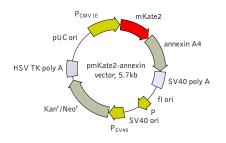


pmKate2-annexin vector

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_{CMV IE}: 1-589 Enhancer region: 59-465 TATA box: 554-560 Transcription start point: 583 mKate2-annexin fusion: 603-2313 mKate2: 603-1314 Start codon (ATG): 603-605 Last amino acid in mKate2: 1312-1314 Annexin A4: 1351-2313 Stop codon: 2314-2316 SV40 early mRNA polyadenylation signal Polyadenylation signals: 2477-2482 & 2506-2511 mRNA 3' ends: 2515 & 2527 f1 single-strand DNA origin: 2574-3029 Bacterial promoter for expression of Kan^r gene -35 region: 3091-3096; -10 region: 3114-3119 Transcription start point: 3126 SV40 origin of replication: 3370-3505 SV40 early promoter Enhancer (72-bp tandem repeats): 3203-3274 & 3275-3346 21-bp repeats: 3350-3370, 3371-3391 & 3393-3413 Early promoter element: 3426-3432 Major transcription start points: 3422, 3460, 3466 & 3471

Kanamycin/neomycin resistance gene

Neomycin phosphotransferase coding sequences: Start codon (ATG): 3554-3556; Stop codon: 4346-4348 G->A mutation to remove Pst I site: 3736 C->A (Arg to Ser) mutation to remove BssH II site: 4082

Herpes simplex virus (HSV) thymidine kinase (TK) polyadenylation signal Polyadenylation signals: 4584-4589 & 4597-4602

pUC plasmid replication origin: 4933-5576

Product	Cat.#	Size	
pmKate2-annexin vector	FP321	20 μ g	
Vector type	mammalian expression vector		
Reporter	mKate2		
Reporter codon usage	mammalian		
Promoter for mKate2	P _{CMV IE}		
Host cells	mammalian		
Selection	prokaryotic - kanamycin eukaryotic - neomycin (G418)		
Replication	prokaryotic - pUC ori eukaryotic - SV40 ori		
Use	far-red fluorescent labeling of annexin A4		

Vector description

pmKate2-annexin is a mammalian expression vector encoding mKate2-annexin fusion protein. The vector can be used for fluorescent labeling of annexin A4 in living cells.

mKate2 codon usage is optimized for high expression in mammalian cells (humanized) [Haas et al. 1996]. Human annexin A4 is fused to the mKate2 C-terminus. To increase mRNA translation efficiency, Kozak consensus translation initiation site is generated upstream of the mKate2-annexin coding sequence [Kozak 1987].

pmKate2-annexin vector can be used as a source of mKate2-annexin 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⁺-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⁻ 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^r) expression to select stably transfected eukaryotic cells using G418. Bacterial promoter (P) provides kanamycin resistance gene expression (Kan^r) in *E. coli*. Kan^r/Neo^r gene is linked with herpes simplex virus (HSV) thymidine kinase (TK) polyadenylation signals.

Expression in mammalian cells

pmKate2-annexin vector can be transfected into mammalian cells by any known transfection method. CMV promoter provides strong, constitutive expression of the mKate2-annexin 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 μ 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

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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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