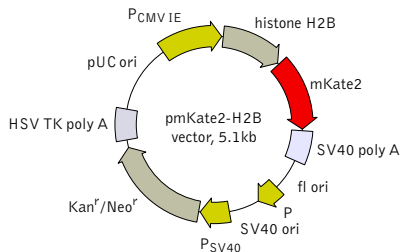


pmKate2-H2B vector

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/support/vector-info.shtml>

Location of features

PCMV IE: 1-589
 Enhancer region: 59-465
 TATA box: 554-560
 Transcription start point: 583
 Histone H2B protein: 657-1034
 TagRFP: 1053-1751
 SV40 early mRNA polyadenylation signal
 Polyadenylation signals: 1904-1909 & 1933-1938
 mRNA 3' ends: 1942 & 1954
 f1 single-strand DNA origin: 2001-2456
 Bacterial promoter for expression of Kan^r gene
 -35 region: 2518-2523; -10 region: 2541-2546
 Transcription start point: 2553
 SV40 origin of replication: 2797-2932
 SV40 early promoter
 Enhancer (72-bp tandem repeats): 2630-2701 & 2702-2773
 21-bp repeats: 2777-2797, 2798-2818 & 2820-2840
 Early promoter element: 2853-2859
 Major transcription start points: 2849, 2887, 2893 & 2898
 Kanamycin/neomycin resistance gene
 Neomycin phosphotransferase coding sequences:
 Start codon (ATG): 2981-2983; Stop codon: 3773-3775
 G->A mutation to remove Pst I site: 3163
 C->A (Arg to Ser) mutation to remove BssH II site: 3509
 Herpes simplex virus (HSV) thymidine kinase (TK) polyadenylation signal
 Polyadenylation signals: 4011-4016 & 4024-4029
 pUC plasmid replication origin: 4360-5003

Product	Cat.#	Size
pmKate2-H2B vector	FP311	20 µg

The price does not include delivery. The price varies in different countries. Please contact your local distributor for exact prices and delivery information.

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 histone H2B

Vector description

pmKate2-H2B is a mammalian expression vector encoding mKate2-H2B fusion protein. The vector can be used for fluorescent labeling of histone H2B in living cells.

mKate2 codon usage is optimized for high expression in mammalian cells, i.e. humanized (Haas et al. 1996). Human histone H2B is fused to the mKate2 N-terminus.

pmKate2-H2B can be used as a source of mKate2-H2B 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 also 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-H2B can be transfected into mammalian cells by any known transfection method. CMV promoter provides strong, constitutive expression of the mKate2-H2B 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 µg/ml) to *E. coli* hosts. Copy number in *E. coli* is about 500.

References

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

Notice to Purchaser:

Evrogen Fluorescent Protein Products (the Products) are intended for research use only. The Products are covered by U.S. Pat. 7,417,131 and other Evrogen Patents and/or Patent applications pending. By use of these Products, you accept the terms and conditions of the applicable Limited Use Label License.

The CMV promoter is covered under U.S. Patents 5,168,062 and 5,385,839, and its use is permitted for research purposes only. Any other use of the CMV promoter requires a license from the University of Iowa Research Foundation, 214 Technology Innovation Center, Iowa City, IA 52242.

MATERIAL SAFETY DATA SHEET INFORMATION: To the best of our knowledge, these products do not require a Material Safety Data Sheet. However, all the properties of these products (and, if applicable, each of their components) have not been thoroughly investigated. Therefore, we recommend that you use gloves and eye protection, and wear a laboratory coat when working with these products.