

## Product datasheet for **RC220558L1V**

### Mineralocorticoid Receptor (NR3C2) (NM\_000901) Human Tagged ORF Clone Lentiviral Particle

#### Product data:

Product Type:	Lentiviral Particles
Product Name:	Mineralocorticoid Receptor (NR3C2) (NM_000901) Human Tagged ORF Clone Lentiviral Particle
Symbol:	Mineralocorticoid Receptor
Synonyms:	MCR; MLR; MR; NR3C2VIT
Mammalian Cell Selection:	None
Vector:	pLenti-C-Myc-DDK (PS100064)
Tag:	Myc-DDK
ACCN:	NM_000901
ORF Size:	2952 bp
ORF Nucleotide Sequence:	The ORF insert of this clone is exactly the same as(RC220558).
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. <a href="#">More info</a>
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:	<a href="#">NM_000901.1</a>
RefSeq Size:	5749 bp
RefSeq ORF:	2955 bp
Locus ID:	4306
UniProt ID:	<a href="#">P08235</a>
Cytogenetics:	4q31.23
Domains:	HOLI, zf-C4



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**Protein Families:** Druggable Genome, Nuclear Hormone Receptor, Transcription Factors

**MW:** 106.9 kDa

**Gene Summary:** This gene encodes the mineralocorticoid receptor, which mediates aldosterone actions on salt and water balance within restricted target cells. The protein functions as a ligand-dependent transcription factor that binds to mineralocorticoid response elements in order to transactivate target genes. Mutations in this gene cause autosomal dominant pseudohypoaldosteronism type I, a disorder characterized by urinary salt wasting. Defects in this gene are also associated with early onset hypertension with severe exacerbation in pregnancy. Alternative splicing results in multiple transcript variants. [provided by RefSeq, Oct 2009]