

## 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

## Product datasheet for RC204031L2V

## NC2 alpha (DRAP1) (NM\_006442) Human Tagged ORF Clone Lentiviral Particle

## **Product data:**

Product Type:	Lentiviral Particles
Product Name:	NC2 alpha (DRAP1) (NM_006442) Human Tagged ORF Clone Lentiviral Particle
Symbol:	NC2 alpha
Synonyms:	NC2-alpha
Mammalian Cell Selection:	None
Vector:	pLenti-C-mGFP (PS100071)
Tag:	mGFP
ACCN:	NM_006442
ORF Size:	615 bp
ORF Nucleotide Sequence:	The ORF insert of this clone is exactly the same as(RC204031).
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 006442.2</u>
RefSeq Size:	1022 bp
RefSeq ORF:	618 bp
Locus ID:	10589
UniProt ID:	<u>Q14919</u>
Cytogenetics:	11q13.1
Protein Families:	Transcription Factors
MW:	22.3 kDa



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Gene Summary:	Transcriptional repression is a general mechanism for regulating transcriptional initiation in
	organisms ranging from yeast to humans. Accurate initiation of transcription from eukaryotic
	protein-encoding genes requires the assembly of a large multiprotein complex consisting of
	RNA polymerase II and general transcription factors such as TFIIA, TFIIB, and TFIID. DR1 is a
	repressor that interacts with the TATA-binding protein (TBP) of TFIID and prevents the

stabilize the TBP-DR1-DNA complex. [provided by RefSeq, Jul 2008]

formation of an active transcription complex by precluding the entry of TFIIA and/or TFIIB into the preinitiation complex. The protein encoded by this gene is a corepressor of

transcription that interacts with DR1 to enhance DR1-mediated repression. The interaction between this corepressor and DR1 is required for corepressor function and appears to

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