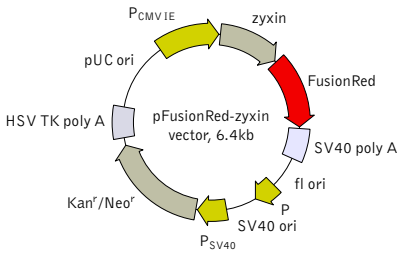
# pFusionRed-zyxin vector

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

| Product | Cat.# | Size |
|---|---|---|
| pFusionRed-zyxin vector | **FP425** | 20 $\mu$g |

| | |
|---|---|
| Vector type | mammalian expression vector |
| Reporter | FusionRed |
| Reporter codon usage | mammalian |
| Promoter for FusionRed | $P_{CMV\ IE}$ |
| Host cells | mammalian |
| Selection | prokaryotic - kanamycin<br>eukaryotic - neomycin (G418) |
| Replication | prokaryotic - pUC ori<br>eukaryotic - SV40 ori |
| Use | red fluorescent labeling of zyxin |

## Location of features

$P_{CMV\ IE}$: 1-589
Enhancer region: 59-465
TATA box: 554-560
Transcription start point: 583
Zyxin: 636-2348
FusionRed: 2370-3068
SV40 early mRNA polyadenylation signal
Polyadenylation signals: 3221-3226 & 3250-3255
mRNA 3' ends: 3259 & 3271
f1 single-strand DNA origin: 3318-3773
Bacterial promoter for expression of $Kan^r$ gene
-35 region: 3835-3840; -10 region: 3858-3863
Transcription start point: 3870
SV40 origin of replication: 4114-4249
SV40 early promoter
Enhancer (72-bp tandem repeats): 3947-4018 & 4019-4090
21-bp repeats: 4094-4114, 4115-4135 & 4137-4157
Early promoter element: 4170-4176
Major transcription start points: 4166, 4204, 4210 & 4215
Kanamycin/neomycin resistance gene
Neomycin phosphotransferase coding sequences:
Start codon (ATG): 4298-4300; Stop codon: 5090-5092
G->A mutation to remove Pst I site: 4480
C->A (Arg to Ser) mutation to remove BssH II site: 4826
Herpes simplex virus (HSV) thymidine kinase (TK) polyadenylation signal
Polyadenylation signals: 5328-5333 & 5341-5346
pUC plasmid replication origin: 5677-6320

## Vector description

pFusionRed-zyxin is a mammalian expression vector encoding FusionRed-zyxin fusion protein. The vector can be used for fluorescent labeling of zyxin in living cells.

FusionRed codon usage is optimized for high expression in mammalian cells (humanized) [Haas et al. 1996]. Human zyxin is fused to the FusionRed N-terminus.

pFusionRed-zyxin vector can be used as a source of FusionRed-zyxin 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

pFusionRed-zyxin vector can be transfected into mammalian cells by any known transfection method. CMV promoter provides strong, constitutive expression of the FusionRed-zyxin 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