# pFUSE-SEAP-mG3Fc 

Plasmid designed for the expression of a SEAP-Fc Fusion protein
Catalog \# pfuse-mg3sp
For research use only
Version 20K05-MM

## PRODUCT INFORMATION

## Content:

- $20 \mu \mathrm{~g}$ of pFUSE-SEAP-mG3Fc plasmid provided as lyophilized DNA
-1 ml of Zeocin ${ }^{\mathrm{TM}}(100 \mathrm{mg} / \mathrm{ml})$


## Storage and Stability:

- Product is shipped at room temperature.
- Lyophilized DNA should be stored at $-20^{\circ} \mathrm{C}$ and is stable 3 months.
- Resuspended DNA should be stored at $-20^{\circ} \mathrm{C}$ and is stable up to 1 year.
- Store Zeocin ${ }^{\mathrm{TM}}$ at $4^{\circ} \mathrm{C}$ or at $-20^{\circ} \mathrm{C}$. The expiry date is specified on the product label.


## Quality control:

- Plasmid construct has been confirmed by restriction analysis and sequencing.
- Plasmid DNA was purified by ion exchange chromatography and lyophilized.
- Expression of SEAP-mG3Fc was confirmed by using QUANTI-Blue ${ }^{\text {ru }}$ Solution.
- SEAP-mG3Fc protein was purified using protein G affinity chromatography following manufacturer's protocol.


## GENERAL PRODUCT USE

pFUSE-SEAP-Fc plasmids express a SEAP-Fc fusion protein generated by fusing the gene encoding for human secreted alkaline phosphatase (SEAP) and the Fc region of an immunoglobulin G (IgG).
pFUSE-SEAP-Fc plasmids yield high levels of Fc-Fusion proteins. The level of expression is usually in the $\mu \mathrm{g} / \mathrm{mL}$ range. They can be transfected in a variety of mammalian cells, including myeloma cell lines, Chinese hamster ovary ( CHO ) cells, monkey COS cells and human embryonic kidney (HEK)293 cells. These cells are commonly used in protein purification systems.
SEAP-Fc fusion proteins are secreted and can be easily detected in the supernatant of pFUSE-SEAP-Fc-transfected cells by using QUANTI-Blue ${ }^{\text {mu }}$ Solution, a SEAP detection medium. SEAP-Fc fusion proteins can be easily purified by single-step protein A or protein $G$ affinity chromatography.

## PLASMID FEATURES

- hEF1-HTLV prom is a composite promoter comprising the Elongation Factor- $1 \alpha(\mathrm{EF}-1 \alpha)$ core promoter ${ }^{1}$ and the R segment and part of the U5 sequence (R-U5') of the Human T-Cell Leukemia Virus (HTLV) Type 1 Long Terminal Repeat ${ }^{2}$. The EF-1 $\alpha$ promoter exhibits a strong activity and yields long lasting expression of a transgene in vivo. The R-U5' has been coupled to the EF-1 $\alpha$ core promoter to enhance stability of RNA.
- SEAP-mG3Fc was generated by fusing the gene encoding for human SEAP with the Fc region of mouse IgG3. This region comprises the CH 2 and CH 3 domains of the IgG heavy chain and the hinge region. The hinge serves as a flexible spacer between the SEAP and Fc moieties, allowing each part of the molecule to function independently.
- SV40 pAn: The Simian Virus 40 late polyadenylation signal enables efficient cleavage and polyadenylation reactions resulting in high levels of steady-state mRNA ${ }^{3}$.
- ori: A minimal $E$. coli origin of replication to limit vector size, but with the same activity as the longer Ori.
- CMV enh / hFerL prom: This composite promoter combines the human cytomegalovirus immediate-early gene 1 enhancer and the core promoter of the human ferritin light chain gene. This ubiquitous promoter drives the expression of the Zeocin ${ }^{\text {n" }}$-resistance gene in mammalian cells.
- EM2KC is a bacterial promoter that enables the constitutive expression of the antibiotic resistance gene in E. coli. EM2KC is located within an intron and is spliced out in mammalian cells.
- Zeo: Resistance to Zeocin ${ }^{\text {™ }}$ is conferred by the Sh ble gene from Streptoalloteichus hindustanus The same resistance gene confers selection in both mammalian cells and E. coli.
- BGIo pAn: The human beta-globin 3'UTR and polyadenylation sequence allows efficient arrest of the transgene transcription ${ }^{4}$.

1. Kim DW et al. 1990. Use of the human elongation factor 1 alpha promoter as a versatile and efficient expression system. 91(2):217-23.
2. Takebe Y. et al. 1988. SR alpha promoter: an efficient and versatile mammalian cDNA expression system composed of the simian virus 40 early promoter and the R-U5 segment of human T-cell leukemia virus type 1 long terminal repeat. Mol Cell Biol. 8(1):466-72. 3. Carswell S. \& Alwine JC. 1989. Efficiency of utilization of the simian virus 40 late polyadenylation site: effects of upstream sequences. Mol Cell Biol. 9(10):4248-58.
3. Yu J. \& Russell JE. 2001. Structural and functional analysis of an mRNP complex that mediates the high stability of human beta-globin mRNA. Mol Cell Biol. 21(17):5879-88.

## METHODS

## Plasmid resuspension

Quickly spin the tube containing the lyophilized plasmid to pellet the DNA. To obtain a plasmid solution at $1 \mu \mathrm{~g} / \mu \mathrm{l}$, resuspend the DNA in $20 \mu \mathrm{l}$ of sterile $\mathrm{H}_{2} \mathrm{O}$. Store resuspended plasmid at $-20^{\circ} \mathrm{C}$.

## Plasmid amplification and cloning

Plasmid amplification and cloning can be performed in E. coli GT116 or in other commonly used laboratory E. coli strains, such as DH5a.

## Zeocin ${ }^{\text {TM }}$ usage

This antibiotic can be used for $E$. coli at $25 \mu \mathrm{~g} / \mathrm{ml}$ in liquid or solid media and at $50-200 \mu \mathrm{~g} / \mathrm{ml}$ to select Zeocin ${ }^{\text {TM }}$-resistant mammalian cells.

## Purification of SEAP-mG3Fc protein

The following protocol describes the purification of SEAP-mG3Fc protein produced by 293 cells using Protein G affinity chromatography.

1- Seed $3.5 \times 10^{6} 293$ cells in a 100 mm plate containing 6 ml of DMEM supplemented with $10 \%$ FBS.
2- Transfect cells with $750 \mu \mathrm{l}$ of $\mathrm{pFUSE}-\mathrm{SEAP}-\mathrm{mG} 3 \mathrm{Fc} /$ LyoVec ${ }^{\text {™ }}$ complexes at a ratio of 1:6 prepared by mixing $7.5 \mathrm{\mu g}$ pFUSE-SEAP-mG3Fc and $750 \mu$ reconstituted LyoVec ${ }^{\text {" }}$ following the LyoVec ${ }^{\text {™ }}$ protocol.
3- After 16 hours transfection, replace the medium with a serum-free medium such as PRO 293a-CDM (Biowithaker-Cambrex).
4- After 72 hours transfection, collect supernatant.
5- Purifiy protein using Protein G affinity chromatography such as Hi Trap Protein G HP (Amersham Biosciences) following manufacturer's protocol.

## RELATED PRODUCTS

| Product | Catalog Code |
| :--- | :--- |
| LyoVec $^{\text {™ }}$ | lyec-12 |
| QUANTI-Blue ${ }^{\text {TM }}$ Solution | rep-qbs |



# PvuI (11) 

Sgfi (11)
1 GGATCTGCGATCGCTCCGGTGCCCGTCAGTGGGCAGAGCGCACATCGCCCACAGTCCCCGAGAAGTTGGGGGGAGGGGTCGGCAATTGAACGGGTGCCTA
101 GAGAAGGTGGCGCGGGGTAAACTGGGAAAGTGATGTCGTGTACTGGCTCCGCCTTTTTTCCCGAGGGTGGGGGAGAACCGTATATAAGTGCAGTAGTCGCC

8. $L$ 701 GCCCTGGGTGCCGCCAAGAAGCTGCAGCCTGCACAGACAGCCGCCAAGAACCTCATCATCTTCCTGGGCGATGGGATGGGGGTGTCTACGGTGACAGCTG


```
    BamHI (804)
```

801 CCAGGATCCTAAAAGGGCAGAAGAAGGACAAACTGGGGCCTGAGATACCCCTGGCTATGGACCGCTTCCCATATGTGGCTCTGTCCAAGACATACAATGT
 901 AGACAAACATGTGCCAGACAGTGGAGCCACAGCCACGGCCTACCTGTGCGGGGTCAAGGGCAACTTCCAGACCATTGGCTTGAGTGCAGCCGCCCGCTTT
 BstEII (1081)
1001 AACCAGTGCAACACGACACGCGGCAACGAGGTCATCTCCGTGATGAATCGGGCCAAGAAAGCAGGGAAGTCAGTGGGAGTGGTAACCACCACACGAGTGC
 1101 AGCACGCCTCGCCAGCCGGCACCTACGCCCACACGGTGAACCGCAACTGGTACTCGGACGCCGACGTGCCTGCCTCGGCCCGCCAGGAGGGGTGCCAGGA
 1201 CATCGCTACGCAGCTCATCTCCAACATGGACATTGATGTGATCCTGGGTGGAGGCCGAAAGTACATGTTTCGCATGGGAACCCCAGACCCTGAGTACCCA


1301 GATGACTACAGCCAAGGTGGGACCAGGCTGGACGGGAAGAATCTGGTGCAGGAATGGCTGGCGAAGCGCCAGGGTGCCCGGTATGTGTGGAACCGCACTG
 1401 AGCTCATGCAGGCTTCCCTGGACCCGTCTGTGACCCATCTCATGGGTCTCTTTGAGCCTGGAGACATGAAATACGAGATCCACCGAGACTCCACACTGGA

2751E L $\quad$ M $\quad$ Q $A$ SacII (1558) PshAI (1591) 1501 CCCCTCCCTGATGGAGATGACAGAGGCTGCCCTGCGCCTGCTGAGCAGGAACCCCCGCGGCTTCTTCCTCTTCGTGGAGGGTGGTCGCATCGACCACGGT
 1601 CATCACGAAAGCAGGGCTTACCGGGCACTGACTGAGACGATCATGTTCGACGACGCCATTGAGAGGGCGGGCCAGCTCACCAGCGAGGAGGACACGCTGA

 XcmI (1771)<br>SacI (1763)

1701 GCCTCGTCACTGCCGACCACTCCCACGTCTTCTCCTTCGGAGGCTACCCCCTGCGAGGGAGCTCCATCTTCGGGCTGGCCCCTGGCAAGGCCCGGGACAG
 StuI (1805)

BsrBI (1884)
1801 GAAGGCCTACACGGTCCTCCTATACGGAAACGGTCCAGGCTATGTGCTCAAGGACGGCGCCCGGCCGGATGTTACCGAGAGCGAGAGCGGGAGCCCCGAG
 BssHII (1967)
1901 TATCGGCAGCAGTCAGCAGTGCCCCTGGACGAAGAGACCCACGCAGGCGAGGACGTGGCGGTGTTCGCGCGCGGCCCGCAGGCGCACCTGGTTCACGGCG
 2001 TGCAGGAGCAGACCTTCATAGCGCACGTCATGGCCTTCGCCGCCTGCCTGGAGCCCTACACCGCCTGCGACCTGGCGCCCCCCGCCGGCACCACCGACGC
 BgIII (2138)
2101 CGCGCACCCGGGGCGGTCCCGGTCCAAGCGTCTGGATAGATCTCCTAGAATACCCAAGCCCAGTACCCCCCCAGGTTCTTCATGCCCACCTGGTAACATC 1. $\mathrm{P} \quad \mathrm{R} \quad \mathrm{I} \quad \mathrm{P} \quad \mathrm{K} \quad \mathrm{P}$
 2201 TTGGGTGGACCATCCGTCTTCATCTTCCCCCCAAAGCCCAAGGATGCACTCATGATCTCCCTAACCCCCAAGGTTACGTGTGTGGTGGTGGATGTGAGCG 20. $\mathrm{L} \quad \mathrm{G} \quad \mathrm{G} \quad \mathrm{P} \quad \mathrm{S} \quad \mathrm{V}$ 542 L $\quad \mathrm{G} \quad \mathrm{G} \quad \mathrm{P} \quad \mathrm{S} \quad \mathrm{V}$ 2301 AGGATGACCCAGATGTCCATGTCAGCTGGTTTGTGGACAACAAAGAAGTACACACAGCCTGGACACAGCCCCGTGAAGCTCAGTACAACAGTACCTTCCG 53 E D D P $\quad \mathrm{D}$
 2401 AGTGGTCAGTGCCCTCCCCATCCAGCACCAGGACTGGATGAGGGGCAAGGAGTTCAAATGCAAGGTCAACAACAAAGCCCTCCCAGCCCCCATCGAGAGA 86. V V V
 Bst1107I (2539)
2501 ACCATCTCAAAACCCAAAGGAAGAGCCCAGACACCTCAAGTATACACCATACCCCCACCTCGTGAACAAATGTCCAAGAAGAAGGTTAGTCTGACCTGCC



2601 TGGTCACCAACTTCTTCTCTGAAGCCATCAGTGTGGAGTGGGAAAGGAACGGAGAACTGGAGCAGGATTACAAGAACACTCCACCCATCCTGGACTCAGA

 2701 TGGGACCTACTTCCTCTACAGCAAGCTCACTGTGGATACAGACAGTTGGTTGCAAGGAGAAATTTTTACCTGCTCCGTGGTGCATGAGGCTCTCCATAAC
 708. G T Y F L Y S K L T V D T D S W L Q G E I F T C S Ball (2851)
NheI (2843)
2801 CACCACACACAGAAGAACCTGTCTCGCTCCCCTGGTAAATGAGCTAGCTGGCCAGACATGATAAGATACATTGATGAGTTTGGACAAACCACAACTAGAA 220. H H T Q K N L S R S P G K •

742 H H T Q K N L S R S P G K •
HpaI (2983)
2901 TGCAGTGAAAAAAATGCTTTATTTGTGAAATTTGTGATGCTATTGCTTTATTTGTAACCATTATAAGCTGCAATAAACAAGTTAACAACAACAATTGCAT AseI (3080) XmnI (3076)

|  | AseI (3080) |
| :---: | :---: |
| 3001 | XmnI (3076) <br> TCATTTTATGTTTCAGGTTCAGGGGGAGGTGTGGGAGGTTTTTTAAAGCAAGTAAAACCTCTACAAATGTGGTATGGAATTAATTCTAAAATACAGCATA |
|  | $\rightarrow \quad 4$ |
| 3101 | GCAAAACTTTAACCTCCAAATCAAGCCTCTACTTGAATCCTTTTCTGAGGGATGAATAAGGCATAGGCATCAGGGGCTGTTGCCAATGTGCATTAGCTGT |
| 3201 | TTGCAGCCTCACCTTCTTTCATGGAGTTTAAGATATAGTGTATTTTCCCAAGGTTTGAACTAGCTCTTCATTTCTTTATGTTTTAAATGCACTGACCTCC |
| 3301 | $\begin{array}{rr\|r\|r\|cc\|} \hline \mathbf{S s p I} \mathbf{( 3 3 2 2 )} & \text { SwaI (3337) } \\ \text { CACATTCCCTTTTAGTAAAATATTCAGAAATAATTTAAATACATCATTGCAATGAAAATAAATGTTTTTTATTAGGCAGAATCCAGATGCTCAAGGCCC } \end{array}$ |
| 3401 | TTCATAATATCCCCCAGTTTAGTAGTTGGACTTAGGGAACAAAGGAACCTTTAATAGAAATTGGACAGCAAGAAAGCGAGCTTCTAGCTTATCCTCAGTC |
|  | ApaLI (3519) |

3501 CTGCTCCTCTGCCACAAAGTGCACGCAGTTGCCGGCCGGGTCGCGCAGGGCGAACTCCCGCCCCCACGGCTGCTCGCCGATCTCGGTCATGGCCGGCCCG

3601 GAGGCGTCCCGGAAGTTCGTGGACACGACCTCCGACCACTCGGCGTACAGCTCGTCCAGGCCGCGCACCCACACCCAGGCCAGGGTGTTGTCCGGCACCA
 SgrAI (3750)
3701 CCTGGTCCTGGACCGCGCTGATGAACAGGGTCACGTCGTCCCGGACCACACCGGCGAAGTCGTCCTCCACGAAGTCCCGGGAGAACCCGAGCCGGTCGGT 561 Q D D BsrBI (3817) BssHII (3828) Ball (3865)
3801 CCAGAACTCGACCGCTCCGGCGACGTCGCGCGCGGTGAGCACCGGAACGGCACTGGTCAACTTGGCCATGATGGCTCCTCctgtcaggagaggaaagaga
 AseI (3964)
3901 agaaggttagtacaattgCTATAGTGAGTTGTATTATACTATGCAGATATACTATGCCAATGATTAATTGTCAAACTAGGGCTGCAgggttcatagtgcc HindIII (4090)
4001 acttttcctgcactgccccatctcctgcccaccctttcccaggcatagacagtcagtgacttacCAAACTCACAGGAGGGAGAAGGCAGAAGCTTGAGAC
SacII (4108)
StuI (4192)

4101 AGACCCGCGGGACCGCCGAACTGCGAGGGGACGTGGCTAGGGCGGCTTCTTTTATGGTGCGCCGGCCCTCGGAGGCAGGGCGCTCGGGGAGGCCTAGCGG
BspEI (4248)
4201 CCAATCTGCGGTGGCAGGAGGCGGGGCCGAAGGCCGTGCCTGACCAATCCGGAGCACATAGGAGTCTCAGCCCCCCGCCCCAAAGCAAGGGGAAGTCACG

## SpeI (4355)

Bsp120I (4347)
4301 CGCCTGTAGCGCCAGCGTGTTGTGAAATGGGGGCTTGGGGGGGTTGGGGCCCTGACTAGTCAAAACAAACTCCCATTGACGTCAATGGGGTGGAGACTTG
 NdeI (4589)
4501 AGTAGGAAAGTCCCATAAGGTCATGTACTGGGCATAATGCCAGGCGGGCCATTTACCGTCATTGACGTCAATAGGGGGCGTACTTGGCATATGATACACT
4601 TGATGTACTGCCAAGTGGGCAGTTTACCGTAAATACTCCACCCATTGACGTCAATGGAAAGTCCCTATTGGCGTTACTATGGGAACATACGTCATTATTG PacI (4778)
SdaI (4771)
4701 ACGTCAATGGGCGGGGGTCGTTGGGCGGTCAGCCAGGCGGGCCATTTACCGTAAGTTATGTAACGCCTGCAGGTTAATTAAGAACATGTGAGCAAAAGGC
4801 CAGCAAAAGGCCAGGAACCGTAAAAAGGCCGCGTTGCTGGCGTTTTTCCATAGGCTCCGCCCCCCTGACGAGCATCACAAAAATCGACGCTCAAGTCAGA
4901 GGTGGCGAAACCCGACAGGACTATAAAGATACCAGGCGTTTCCCCCTGGAAGCTCCCTCGTGCGCTCTCCTGTTCCGACCCTGCCGCTTACCGGATACCT

## ApaLI (5098)

5001 GTCCGCCTTTCTCCCTTCGGGAAGCGTGGCGCTTTCTCATAGCTCACGCTGTAGGTATCTCAGTTCGGTGTAGGTCGTTCGCTCCAAGCTGGGCTGTGTG
5101 CACGAACCCCCCGTTCAGCCCGACCGCTGCGCCTTATCCGGTAACTATCGTCTTGAGTCCAACCCGGTAAGACACGACTTATCGCCACTGGCAGCAGCCA
5201 CTGGTAACAGGATTAGCAGAGCGAGGTATGTAGGCGGTGCTACAGAGTTCTTGAAGTGGTGGCCTAACTACGGCTACACTAGAAGAACAGTATTTGGTAT
5301 CTGCGCTCTGCTGAAGCCAGTTACCTTCGGAAAAAGAGTTGGTAGCTCTTGATCCGGCAAACAAACCACCGCTGGTAGCGGTGGTTTTTTTGTTTGCAAG
5401 CAGCAGATTACGCGCAGAAAAAAAGGATCTCAAGAAGATCCTTTGATCTTTTCTACGGGGTCTGACGCTCAGTGGAACGAAAACTCACGTTAAGGGATTT PacI (5518) SwaI (5526) NotI (5534)
5501 TGGTCATGGCTAGTTAATTAACATTTAAATCAGCGGCCGCAATAAAATATCTTTATTTTCATTACATCTGTGTGTTGGTTTTTTGTGTGAATCGTAACTA
5601 ACATACGCTCTCCATCAAAACAAAACGAAACAAAACAAACTAGCAAAATAGGCTGTCCCCAGTGCAAGTGCAGGTGCCAGAACATTTCTCTATCGAA

