGGAGCAACATGCTCAGGCCTGGCCCAACAN

**Restriction Sites:**

NotI-NotI

**ACCN:**

NM\_000135

**Insert Size:**

4640 bp

**OTI Disclaimer:**

Our molecular clone sequence data has been matched to the reference identifier above as a point of reference. Note that the complete sequence of our molecular clones may differ from the sequence published for this corresponding reference, e.g., by representing an alternative RNA splicing form or single nucleotide polymorphism (SNP).

**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).

<b>Reconstitution Method:</b>	<ol style="list-style-type: none"><li>1. Centrifuge at 5,000xg for 5min.</li><li>2. Carefully open the tube and add 100ul of sterile water to dissolve the DNA.</li><li>3. Close the tube and incubate for 10 minutes at room temperature.</li><li>4. Briefly vortex the tube and then do a quick spin (less than 5000xg) to concentrate the liquid at the bottom.</li><li>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.</li></ol>
<b>RefSeq:</b>	<u>NM_000135.2</u> , <u>NP_000126.2</u>
<b>RefSeq Size:</b>	5460 bp
<b>RefSeq ORF:</b>	4368 bp
<b>Locus ID:</b>	2175
<b>UniProt ID:</b>	<u>O15360</u>
<b>Cytogenetics:</b>	16q24.3
<b>Domains:</b>	Fanconia
<b>Protein Families:</b>	Druggable Genome
<b>Gene Summary:</b>	<p>The Fanconi anemia complementation group (FANC) currently includes FANCA, FANCB, FANCC, FANCD1 (also called BRCA2), FANCD2, FANCE, FANCF, FANCG, FANCI, FANCJ (also called BRIP1), FANCL, FANCM and FANCN (also called PALB2). The previously defined group FANCH is the same as FANCA. Fanconi anemia is a genetically heterogeneous recessive disorder characterized by cytogenetic instability, hypersensitivity to DNA crosslinking agents, increased chromosomal breakage, and defective DNA repair. The members of the Fanconi anemia complementation group do not share sequence similarity; they are related by their assembly into a common nuclear protein complex. This gene encodes the protein for complementation group A. Alternative splicing results in multiple transcript variants encoding different isoforms. Mutations in this gene are the most common cause of Fanconi anemia. [provided by RefSeq, Jul 2008]</p> <p>Transcript Variant: This variant (1) represents the longest transcript and encodes the longest isoform (a).</p>