

Product datasheet for RC219592L3V

OriGene Technologies, Inc.

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Axin 1 (AXIN1) (NM_003502) Human Tagged ORF Clone Lentiviral Particle

Product data:

Product Type: Lentiviral Particles

Product Name: Axin 1 (AXIN1) (NM_003502) Human Tagged ORF Clone Lentiviral Particle

Symbol: Axin 1

Synonyms: AXIN; PPP1R49

Mammalian Cell

Selection:

ACCN:

Puromycin

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

NM 003502

Tag: Myc-DDK

ORF Size: 2586 bp

ORF Nucleotide

The ORF insert of this clone is exactly the same as(RC219592).

OTI Disclaimer:

Sequence:

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 003502.2

 RefSeq Size:
 3477 bp

 RefSeq ORF:
 2589 bp

 Locus ID:
 8312

 UniProt ID:
 015169

Cytogenetics: 16p13.3

Protein Families: Druggable Genome, ES Cell Differentiation/IPS, Stem cell relevant signaling - Wnt Signaling

pathway





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Protein Pathways: Basal cell carcinoma, Colorectal cancer, Endometrial cancer, Pathways in cancer, Wnt

signaling pathway

MW: 95.5 kDa

Gene Summary: This gene encodes a cytoplasmic protein which contains a regulation of G-protein signaling

(RGS) domain and a dishevelled and axin (DIX) domain. The encoded protein interacts with adenomatosis polyposis coli, catenin beta-1, glycogen synthase kinase 3 beta, protein phosphate 2, and itself. This protein functions as a negative regulator of the wingless-type MMTV integration site family, member 1 (WNT) signaling pathway and can induce apoptosis. The crystal structure of a portion of this protein, alone and in a complex with other proteins,

has been resolved. Mutations in this gene have been associated with hepatocellular

carcinoma, hepatoblastomas, ovarian endometriod adenocarcinomas, and

medullablastomas. Alternative splicing results in multiple transcript variants. [provided by

RefSeq, Jan 2016]