

# pFUSE-rIgG-Fc2 (IL2ss)

Plasmid designed for the construction of Fc-Fusion proteins

Catalog # pfuse-rfc2

## For research use only

Version 20K05-MM

## PRODUCT INFORMATION

### Content:

- 20 µg of pFUSE-rIgG-Fc2 (IL2ss) plasmid provided as lyophilized DNA
- 1 ml of Zeocin™ (100 mg/ml)

### Storage and Stability:

- Product is shipped at room temperature.
- Lyophilized DNA should be stored at -20°C and is stable 3 months.
- Resuspended DNA should be stored at -20°C and is stable up to 1 year.
- Store Zeocin™ at 4 °C or at -20 °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.

## GENERAL PRODUCT USE

pFUSE-Fc is a family of plasmid developed to facilitate the construction of Fc-Fusion proteins by fusing a sequence encoding a given protein to the Fc region of an immunoglobulin.

pFUSE-Fc plasmids yield high levels of Fc-Fusion proteins. The level of expression is usually in the µg/mL range. They can be transfected in a variety of mammalian cells, including myeloma cell lines, CHO cells, monkey COS cells and human embryonic kidney (HEK)293 cells. These cells are commonly used in protein purification systems.

pFUSE-Fc2 (IL2ss) plasmids allow the secretion of Fc-Fusion proteins. They contain the IL2 signal sequence (IL2ss) for the generation of Fc-Fusion proteins derived from proteins that are not naturally secreted. As Fc-Fusion proteins are secreted, they can be easily detected in the supernatant of pFUSE-Fc-transfected cells by SDS-PAGE. Furthermore, functional domains can be identified by immunoblotting and ligand blotting.

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-1α (EF-1α) core promoter<sup>1</sup> 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<sup>2</sup>. The EF-1α promoter exhibits a strong activity and yields long lasting expression of a transgene *in vivo*. The R-U5' has been coupled to the EF-1α core promoter to enhance stability of RNA.

• **MCS:** The multiple cloning site contains several restriction sites that are compatible with many other enzymes, thus facilitating cloning.

• **IL2 ss:** The IL2 signal sequence contains 20 amino acids and share common characteristics with signal peptides of other secretory proteins. The intracellular cleavage of the IL2 signal peptide occurs after Ser20 and leads to the secretion of the antigenic protein.

• **rIgG Fc (rabbit):** The Fc region comprises the CH2 and CH3 domains of the IgG heavy chain and the hinge region. The hinge serves as a flexible spacer between the two parts of the Fc-Fusion protein, 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<sup>3</sup>.

• **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™-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™ is conferred by the *Sh ble* gene from *Streptallocteichus hindustanus*. The same resistance gene confers selection in both mammalian cells and *E. coli*.

• **BGlo pAn:** The human beta-globin 3'UTR and polyadenylation sequence allows efficient arrest of the transgene transcription<sup>4</sup>.

## METHODS

### **Plasmid resuspension**

Quickly spin the tube containing the lyophilized plasmid to pellet the DNA. To obtain a plasmid solution at 1 µg/µl, resuspend the DNA in 20 µl of sterile H<sub>2</sub>O. Store resuspended plasmid at -20°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 DH5α.

### **Zeocin™ usage**

This antibiotic can be used for *E. coli* at 25 µg/ml in liquid or solid media and at 50-200 µg/ml to select Zeocin™-resistant mammalian cells.

### **References:**

1. Kim, D.W. *et al.* (1990). Gene 2: 217-223.
2. Takebe, Y. *et al.* (1988). Mol. Cell Biol. 1: 466-472.
3. Carswell, S., and Alwine, J.C. (1989). Mol. Cell Biol. 10: 4248-4258.
4. Yu J & Russell JE. (2001). Mol Cell Biol, 21(17):5879-88.

## TECHNICAL SUPPORT

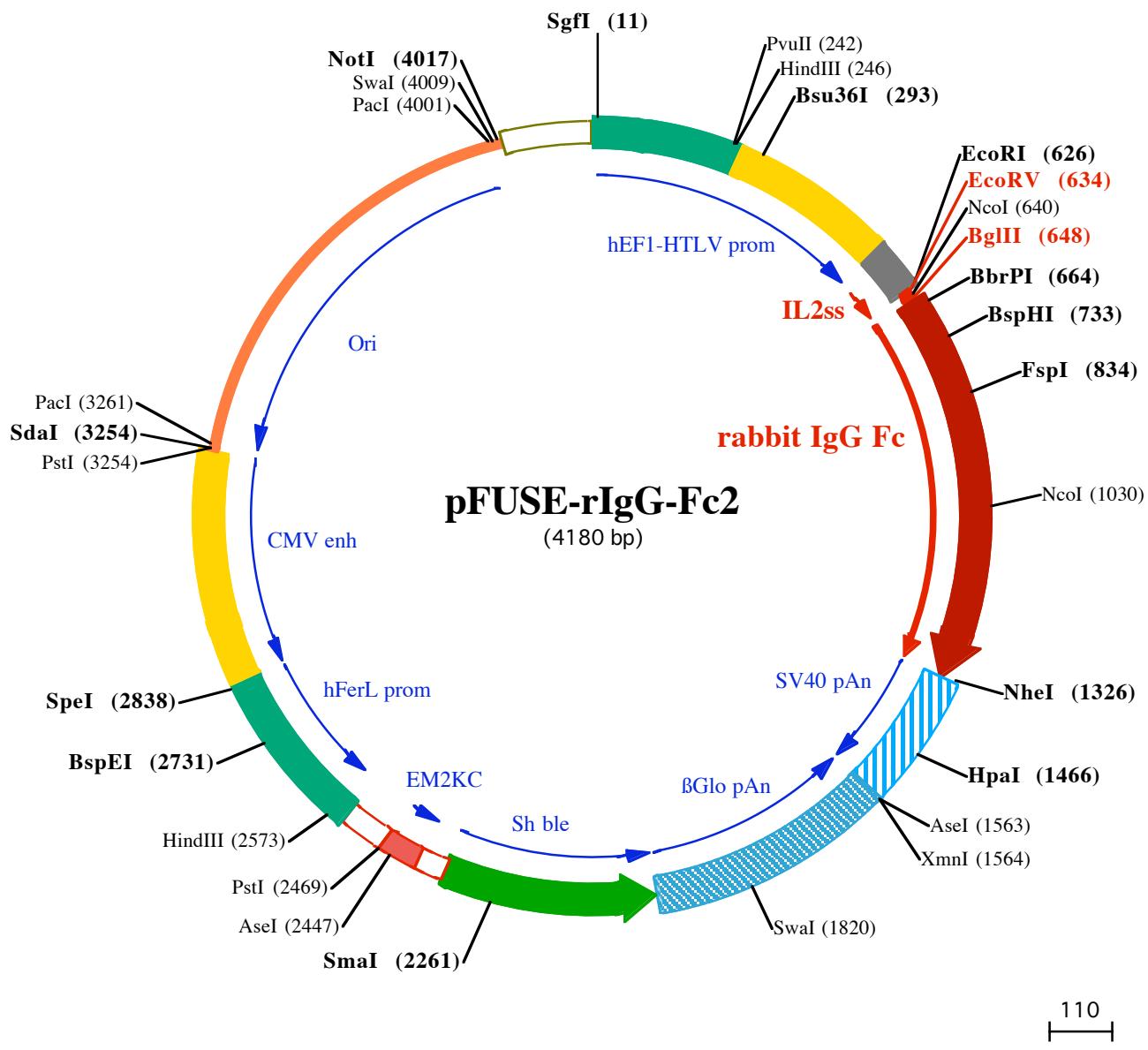
InvivoGen USA (Toll-Free): 888-457-5873

InvivoGen USA (International): +1 (858) 457-5873

InvivoGen Europe: +33 (0) 5-62-71-69-39

InvivoGen Hong Kong: +852 3622-3480

E-mail: info@invivogen.com



**SgfI (11)**

1 GGATCTCGATCGTCCGGTCCCCGTCAAGTGGGAGAGCGCACATGCCACAGTCCCAGAAGTTGGGGAGGGTCGGCAATTGAACGGTGCCTA

101 GAGAAGGTGGCGGGGTAAACTGGAAAGTGATTCGTACTGGCTCCCTTCCGAGGGTGGGGAGAACGTATAAGTCAGTAGTCGC

HindIII (246)

PvuII (242)

201 GTAACGTTCTTTCGCAACGGTTGCCAGAACACAGCTGAGCTCGAGGGGCTGCATCTCCTCACGCCGCCCTACCTGAGGCC

Bsu36I (293)

301 GCCATCCACGCCGGTGGAGTCGCGTCTGCCCTCCGCTGTGGTGCCTCTGAACCTCGTCCCGCTAGTAAAGCTCAGTCAGGACC

401 GGGCTTGTCCGGCCTCCCTGGACCTACCTAGACTCAGCGGCTCTCACGCTTGCTGACCCCTGCTCAACTCTACGCTTTGTTGTT

501 TCTGTTCTCGCCGTTACAGATCCAAGCTGACCGCGCTACCTGAGATCACccgcGAAGGAGGGCACCATGTACAGGATCACTCTGTCTGCA

1►MetTyrArgMetGI nLeuLeuSer CysI

EcoRV (634)

EcoRI (626) NcoI (640) BglIII (648) BbrPI (664)

601 TTGCACTAAGTCTTGCACTTGTCAACGAAATCGATATCGCCATGGTTAGATCTAGCAAGGCCACGTGCCACCCCTGAACCTCTGGGGGACCGTGTG

10►IeAl iLeuSer LeuAl iLeuVal Thr AsnSer 1►Ser LysProThr CysProProProGl uLeuLeuGl yGl yProSer Va

BspHI (733)

701 CTTCATCTTCCCCAAAACCAAGGACACCCCTCATGATCTCACGCCACCCCGAGGTACATGCGTGGTGGACGTGAGCCAGGATGACCCGAGGTG

16►IPhelePheProProLysProLysAspThrLeuMetIleSerArgThrProGl uVal Thr CysVal Val Val AspVal Ser Gl nAspAspProGl uVal

FspI (834)

801 CAGTTACATGGTACATAAACAAACGAGCAGGTGCCACGCCGGCGCTACGGAGCAGCAGTTACACAGCACGATCCGCGTGGTCAACCCCTCC

50►Gl nPheThr TrpTyr IleAsnGl uGl nVal ArgThr Al aArgProProLeuArgGl uGl nGl nPheAsnSerThr IleArgVal Val Ser Thr LeuP

901 CCATCGCGCACGAGCTGGCTGGGGCAAGGAGTCAGTCAAAGTCCACAAACAGGACTCCGGCCCATCGAGAAAACCATCTCAAAGCCAG

83►rolleAl aHi sGl nAspTrpLeuArgGl yLysGl uPheLysCysValHi sAsnLysAl aLeuProAl aProI leGl uLysThr IleSer LysAl aAr

NcoI (1030)

1001 AGGGCAGCCCCCTGGAGCGAAGGTACACCATGGCCCTCCCGGGAGGAGCTGAGCAGGTGGTCAGCCTGACCTGCATGATCAACGGCTTCTAC

116►gGl yGl nProLeuGl uProLysVal IleTyrThr MetIleTyrProArgGl uGl uLeuSer Ser ArgSer Val IleSer LeuThr CysMetIleAsnGl yPheTyr

1101 CCTTCGGACATCTGGTGGAGTGGAGAAGACGGAGGGACAGGACAACATAAGACCACGCCGGCTGCTGGACAGCGACGGCTCTACTTCTCT

150►ProSerAspIleSerValGl uTrpGl uLysAsnGl yLysAl aGl uAspAsnTyrLysThr Thr ProAl aVal LeuAspSerAspGl ySer TyrPheLeuT

1201 ACAGCAAGCTCTAGTCCACAGAGTGGAGCTGGCAGCGGGCGACGCTTCACTGCTCGTGTAGTGACGAGGCCCTGCACAACACTACACGAGACT

183►ySerLysLeuSerValProThrVal uTrpGl nArgGl yAspVal PheThr CysSer Val MetIleHisGl uAl aLeuHi sAsnHisTyrThr Gl nLysSe

NheI (1326)

1301 CATCTCCGCTCTCCGGTAAATGAGCTAGTGGCAGACATGATAAGATACTATTGATGAGTTGGACAAACACAACACTAGAATGAGTGAACAAATGCATTCTTTATGTTCAAG

216►rIleSerArgSerProGl yLys•••

HpaI (1466)

1401 TTTATTGTGAAATTGTGATGCTATTGCTTATTGTAACCATTATAAGCTGAAATAAACAGTTAACACAATTGATTCTAGTGGTCAACCTTACCTTATGTTCAAG

AseI (1563)

XmnI (1564)

1501 TTCAGGGGAGGTGGAGGTTAAAGCAAGTAAACCTCTACAAATGTGGTATGGAATTAAATTCTAAACATAGCAAAACTTAACTCC

1601 AAATCAAGCTCTACTTGAATCTTCTGAGGGATGAATAAGGCATAGGCATAGGGCTGTTGCAATGTCATTAGCTGTTGAGCCTCACCTCT

1701 TTCATGGAGTTAACATATAGTGTATTTCCAAGGTTGAACTAGCTCTCATTCTTATGTTAAATGCACTGACCTCCACATTCCCTTTAGT

Swal (1820)

1801 AAAATATTAGAAATAATTAAATACATCATTGCAATGAAAATAATGTTTATTAGGCAGAACATGCTCAAGGCCCTCATATAATCCCCAG

1901 TTTAGTAGTTGACTTAGGGAACAAAGAACCTTAAAGAAATTGGACAGCAAGAACGGAGCTTAGCTTACCTCTAGCTCTGCTCTGCCAAC

125►•••AspGl nGl uGl uAl aVal Ph

2001 AGTCACCGAGTGGCCGGGTCGCGAGGGCAACTCCGCCACGGCTGCTCGCCATCGGTATGGCAGGCGCCGGAGGGCTCCGGAAAGT

117►eHi sVal CysAsnGl yAl aProAspArgLeuAl aPheGl uArgGl uTrpGl nGl uGl yIleGl uThr MetAl aProGl ySer Al aAspArgPheAsn

2101 CGTGGACACGACTCCGACCACTGGCTACAGCTGTCAGGGCCACCCACACCCAGGGCTGTTGCGCACCCATGGTCTGGACCCGG

84►ThrSerValValGl uSerTrpGl uAl aTyrLeuGl uAspLeuGl yArgValTrpValTrpAl aLeuThrAsnAspProValValGl nAspGl nValAl aS

SmaI (2261)

2201 CTGATGACAGGGTCAGTCGTCCTCCGGACACACCCGGCAAGTCGCTCTCCAGAAGTCCGGAGACCCGAGCCGGTGGCCAGACTCGACCGCTC

50►erIlePheLeuThrValAspArgValValGl yAl aPheAspAspArgVal uVal PheAspArgSerPheGl yLeuArgAspThrTrpPheGl uValAl aGl

2301 CGGGCACGTGCGCGGGTGGACCCGGCACTGGCATGTTCTctgtcaggagagggaaagagaagaaagttatcatatt

17►yAl aValAspArgAl aThrLeuValProValAl aSerThrLeuLysAl aMet

AseI (2447) PstI (2469)

2401 gCTATAGTGAGTTGATTATACTATGAGATATACTATGCCAATGATTAATTGTCAGGGCTGCAgggttcatagtgccactttctgcactgc

HindIII (2573)

2501 ccatctccgtcccccaccccccattccaggcatagacagtcaagtactacCAAACCTACAGGAGGGAGAAGGCAGAACGTTGAGACAGACCCGGACGCC

2601 GAACTGCGAGGGGAGTGGCTAGGGCGCTTCTTTATGGTGCGCCCTCGGAGGCAGGGCGCTGGGGAGGCTAGCGCCAACTGCGGTGGCAG

BspEI (2731)

2701 GAGCGGGGGCGAAGGCCGTGCCTGACCATCCGGAGCACATAGGAGTCTCAGCCCCCGCCCAAAGCAAGGGAGTCACGCCCTGTAGGCCAGCG

2801 TGTTGAAATGGGGCTGGGGGGCTGACTAGTCAGTCAGTCATGGTAACTGGGAGACTTGGAAATCCCCGTGAGTC

2901 AACCGCTATCCACCCCCATTGATGACTGCCAAACCGCATCATGGTAATAGCGATGACTAACGTTAGATGACTGCCAGTAGGAAAGTCCATA

3001 AGGTCTGACTGGCATAATGCCAGGGGGCATTACCGTCATTGAGCTCAATAGGGGCTACTTGGCATATGATACACTTGATGACTGCCAGTG

3101 GGCAGTTACCGTAAACTCCACCCATTGACGTCAATGAAAGTCCATTGGCTTACTATGGAACATACGTCAATTGACGTCAATGGCGGGGG

PacI (3261)  
PstI (3254)  
**SdaI (3254)**

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3201 TCGTGGCGGTAGCCAGGCGGGCATTACCGTAAGTTATGTAACGCCCTGCAGGTTAATAAGAACATGTGAGCAAAAGGCCAGCAAAGGCCAGGA  
 3301 CCGTAAAAGGCCCGTGGCTGGCTTTCCATAGGCTCGCCCCCTGACGAGCATCACAAAATCGACGCTCAAGTCAGAGGTGGCAGAACCCGACA  
 3401 GGAATATAAGATACCAGGCCTTCCCCCTGGAAGCTCCCTCGCGCTCTCTGTTCCGACCCCTGCCGCTTACCGGATACCTGTCGCCCTTCTCCCT  
 3501 CGGGAAAGCGTGGCGCTTCTCATAGCTCACGCTGTAGGTATCTCAGTTGGTGTAGGTGTTGCTCCAAGCTGGCTGTGCACGAACCCCCGTTCA  
 3601 GCCCGACCGCTGCCCTATCGGTAACTATCGTCTGAGTCCAACCCGTAAGACACGACTTATGCCACTGGCAGCAGCCACTGGTAACAGGATTAGC  
 3701 AGAGCGAGGTATGAGCGGTGCTACAGAGTTCTGAAGTGGTGGCTAACTACGGCTACACTAGAAGAACAGTATTGGTATCTGCCTGCTGAAGC  
 3801 CAGTTACCTCGAAAAAGAGTTGGTAGCTCTTGATCCGCAAACAAACCACCGCTGGTAGCGGTGGTTTGTCAAGCAGCAGATTACGCCAG  


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PacI (4001)

3901 AAAAAAAGGATCTAAGAAGATCCTTGATCTTCTACGGGTCTGACGCTAGTGGAACGAAACTCACGTTAAGGGATTTGGTCAAGCTGGCTAGTTAA

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Swal (4009)    **NotI (4017)**

4001 TTAACATTTAAATCAGCGGCCGAAATAAAATATCTTATTTCTTACATCTGTGTTGGTTTTGTGAATGTAACATACGCTCTCCATCA  
 4101 AAACAAAAGAAACAAACAAACTAGCAAAATAGGCTGCCCCAGTCAAGTCAGGTGCCAGAACATTCTATCGAA