

## Product datasheet for RC206273L3V

## OriGene Technologies, Inc.

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## HAAO (NM 012205) Human Tagged ORF Clone Lentiviral Particle

**Product data:** 

**Product Type:** Lentiviral Particles

**Product Name:** HAAO (NM 012205) Human Tagged ORF Clone Lentiviral Particle

Symbol:

3-HAO; h3HAO; HAO; VCRL1 Synonyms:

**Mammalian Cell** 

Selection:

ACCN:

Puromycin

Vector: pLenti-C-Myc-DDK-P2A-Puro (PS100092)

Tag: Myc-DDK NM 012205

**ORF Size:** 858 bp

**ORF Nucleotide** 

The ORF insert of this clone is exactly the same as(RC206273).

Sequence:

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 012205.1

RefSeq Size: 1301 bp RefSeq ORF: 861 bp Locus ID: 23498 **UniProt ID:** P46952 Cytogenetics: 2p21

**Protein Pathways:** Metabolic pathways, Tryptophan metabolism

MW: 32.6 kDa







## **Gene Summary:**

3-Hydroxyanthranilate 3,4-dioxygenase is a monomeric cytosolic protein belonging to the family of intramolecular dioxygenases containing nonheme ferrous iron. It is widely distributed in peripheral organs, such as liver and kidney, and is also present in low amounts in the central nervous system. HAAO catalyzes the synthesis of quinolinic acid (QUIN) from 3-hydroxyanthranilic acid. QUIN is an excitotoxin whose toxicity is mediated by its ability to activate glutamate N-methyl-D-aspartate receptors. Increased cerebral levels of QUIN may participate in the pathogenesis of neurologic and inflammatory disorders. HAAO has been suggested to play a role in disorders associated with altered tissue levels of QUIN. [provided by RefSeq, Jul 2008]