

Product datasheet for SC334770

ERCC8 (NM 001290285) Human Untagged Clone

Product data:

Product Type: Expression Plasmids

Product Name: ERCC8 (NM_001290285) Human Untagged Clone

Tag: Tag Free Symbol: ERCC8

Synonyms: CKN1; CSA; UVSS2

Mammalian Cell

Selection:

Neomycin

Vector:pCMV6-Entry (PS100001)E. coli Selection:Kanamycin (25 ug/mL)

Fully Sequenced ORF: >NCBI ORF sequence for NM_001290285, the custom clone sequence may differ by one or

more nucleotides

Restriction Sites: Sgfl-Mlul

ACCN: NM 001290285

OTI Disclaimer: Our molecular clone sequence data has been matched to the reference identifier above as a

point of reference. Note that the complete sequence of our molecular clones may differ from the sequence published for this corresponding reference, e.g., by representing an alternative

RNA splicing form or single nucleotide polymorphism (SNP).



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ERCC8 (NM_001290285) Human Untagged Clone - SC334770

Components: The ORF clone is ion-exchange column purified and shipped in a 2D barcoded Matrix tube

containing 10ug of transfection-ready, dried plasmid DNA (reconstitute with 100 ul of water).

Reconstitution Method: 1. Centrifuge at 5,000xg for 5min.

2. Carefully open the tube and add 100ul of sterile water to dissolve the DNA.

3. Close the tube and incubate for 10 minutes at room temperature.

4. Briefly vortex the tube and then do a quick spin (less than 5000xg) to concentrate the liquid

at the bottom.

5. Store the suspended plasmid at -20°C. The DNA is stable for at least one year from date of

shipping when stored at -20°C.

RefSeq: <u>NM 001290285.1</u>, <u>NP 001277214.1</u>

RefSeq Size: 1962 bp
RefSeq ORF: 732 bp
Locus ID: 1161
Cytogenetics: 5q12.1

Protein Families: Druggable Genome, Transcription Factors

Protein Pathways: Nucleotide excision repair, Ubiquitin mediated proteolysis

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]

Transcript Variant: This variant (4) lacks an alternate internal exon and uses a downstream AUG start codon compared to variant 1. The resulting isoform (4) has a shorter and distinct N-

terminus compared to isoform 1.