

Product datasheet for RC214392L1V

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

ERCC8 (NM_000082) Human Tagged ORF Clone Lentiviral Particle

Product data:

Product Type: Lentiviral Particles

Product Name: ERCC8 (NM_000082) Human Tagged ORF Clone Lentiviral Particle

Symbol: ERCC8

Synonyms: CKN1; CSA; UVSS2

Mammalian Cell

Selection:

None

Vector: pLenti-C-Myc-DDK (PS100064)

Tag: Myc-DDK
ACCN: NM 000082

ORF Size: 1188 bp

ORF Nucleotide

The ORF insert of this clone is exactly the same as(RC214392).

Sequence:

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. 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 000082.2

RefSeq Size: 2031 bp
RefSeq ORF: 1191 bp
Locus ID: 1161
UniProt ID: Q13216

Cytogenetics: 5q12.1

Domains: WD40

Protein Families: Druggable Genome, Transcription Factors





ERCC8 (NM_000082) Human Tagged ORF Clone Lentiviral Particle - RC214392L1V

Protein Pathways: Nucleotide excision repair, Ubiquitin mediated proteolysis

MW: 43.9 kDa

Gene Summary: This gene encodes a WD repeat protein, which interacts with Cockayne syndrome type B

(CSB) protein and with p44 protein, a subunit of the RNA polymerase II transcription factor IIH. Mutations in this gene have been identified in patients with hereditary disease Cockayne syndrome (CS). CS cells are abnormally sensitive to ultraviolet radiation and are defective in the repair of transcriptionally active genes. Several transcript variants encoding different

isoforms have been found for this gene. [provided by RefSeq, Mar 2014]