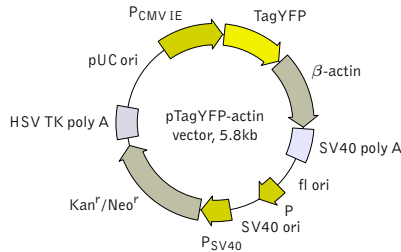


pTagYFP-actin vector

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
 Kozak consensus translation initiation site: 606-616
 TagYFP
 Start codon (ATG): 613-615
 Last amino acid in TagYFP: 1327-1329
 Beta-Actin: 1351-2478
 Stop codon: 2476-2478
 SV40 early mRNA polyadenylation signal
 Polyadenylation signals: 2639-2644 & 2668-2673
 mRNA 3' ends: 2677 & 2689
 f1 single-strand DNA origin: 2736-3191
 Eukaryotic promoter for expression of Kan^r gene
 -35 region: 3253-3258; -10 region: 3276-3281
 Transcription start point: 3288
 SV40 origin of replication: 3532-3667
 SV40 early promoter
 Enhancer (72-bp tandem repeats): 3365-3436 & 3437-3508
 21-bp repeats: 3512-3532, 3533-3553 & 3555-3575
 Early promoter element: 3588-3594
 Major transcription start points: 3584, 3622, 3628 & 3633
 Kanamycin/neomycin resistance gene
 Neomycin phosphotransferase coding sequences:
 Start codon (ATG): 3716-3718; Stop codon: 4508-4510
 G->A mutation to remove Pst I site: 3898
 C->A (Arg to Ser) mutation to remove BssH II site: 4244
 Herpes simplex virus (HSV) thymidine kinase (TK) polyadenylation signal
 Polyadenylation signals: 4746-4751 & 4759-4764
 pUC plasmid replication origin: 5095-5738

Product	Cat.#	Size
pTagYFP-actin vector	FP134	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	TagYFP	
Reporter codon usage	mammalian	
Promoter for TagYFP	P _{CMV IE}	
Host cells	mammalian	
Selection	prokaryotic - kanamycin eukaryotic - neomycin (G418)	
Replication	prokaryotic - pUC ori eukaryotic - SV40 ori	
Use	yellow fluorescent labeling of β-actin filaments	

Vector description

pTagYFP-actin is a mammalian expression vector encoding TagYFP-actin fusion protein. The vector can be used for fluorescent labeling of β-actin in living cells.

TagYFP codon usage is optimized for high expression in mammalian cells, i.e. humanized [Haas et al. 1996]. Human cytoplasmic β-actin is fused to the TagYFP C-terminus. To increase mRNA translation efficiency, Kozak consensus translation initiation site is generated upstream of TagYFP-actin coding sequence [Kozak 1987].

pTagYFP-actin can be used as a source of TagYFP-actin 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

pTagYFP-actin can be transfected into mammalian cells by any known transfection method. CMV promoter provides strong, constitutive expression of the TagYFP-actin 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
- Kozak (1987) "An analysis of 5'-noncoding sequences from 699 vertebrate messenger RNAs." *Nucleic Acids Res*, 15 (20): 8125-8148 / pmid: 3313277

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.