

## Product datasheet for **SC206231**

### ABHD11 (NM\_148913) Human 3' UTR Clone

#### Product data:

**Product Type:** 3' UTR Clones  
**Product Name:** ABHD11 (NM\_148913) Human 3' UTR Clone  
**Symbol:** ABHD11  
**Synonyms:** PP1226; WBSCR21  
**Mammalian Cell Selection:** Neomycin  
**Vector:** pMirTarget (PS100062)  
**ACCN:** NM\_148913  
**Insert Size:** 495 bp  
**Insert Sequence:** >SC206231 3'UTR clone of NM\_148913  
The sequence shown below is from the reference sequence of NM\_148913. The complete sequence of this clone may contain minor differences, such as SNPs.  
**Blue**=Stop Codon **Red**=Cloning site

```
GGCAAGTTGGACGCCCGCAAGATCCGCGAGATTCTCATTAAAGCCAAGAAGGGCGGAAAGATCGCCGTG  
TAACAATTGGCAGAGCTCAGAATTCAAGCGATCGCC  
ATAGCTGCCATCCGAGGCTTCTGGTCTAAGAGTTGCTGGCAAGAAGATGGCCGGGCGTGGTGGCTCAT  
GCCTGTAATTCCAGCACTTTGGGAGGCTAAGGCGGGAGGATGACTTGAGGCCAGGAGTTGGAGACCAGC  
CTGGCCAACATGGTGAACCCCTGTCTCTACTAAAAATACAAAAATTAGCCTGGCGTGGTGGTGCACACC  
TGTAATCCCAGCTACTCTGGAGGCTGAGGCAGGAGAATCACTTGAACCCTGGAGGCAGAGGTTGCAATG  
AGCCGAGATCACACCACTACACTCCAGCCTAGGCAACAGAGCAAGACTCTGTCTCAAAAAACAAAAC  
AAAAAGGAGGCACAAAACCCAGGCTTCAAGTCTCTGCAGCCTGCTCCACATTTGGGCACAGAAGGACT  
CAGACAGGCACTGTGTGGGCACGAGTTTTACAGGGGTGGTCAGACCTCAGGCTTTAATGAATAAAGAC  
ACTACTCCCAA  
ACGCGTAAGCGGCCGCGCATCTAGATTCAAGAAAATGACCGACCAAGCGACGCCAACCTGCCATCA  
CGAGATTCGATTCCACCGCCGCTTCTATGAAAGG
```

**Restriction Sites:** SgfI-MluI  
**OTI Disclaimer:** Our molecular clone sequence data has been matched to the sequence identifier above as a point of reference. Note that the complete sequence of this clone is largely the same as the reference sequence but may contain minor differences, e.g., single nucleotide polymorphisms (SNPs).



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<b>Components:</b>	The cDNA clone is shipped in a 2-D bar-coded Matrix tube as 10 ug dried plasmid DNA. The package also includes 100 pmols of both the corresponding 5' and 3' vector primers in separate vials.
<b>RefSeq:</b>	<u><a href="#">NM_148913.4</a></u>
<b>Summary:</b>	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]
<b>Locus ID:</b>	83451
<b>MW:</b>	18.1