

OriGene Technologies, Inc.

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Product datasheet for RC206679L1V

ADA (NM_000022) Human Tagged ORF Clone Lentiviral Particle

Product data:

Product Type:	Lentiviral Particles
Product Name:	ADA (NM_000022) Human Tagged ORF Clone Lentiviral Particle
Symbol:	ADA
Synonyms:	ADA1
Mammalian Cell Selection:	None
Vector:	pLenti-C-Myc-DDK (PS100064)
Tag:	Myc-DDK
ACCN:	NM_000022
ORF Size:	1089 bp
ORF Nucleotide Sequence:	The ORF insert of this clone is exactly the same as(RC206679).
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. <u>More info</u>
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:	<u>NM 000022.2</u>
RefSeq Size:	1566 bp
RefSeq ORF:	1092 bp
Locus ID:	100
UniProt ID:	<u>P00813</u>
Cytogenetics:	20q13.12
Domains:	A_deaminase
Protein Families:	Druggable Genome



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GRIGENE ADA (NM_000022) Human Tagged ORF Clone Lentiviral Particle – RC206679L1V	
Protein Pathways:	Metabolic pathways, Primary immunodeficiency, Purine metabolism
MW:	40.8 kDa
Gene Summary:	This gene encodes an enzyme that catalyzes the hydrolysis of adenosine to inosine in the purine catabolic pathway. Various mutations have been described for this gene and have been linked to human diseases related to impaired immune function such as severe combined immunodeficiency disease (SCID) which is the result of a deficiency in the ADA enzyme. In ADA-deficient individuals there is a marked depletion of T, B, and NK lymphocytes, and consequently, a lack of both humoral and cellular immunity. Conversely, elevated levels of this enzyme are associated with congenital hemolytic anemia. [provided by RefSeq, Sep 2019]

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