

Product datasheet for **RC226357L4V**

AHI1 (NM_001134830) Human Tagged ORF Clone Lentiviral Particle

Product data:

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|---------------------------|--|
| Product Type: | Lentiviral Particles |
| Product Name: | AHI1 (NM_001134830) Human Tagged ORF Clone Lentiviral Particle |
| Symbol: | AHI1 |
| Synonyms: | AHI-1; dj71N10.1; JBTS3; ORF1 |
| Mammalian Cell Selection: | Puromycin |
| Vector: | pLenti-C-mGFP-P2A-Puro (PS100093) |
| Tag: | mGFP |
| ACCN: | NM_001134830 |
| ORF Size: | 3588 bp |
| ORF Nucleotide Sequence: | The ORF insert of this clone is exactly the same as(RC226357). |
| 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. |
| RefSeq: | NM_001134830.1 , NP_001128302.1 |
| RefSeq ORF: | 3591 bp |
| Locus ID: | 54806 |
| UniProt ID: | Q8N157 |
| Cytogenetics: | 6q23.3 |
| MW: | 136.9 kDa |



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Gene 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]