

## **Product datasheet for SC202460**

## AHI1 (NM 001134832) Human 3' UTR Clone

**Product data:** 

**Product Type:** 3' UTR Clones

Product Name: AHI1 (NM\_001134832) Human 3' UTR Clone

Symbol: AHI1

Synonyms: AHI-1; dJ71N10.1; JBTS3; ORF1

**Mammalian Cell** 

Selection:

Neomycin

**Vector:** pMirTarget (PS100062)

**ACCN:** NM\_001134832

**Insert Size:** 227 bp

Insert Sequence: >SC202460 3'UTR clone of NM\_001134832

The sequence shown below is from the reference sequence of NM\_001134832. The complete

sequence of this clone may contain minor differences, such as SNPs.

Blue=Stop Codon Red=Cloning site

GGCAAGTTGGACGCCCGCAAGATCCGCGAGATTCTCATTAAGGCCAAGAAGGGCGGAAAGATCGCCGTG

TAACAATTGGCAGAGCTCAGAATTCAAGCGATCGCC

GAATAAACAGTATGAGGTTA

CGAGATTTCGATTCCACCGCCGCCTTCTATGAAAGG

**Restriction Sites:** Sgfl-Mlul

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).

**Components:** 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.

**RefSeq:** <u>NM 001134832.2</u>



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## AHI1 (NM\_001134832) Human 3' UTR Clone - SC202460

**Summary:** This gene is apparently required for both cerebellar and cortical development in humans.

This gene mutations cause specific forms of Joubert syndrome-related disorders. Joubert syndrome (JS) is a recessively inherited developmental brain disorder with several identified causative chromosomal loci. Alternatively spliced transcript variants encoding different

isoforms have been identified. [provided by RefSeq, Oct 2008]

**Locus ID:** 54806

MW: 8.8