

Product datasheet for **RC222991L4V**

AOC2 (NM_009590) Human Tagged ORF Clone Lentiviral Particle

Product data:

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| Product Type: | Lentiviral Particles |
| Product Name: | AOC2 (NM_009590) Human Tagged ORF Clone Lentiviral Particle |
| Symbol: | AOC2 |
| Synonyms: | DAO2; RAO; SSAO |
| Mammalian Cell Selection: | Puromycin |
| Vector: | pLenti-C-mGFP-P2A-Puro (PS100093) |
| Tag: | mGFP |
| ACCN: | NM_009590 |
| ORF Size: | 2268 bp |
| ORF Nucleotide Sequence: | The ORF insert of this clone is exactly the same as(RC222991). |
| 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_009590.2 |
| RefSeq Size: | 2681 bp |
| RefSeq ORF: | 2271 bp |
| Locus ID: | 314 |
| UniProt ID: | O75106 |
| Cytogenetics: | 17q21.31 |
| Protein Families: | Transmembrane |



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| Protein Pathways: | beta-Alanine metabolism, Glycine, serine and threonine metabolism, Metabolic pathways, Phenylalanine metabolism, Tyrosine metabolism |
| MW: | 83.5 kDa |
| Gene Summary: | Copper amine oxidases catalyze the oxidative conversion of amines to aldehydes and ammonia in the presence of copper and quinone cofactor. This gene shows high sequence similarity to copper amine oxidases from various species ranging from bacteria to mammals. The protein contains several conserved motifs including the active site of amine oxidases and the histidine residues that likely bind copper. It may be a critical modulator of signal transmission in retina, possibly by degrading the biogenic amines dopamine, histamine, and putrescine. This gene may be a candidate gene for hereditary ocular diseases. Alternate splicing results in multiple transcript variants. [provided by RefSeq, Jul 2008] |