

Product datasheet for RC206868L2V

OriGene Technologies, Inc.

9620 Medical Center Drive, Ste 200 Rockville, MD 20850, US Phone: +1-888-267-4436 https://www.origene.com techsupport@origene.com EU: info-de@origene.com CN: techsupport@origene.cn

HHIP (NM_022475) Human Tagged ORF Clone Lentiviral Particle

Product data:

Product Type: Lentiviral Particles

Product Name: HHIP (NM_022475) Human Tagged ORF Clone Lentiviral Particle

Symbol: HHIP
Synonyms: HIP

Mammalian Cell None

Selection:

Vector: pLenti-C-mGFP (PS100071)

Tag: mGFP

ACCN: NM_022475 **ORF Size:** 2100 bp

ORF Nucleotide

The ORF insert of this clone is exactly the same as(RC206868).

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.

RefSeg: NM 022475.1

 RefSeq Size:
 3555 bp

 RefSeq ORF:
 2103 bp

 Locus ID:
 64399

 UniProt ID:
 Q96QV1

 Cytogenetics:
 4q31.21

Domains: EGF

Protein Families: Secreted Protein





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Protein Pathways: Basal cell carcinoma, Hedgehog signaling pathway, Pathways in cancer

MW: 78.9 kDa

Gene Summary: This gene encodes a member of the hedgehog-interacting protein (HHIP) family. The

hedgehog (HH) proteins are evolutionarily conserved protein, which are important morphogens for a wide range of developmental processes, including anteroposterior patterns of limbs and regulation of left-right asymmetry in embryonic development. Multiple cell-surface receptors are responsible for transducing and/or regulating HH signals. The HHIP encoded by this gene is a highly conserved, vertebrate-specific inhibitor of HH signaling. It interacts with all three HH family members, SHH, IHH and DHH. Two single nucleotide

obstructive pulmonary disease (COPD). A single nucleotide polymorphism in this gene is also

polymorphisms (SNPs) near this gene are significantly associated with risk of chronic

strongly associated with human height.[provided by RefSeq, Feb 2011]