

Product datasheet for RC236163

ABHD11 (NM_001301058) Human Tagged ORF Clone

Product data:

Product Type: Expression Plasmids

Product Name: ABHD11 (NM_001301058) Human Tagged ORF Clone

Tag: Myc-DDK
Symbol: ABHD11

Synonyms: PP1226; WBSCR21

Mammalian Cell Neomycin

Selection:

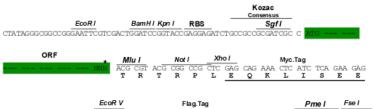
Vector:pCMV6-Entry (PS100001)E. coli Selection:Kanamycin (25 ug/mL)

Restriction Sites: Sgfl-Mlul

Cloning Scheme:

Cloning sites used for ORF Shuttling:





GAT CTG GCA GCA ANT GAT ATC CTG GAT TAC AGG GAT GAC GAC GAT AAG GTT TAA AACGGCCGGCCC

D L A A N D I L L D Y K D D D D K V stop

ACCN: NM_001301058

ORF Size: 477 bp



OriGene Technologies, Inc. 9620 Medical Center Drive, Ste 200

CN: techsupport@origene.cn

Rockville, MD 20850, US Phone: +1-888-267-4436 https://www.origene.com techsupport@origene.com EU: info-de@origene.com

^{*} The last codon before the Stop codon of the ORF



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OTI Disclaimer: 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

OTI Annotation: This clone was engineered to express the complete ORF with an expression tag. Expression

varies depending on the nature of the gene.

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, NP 001287987.1</u>

 RefSeq Size:
 1142 bp

 RefSeq ORF:
 453 bp

 Locus ID:
 83451

 UniProt ID:
 Q8NFV4

Cytogenetics: 7q11.23 **MW:** 17.5 kDa

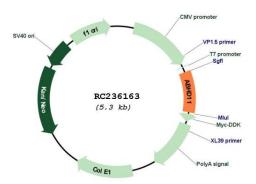
Gene Summary: This gene encodes a protein containing an alpha/beta hydrolase fold domain. This gene is

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

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



Product images:



Circular map for RC236163