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Datasheet for ABIN4927980

Human KRTAP19-2 ORF Clone in Mammalian Expression Vector (DYKDDDDK Tag)

Overview		
Quantity:	10 μg	
Gene:	KRTAP19-2	
Species:	Human	
Fusion tag:	DYKDDDDK Tag	
Insert:	ORF	
Vector:	Mammalian Expression Vector	
Application:	Protein Expression (PExp)	
Product Details		
Purpose:	Expression/transfection ready cDNA ORF clone of Human KRTAP19-2 with C terminal DYKDDDDK tag is ideal for express proteins in E.coli & mammalian cells.	
Brand:	GenEZ™	
Insert Length:	159 bp	
Vector Backbone:	pcDNA3.1+C-(K)-DYK	
Promoter:	CMV Promoter	
Selectable Marker:	Neomycin	
Bacterial Resistance:	Ampicillin	
Expression Type:	Transient, Stable	
Sequence:	ATGTGCTATG GCTACGGCTG TGGATGTGGC AGCTTCTGCA GACTGGGCTA TGGCTGCGGC TATGAAGGAT GCAGATATGG TTGTGGCCAC AGAGGCTGTG GAGATGGCTG CTGCTGCCCA TCATGCTACA GAAGATATAG ATTCACTGGC TTCTACTAA	
Specificity:	ORF Insert Method: CloneEZ® Seamless cloning technology, recombination-based cloning	

Product Details

technology
Gene cDNA ORF clone sequences were retrieved from the NCBI Reference Sequence Database
(RefSeq). These sequences represent the protein coding region of the gene cDNA ORF which is
encoded by the open reading frame (ORF) sequence.
Forward primer: 5'-TAATACGACTCACTATAGGG-3'
Reverse primer: 5'-CCTCGACTGTGCCTTCTA-3'
End-sequenced
The GenEZ ORF clone is delivered as 10 μg of lyophilized plasmid DNA in a vial.
KRTAP19-2
KRTAP19-2 (KRTAP19-2 Products)
337969
NM_181608
For Research Use only
Lyophilized
RT/-20 °C
Keep the vial sealed and store at -20°C for long-term storage.
Before use, centrifuge the vial at 6,000 g x g for 1 minute at 4°C.
 Open the lid and add 100 µl (or other volume depending on your desired final concentration) of distilled water (or TE buffer) to dissolve the DNA.
 If necessary, heat the solution at 50°C for 15 minutes to dissolve the DNA.
Close the lid and vortex the vial for 1 minute.
 Aliquot the dissolved plasmid DNA and store in small aliquots at -20°C.
12 months

Publications

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Product	CITEC	ın.

Johnson, Drugan, Miller, Evans: "38" in: , Vol. 1363, Issue Nucleic acids research, pp. 28-39, (1991)