

Product datasheet for SC203306

ATP13A2 (NM 022089) Human 3' UTR Clone

Product data:

Product Type: 3' UTR Clones

Product Name: ATP13A2 (NM_022089) Human 3' UTR Clone

Symbol: ATP13A2

CLN12; HSA9947; KRPPD; PARK9; SPG78 Synonyms:

Mammalian Cell

Selection:

Neomycin

pMirTarget (PS100062) Vector:

ACCN: NM 022089

Insert Size: 293 bp

Insert Sequence: >SC203306 3'UTR clone of NM_022089

The sequence shown below is from the reference sequence of NM_022089. The complete

sequence of this clone may contain minor differences, such as SNPs.

Blue=Stop Codon Red=Cloning site

GGCAAGTTGGACGCCCGCAAGATCCGCGAGATTCTCATTAAGGCCAAGAAGGGCGGAAAGATCGCCGTG

TAACAATTGGCAGAGCTCAGAATTCAAGCGATCGCC

CCGCCGCTGCCCGCCCGCCCCTGAGGTAGTGCAGGCCCACGGGCACCCCAGACACTGGAACTCCCTGC CTCTGAGCCACCAACTGGACCCCTCTCCAGCAACACCACCGCCACCACCTCCCACATCCCTGAGGTTGG CGACTGTCTACACTCCTCCCCCGAGACCACCCCCACCCTGGGGAAGCGTTGACTACTGTCCCCTACCTT GGACCATCCCGCGTAGGGGTGGCAGCCCCCAGCTCCCCTCAGTGCTGCTGTCAGTGTAGCAAATAAAGT

CATGATATTTTCCTGGC

CGAGATTTCGATTCCACCGCCGCCTTCTATGAAAGG

Restriction Sites: Sgfl-Mlul

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

point of reference. Note that the complete sequence of this clone is largely the same as the

reference sequence but may contain minor differences, e.g., single nucleotide

polymorphisms (SNPs).

Components: The cDNA clone is shipped in a 2-D bar-coded Matrix tube as 10 ug dried plasmid DNA. The

package also includes 100 pmols of both the corresponding 5' and 3' vector primers in

separate vials.

NM 022089.4 RefSeq:



OriGene Technologies, Inc. 9620 Medical Center Drive, Ste 200

CN: techsupport@origene.cn

Rockville, MD 20850, US Phone: +1-888-267-4436 https://www.origene.com techsupport@origene.com EU: info-de@origene.com



ATP13A2 (NM_022089) Human 3' UTR Clone - SC203306

Summary: This gene encodes a member of the P5 subfamily of ATPases which transports inorganic

cations as well as other substrates. Mutations in this gene are associated with Kufor-Rakeb syndrome (KRS), also referred to as Parkinson disease 9. Multiple transcript variants encoding

different isoforms have been found for this gene.[provided by RefSeq, Nov 2008]

Locus ID: 23400

MW: 10.6