

Product datasheet for SC334057

ABHD11 (NM 001301058) Human Untagged Clone

Product data:

Product Type: Expression Plasmids

Product Name: ABHD11 (NM_001301058) Human Untagged Clone

Tag: Tag Free Symbol: ABHD11

Synonyms: PP1226; WBSCR21

Mammalian Cell

Selection:

Neomycin

Vector:pCMV6-Entry (PS100001)E. coli Selection:Kanamycin (25 ug/mL)

Fully Sequenced ORF: >SC334057 representing NM_001301058.

Blue=Insert sequence Red=Cloning site Green=Tag(s)

GATCCGGTACCGAGGAGATCTGCCGCCGCGATCGCC

TACAAGGATGACGACGATAAGGTTTAAACGGCCGGC

Restriction Sites: Sgfl-Mlul



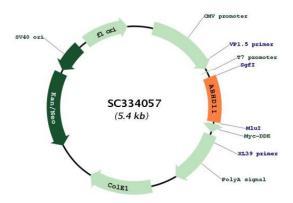
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Plasmid Map:



ACCN: NM_001301058

Insert Size: 480 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).

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: <u>NM 001301058.1</u>



ABHD11 (NM_001301058) Human Untagged Clone - SC334057

 RefSeq Size:
 1142 bp

 RefSeq ORF:
 480 bp

 Locus ID:
 83451

 UniProt ID:
 Q8NFV4

 Cytogenetics:
 7q11.23

 MW:
 17 kDa

Gene Summary: This gene encodes a protein containing an alpha/beta hydrolase fold domain. This gene is

deleted in Williams syndrome, a multisystem developmental disorder caused by the deletion

of contiguous genes at 7q11.23. [provided by RefSeq, Mar 2016]

Transcript Variant: This variant (9) lacks two exons in the central coding region, which results in a frameshift, compared to variant 1. The encoded isoform (9) has a shorter and distinct C-

terminus, compared to isoform 1.