

pTagFP635-Cx32 vector

Cat# FP383

Vector description

pTagFP635-Cx32 is a mammalian expression vector encoding TagFP635-Cx32 fusion protein. The vector can be used for fluorescent labeling of connexin 32 in living cells.

TagFP635 codon usage is optimized for high expression in mammalian cells, i.e. humanized (Haas *et al.*, 1996). Human connexin 32 is fused to the TagFP635 N-terminus.

pTagFP635-Cx32 can be used as a source of TagFP635-Cx32 hybrid sequence. The vector backbone contains unique restriction sites that permit its excision and further insertion into an expression vector of choice.

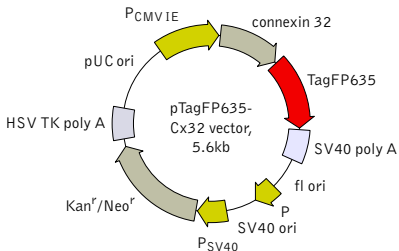
Note: The plasmid DNA was isolated from dam^+ -methylated *E. coli*. Therefore some restriction sites are blocked by methylation. If you wish to digest the vector using such sites you will need to transform the vector into a dam^- host and make fresh DNA.

The vector backbone also contains an immediate early promoter of cytomegalovirus (P_{CMVIE}) 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.

Vector map

For vector sequence, please visit our Web site at <http://www.evrogen.com/support/vector-info.shtml>



Expression in mammalian cells

pTagFP635-Cx32 can be transfected into mammalian cells by any known transfection method. CMV promoter provides strong, constitutive expression of the TagFP635-Cx32 fusion in eukaryotic cells. If required, stable transformants can be selected using G418 [Gorman, 1985].

Location of features

PCMV IE: 1-589
Enhancer region: 59-465
TATA box: 554-560
Transcription start point: 583
Connexin 32: 697-1545
TagFP635: 1567-2280
SV40 early mRNA polyadenylation signal
Polyadenylation signals: 2433-2438 2462-2467
mRNA 3' ends: 2471 2483
f1 single-strand DNA origin: 2530-2985
Bacterial promoter for expression of Kan^r gene
-35 region: 3047-3052
-10 region: 3070-3075
Transcription start point: 3082
SV40 origin of replication: 3326-3461
SV40 early promoter
Enhancer (72-bp tandem repeats): 3159-3230 3231-3302
21-bp repeats: 3306-3326, 3327-3347 3349-3369
Early promoter element: 3382-3388
Major transcription start points: 3378, 3416, 3422 3427
Kanamycin/neomycin resistance gene
Neomycin phosphotransferase coding sequences:
Start codon (ATG): 3510-3512
Stop codon: 4302-4304
G->A mutation to remove Pst I site: 3692
C->A (Arg to Ser) mutation to remove BssH II site: 4038
Herpes simplex virus (HSV) thymidine kinase (TK) polyadenylation signal
Polyadenylation signals: 4540-4545 4553-4558
pUC plasmid replication origin: 4889-5532

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 C. High efficiency gene transfer into mammalian cells. In DNA cloning: A Practical Approach, Vol. II. Ed. D. M. Glover. (IRL Press, Oxford, U.K.). 1985; 143-90.

Haas J, Park EC, Seed B. Codon usage limitation in the expression of HIV-1 envelope glycoprotein. *Curr Biol.* 1996; 6 (3):315-24. / pmid: 8805248

Kozak M. An analysis of 5'-noncoding sequences from 699 vertebrate messenger RNAs. *Nucleic Acids Res.* 1987; 15 (20):8125-48. / pmid: 3313277

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