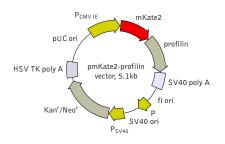


# pmKate2-profilin vector

The vector sequence has been compiled using the information from sequence databases, published literature, and other sources, together with partial sequences obtained by Evrogen. This vector has not been completely sequenced.



For vector sequence, please visit our Web site at http://www.evrogen.com/products/vectors.shtml

## **Location of features**

P<sub>CMV IE</sub>: 1-589 Enhancer region: 59-465 TATA box: 554-560 Transcription start point: 583 mKate2-profilin fusion: 613-1764 mKate2: 613-1314 Start codon (ATG): 613-615 Last amino acid in mKate2: 1312-1314 Profilin: 1345-1764 Stop codon: 1765-1767 SV40 early mRNA polyadenylation signal Polyadenylation signals: 1928-1933 & 1957-1962 mRNA 3' ends: 1966 & 1978 f1 single-strand DNA origin: 2025-2480 Bacterial promoter for expression of Kan<sup>r</sup> gene -35 region: 2542-2547; -10 region: 2565-2570 Transcription start point: 2577 SV40 origin of replication: 2821-2956 SV40 early promoter Enhancer (72-bp tandem repeats): 2654-2725 & 2726-2797 21-bp repeats: 2801-2821, 2822-2842 & 2844-2864 Early promoter element: 2877-2883 Major transcription start points: 2873, 2911, 2917 & 2922 Kanamycin/neomycin resistance gene Neomycin phosphotransferase coding sequences: Start codon (ATG): 3005-3007: Stop codon: 3797-3799

G->A mutation to remove Pst I site: 3187 C->A (Arg to Ser) mutation to remove BssH II site: 3533 Herpes simplex virus (HSV) thymidine kinase (TK) polyadenylation signal

Polyadenylation signals: 4035-4040 & 4048-4053 pUC plasmid replication origin: 4384-5027

Product	Cat.#	Size	
pmKate2-profilin vector	FP320	20 $\mu$ g	
Vector type	mammalian expr	ession vector	
Reporter	mKate2		
Reporter codon usage	mammalian		
Promoter for mKate2	P <sub>CMV IE</sub>		
Host cells	mammalian		
Selection	prokaryotic - kanamycin eukaryotic - neomycin (G418)		
Replication	prokaryotic - pUC ori eukaryotic - SV40 ori		
Use	far-red fluorescent labeling of profilin		

# Vector description

pmKate2-profilin is a mammalian expression vector encoding mKate2-profilin fusion protein. The vector can be used for fluorescent labeling of profilin in living cells.

mKate2 codon usage is optimized for high expression in mammalian cells (humanized) [Haas et al. 1996]. Mouse profilin is fused to the mKate2 C-terminus. To increase mRNA translation efficiency, Kozak consensus translation initiation site is generated upstream of the mKate2-profilin coding sequence [Kozak 1987].

pmKate2-profilin vector can be used as a source of mKate2-profilin hybrid sequence. The vector backbone contains unique restriction sites that permit its excision and further insertion into expression vector of choice. **Note:** The plasmid DNA was isolated from dam<sup>+</sup>-methylated *E.coli*. Therefore some restriction sites are blocked by methylation. If you wish to digest the vector using such sites you will need to transform the vector into a dam<sup>-</sup> host and make fresh DNA.

The vector backbone contains immediate early promoter of cytomegalovirus ( $P_{CMV \, IE}$ ) for protein expression, SV40 origin for replication in mammalian cells expressing SV40 T-antigen, pUC origin of replication for propagation in *E. coli*, and f1 origin for single-stranded DNA production. SV40 polyadenylation signals (SV40 poly A) direct proper processing of the 3'-end of the reporter mRNA.

SV40 early promoter ( $P_{SV40}$ ) provides neomycin resistance gene (Neo<sup>r</sup>) expression to select stably transfected eukaryotic cells using G418. Bacterial promoter (P) provides kanamycin resistance gene expression (Kan<sup>r</sup>) in *E. coli*. Kan<sup>r</sup>/Neo<sup>r</sup> gene is linked with herpes simplex virus (HSV) thymidine kinase (TK) polyadenylation signals.

#### **Expression in mammalian cells**

pmKate2-profilin vector can be transfected into mammalian cells by any known transfection method. CMV promoter provides strong, constitutive expression of the mKate2-profilin fusion in eukaryotic cells. If required, stable transformants can be selected using G418 [Gorman 1985].

#### Propagation in E. coli

Suitable host strains for propagation in *E. coli* include DH5alpha, HB101, XL1-Blue, and other general purpose strains. Plasmid incompatibility group is pMB1/ColE1. The vector confers resistance to kanamycin (30  $\mu$ g/ml) to *E. coli* hosts. Copy number in *E. coli* is about 500.

#### References

Gorman, C. (1985). "High efficiency gene transfer into mammalian cells." In: DNA cloning: A Practical Approach, Vol. II. Ed. by Glover. (IRL Press, Oxford, U.K.) Pp. 143–190.

Haas, J. et al. (1996) "Codon usage limitation in the expression of HIV-1 envelope glycoprotein." Curr Biol, 6 (3): 315–324 / pmid: 8805248

Kozak, M. (1987) "An analysis of 5'-noncoding sequences from 699 vertebrate messenger RNAs." Nucleic Acids Res, 15 (20): 8125–8148 / pmid: 3313277

### Notice to Purchaser:

mKate2-related materials (also referred to as "Products") are intended for research use only.

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MSDS information is available at http://www.evrogen.com/MSDS.shtml

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