

Product datasheet for RC200902L3V

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

RhoGDI (ARHGDIA) (NM 004309) Human Tagged ORF Clone Lentiviral Particle

Product data:

Product Type: Lentiviral Particles

Product Name: RhoGDI (ARHGDIA) (NM_004309) Human Tagged ORF Clone Lentiviral Particle

Symbol:

GDIA1; HEL-S-47e; NPHS8; RHOGDI; RHOGDI-1 Synonyms:

Mammalian Cell

Selection:

Puromycin

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

612 bp

Tag: Myc-DDK

NM 004309 ACCN:

OTI Disclaimer:

ORF Nucleotide

Sequence:

ORF Size:

The ORF insert of this clone is exactly the same as(RC200902).

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 004309.3

RefSeq Size: 1920 bp RefSeq ORF: 615 bp Locus ID: 396 **UniProt ID:** P52565

Cytogenetics: 17q25.3 **Domains:** Rho_GDI

Protein Families: Druggable Genome





Protein Pathways: Neurotrophin signaling pathway

MW: 23.2 kDa

Gene Summary: This gene encodes a protein that plays a key role in the regulation of signaling through Rho

GTPases. The encoded protein inhibits the disassociation of Rho family members from GDP (guanine diphosphate), thereby maintaining these factors in an inactive state. Activity of this protein is important in a variety of cellular processes, and expression of this gene may be altered in tumors. Mutations in this gene have been found in individuals with nephrotic syndrome, type 8. Alternate splicing results in multiple transcript variants. [provided by

RefSeq, Jul 2014]