

pUNO1-hSTING-N200

Expression vector containing N200 isoform human STING (I200N) open reading frame

Catalog code: puno1-hsting-n200

<https://www.invivogen.com/hsting-n200>

For research use only

Version 19K10-MM

PRODUCT INFORMATION

Contents

- 20 µg of lyophilized plasmid DNA
- 2 x 1 ml blasticidin at 10 mg/ml

Storage and Stability

- Product is shipped at room temperature.
- Lyophilized DNA should be stored at -20°C.
- Resuspended DNA should be stored at -20°C and is stable at least for 1 year.
- Store blasticidin at 4°C or -20°C.*

*The expiry date is specified on the product label.

Quality control

- Plasmid construct has been confirmed by restriction analysis and full-length open reading frame (ORF) sequencing.
- Plasmid DNA was purified by ion exchange chromatography.

GENERAL PRODUCT USE

- **Subclone gene into another vector.** Two unique restriction sites flank the gene, allowing convenient excision. The 5' site is BspEI which is compatible with AgeI, XmaI, NgoMIV and SgrAI. The 3' site is NheI which is compatible with XbaI, SpeI, and AvrII.
- **Stable gene expression in mammalian cells.** pUNO1 plasmids can be used directly in transfection experiments both *in vitro* and *in vivo*. pUNO1 plasmids contain the blasticidin-resistance gene (*bsr*) driven by the CMV promoter/enhancer in tandem with the bacterial EM7 promoter. This allows the amplification of the plasmid in *E. coli*, as well as the selection of stable clones in mammalian cells using the same selective antibiotic. pUNO1 allows high levels of expression and secretion of the gene product.

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 water. Store resuspended plasmid at -20°C.

Plasmid amplification and cloning

Plasmid amplification and cloning can be performed in *E. coli* GT116 or other commonly used laboratory *E. coli* strains, such as DH5α.

Blasticidin usage

Blasticidin should be used at 25-100 µg/ml in bacteria and 1-30 µg/ml in mammalian cells. Blasticidin is supplied as a 10 mg/ml colorless solution in HEPES buffer.

PLASMID FEATURES

- **Bsr (blasticidin resistance gene):** The *bsr* gene from *Bacillus cereus* encodes a deaminase that confers resistance to the antibiotic blasticidin. The *bsr* gene is driven by the CMV promoter/enhancer in tandem with the bacterial EM7 promoter. Therefore, blasticidin can be used to select stable mammalian cells transfectedants and *E. coli* transformants.
- **CMV promoter & enhancer** drives the expression of the blasticidin resistance in mammalian cells.

Human STING-N200

ORF size: 1140 bp

Cloning fragment size: 1181 bp

STING (stimulator of interferon genes; also known as TMEM173, MITA, MPYS, and ERIS) is essential for the IFN response to microbial or self-DNA, and acts as a direct sensor of cyclic dinucleotides (CDNs). CDNs are important messengers in bacteria, affecting numerous responses of the prokaryotic cell, but also in mammalian cells, acting as agonists of the innate immune response. Studies have revealed that STING variation can affect CDN recognition and signal transduction. The hSTING-N200 isoform harbors a missense mutation (I200N) equivalent to I199N mutation found in the Goldenticket (*Gt*) mouse strain^{1,2}. Residue I200 is buried in the interior of the STING protomer. The I200N mutation results in a null-phenotype with no detectable STING activity³.

• **EF-1 α /HTLV hybrid promoter** is a composite promoter comprised of the Elongation Factor-1 α (EF-1 α) core promoter³ and the 5' untranslated region of the Human T-Cell Leukemia Virus (HTLV). EF-1 α utilizes a type 2 promoter that encodes for a «house keeping» gene. It is expressed at high levels in all cell cycles and lower levels during G0 phase. The promoter is also non-tissue specific; it is highly expressed in all cell types. The R segment and part of the U5 sequence (R-U5') of the HTLV Type 1 Long Terminal Repeat⁴ has been coupled to the EF-1 α promoter to enhance stability of DNA and RNA. This modification not only increases steady state transcription, but also significantly increases translation efficiency possibly through mRNA stabilization.

• **SV40 pAn:** The Simian Virus 40 late polyadenylation signal enables efficient cleavage and polyadenylation reactions, resulting in high levels of steady-state mRNA⁵.

• **pMB1 ori** is a minimal *E. coli* origin of replication to limit vector size, but with the same activity as the longer Ori.

• **Human beta-Globin polyA** is a strong polyadenylation (pAn) signal placed downstream of *bsr*. The use of beta-globin pAn minimizes interference⁶ and possible recombination events with the SV40 polyadenylation signal.

1. Yin Q. et al., 2012. Cyclic di-GMP sensing via the innate immune signaling protein STING. Mol Cell 46(6):735-45. 2. Sauer JD. et al., 2011. The N-ethyl-N-nitrosourea-induced Goldenticket mouse mutant reveals an essential function of Sting in the in vivo interferon response to Listeria monocytogenes and cyclic dinucleotides. Infect Immun 79(2):688-94. 3. Kim D. et al., 1990. Use of the human elongation factor 1 α promoter as a versatile and efficient expression system. Gene 91(2):217-23. 4. 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. 5. Carswell S. & Alwine J., 1989. Efficiency of utilization of the simian virus 40 late polyadenylation site: effects of upstream sequences. Mol Cell Biol. 9(10):4248-58. 6. Yu J. & Russell J., 2001. Structural and functional analysis of an mRNP complex that mediates the high stability of human β -globin mRNA. Mol Cell Biol. 21(17):5879-88.

RELATED PRODUCTS

Product	Description	Cat. Code
Blasticidin ChemiComp GT116	Selection antibiotic Competent <i>E. coli</i>	ant-bl-1 gt116-11

TECHNICAL SUPPORT

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

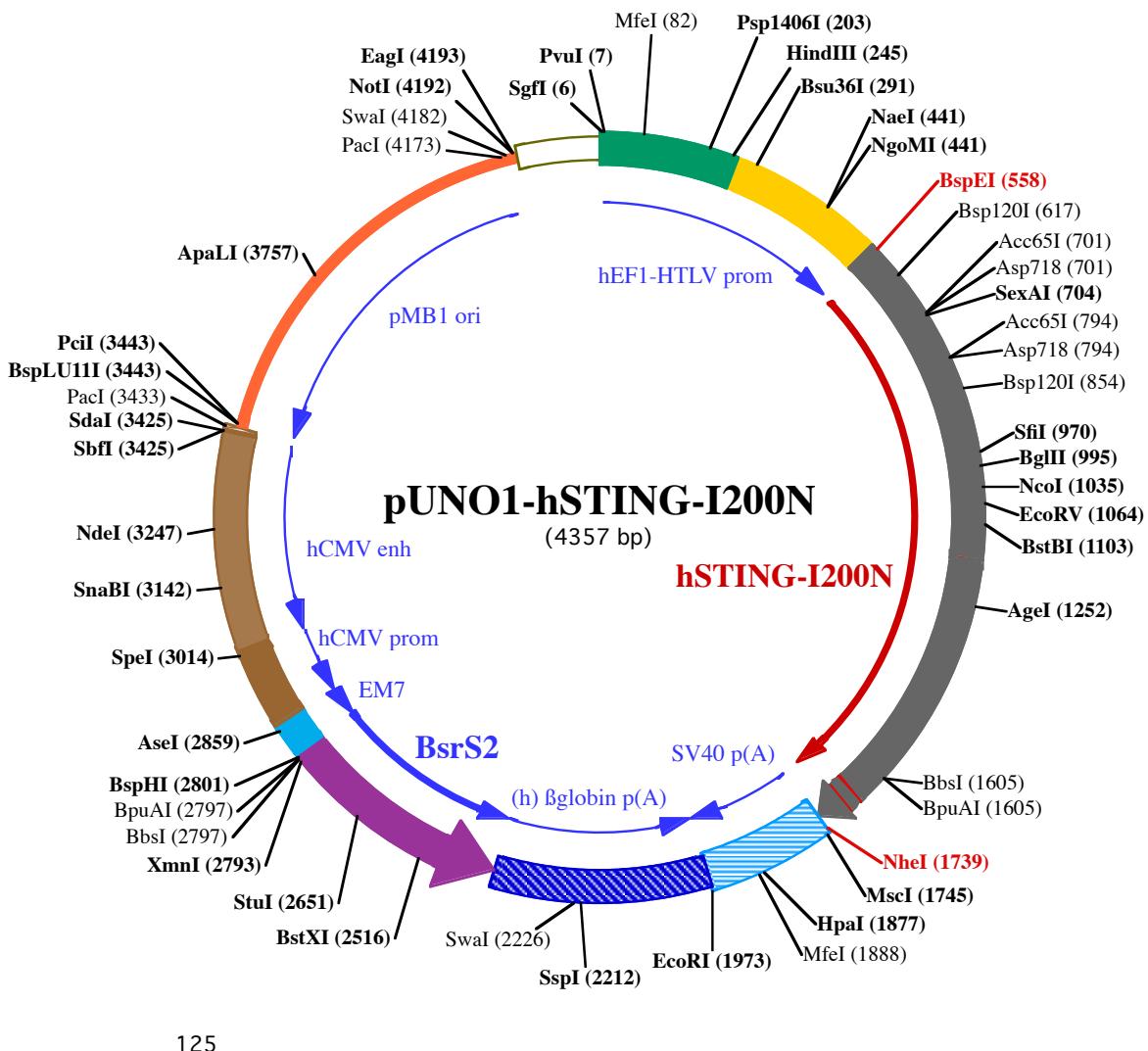
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PvuI (7)
SgI (6)

1 GGATCTCGATCGCTCCGGTCCCCGTAGTGGCAGAGCGCACATGCCACAGTCCCAGAGAAGTTGGGGGAGGGTCGGCAATTGAACCGGTGCTA

101 GAGAAAGTGGCGGGGAAACTGGGAAAGTGTGATGCTGTACTGGCTCCGCTTTCCGAGGGTGGGGAGAACGTATAAGTGCAGTAGTCGC

Psp1406I (203) **HindIII (245)** **Bsu36I (291)**

201 GTGAACGTTCTTTCGCAACGGTTGCCAGAACACAGCTGAAGCTCGAGGGCTCGCATCTCCTCACCGGCCGCCCTACCTGAGGCC

301 GCCATCCACGCCGGTGAGTCGCTCTGCCCTCCGCTGTGGTGCCTCTGAACGTGCTCCCGTAGGTAAGTTAAAGCTAGGTCAGGACC

NgoMI (441)
NaeI (441)

401 GGGCCTTGTCCGGCTCCCTGGAGCCTACCTAGACTCAGCCGCTCTCACGCTTGCTGACCCCTGCTCAACTCTACGTCTTGTCTT

BspEI (558)

501 TCTGTTCTCGCCGTTACAGATCCAAGCTGTGACCGGCCCTACCTGAGATCACCGCTCCGGACAGCATGCCCACTCCAGCCTGCATCCATCCC
11▶ C P R G H G A Q K A A L V L L S A C L V T L W G L G E P P E H T L

Bsp120I (617)

601 GTGTCAGGGGTACGGGGCCAGAACGGCAGCCTGGTCTGCTGAGTGCTGCCCTGGTACCCCTTGGGGCTAGGAGGCCAACAGAGCACACTC
11▶ C P R G H G A Q K A A L V L L S A C L V T L W G L G E P P E H T L

SexAI (704)

701 CGGTACCTGGCTCCACCTAGCTCCCTGAGCTGGACTGCTGTTAACGGGCTGCAGCCTGGTAGGGAGCTGCCACATCCACTCCAGGTACC
45▶ R Y L V L H L A S L Q L G L L N G V C S L A E E L R H I H S R Y

Bsp120I (854)

801 GGGGCAGTACTGGAGGACTGTGGGGCTGCCTGGCTGCCCCCTCCGGCTGGGCTTGTGCTGCTGTCATCTATTCTACTACTCCCTCCAAA
78▶ R G S Y W R T V R A C L G C P L R R G A L L L S I Y F Y Y S L P N

BglIII (995)

901 TCGGGTCGGCCGCCCTCACTGGATGCTTGCCTCCCTGGCCTCTCGAGGCACTGAACATCCTCTGGCCCAAGGGCTGGCCAGCTGAGATC
111▶ A V G P P F T W M L A L L G L S Q A L N I L L G L K G L A P A E I

NcoI (1035) **EcoRV (1064)**

1001 TCTGAGTGTGAAAAAGGAATTCAACGTGGCCATGGCTGGCATGGTATATTACATCGGATATCTGGCTGATCTGCCAGAGCTCAGGCC
145▶ S A V C E K G N F N V A H G L A W S Y Y I G Y L R L I L P E L Q A

BstBI (1103)

1101 GGATTGAACTTACAATCAGCATTACAACACTGCTACGGGTGAGCTGAGCCAGCGCTGTATAATCTCTCCATTGGACTGTGGGTGCTGATAA
178▶ R I R T Y N Q H Y N N L L R G A V S Q R L Y N L L P L D C G V P D N

AgeI (1252)

1201 CCTGAGTATGGCTGACCCAACATTGCTTCTGGATAAACTGCCAGACGGGTGACCGCTGGCATCAAGGATGGTTACAGAACAGCATC
211▶ L S M A D P N I R F L D K L P Q Q T G D R A G I K D R V Y S N S I

1301 TATGAGCTCTGGAGAACGGCAGCGGGCGGGCACCTGTGCTGGAGTACGCCACCCCTTGAGACTTGTGCTGGCATGTACAATACAGTCAGTG
245▶ Y E L L E N G Q R A G T C V L E Y A T P L Q T L F A M S Q Y S Q A

1401 GCTTAGCCGGAGGATAGGCTTGAGCAGGCCAAACTCTCTGGACACTTGAGGACATCTGGAGATGCCCTGAGTCAGAACACTGCCCT
278▶ G F S R E D R L E Q A K L F C R T L E D I L A D A P E S Q N N C R L

1501 CATTGCTTACAGGAACCTGAGATGACAGCAGCTCTCGCTGCTCCAGGAGGTTCTGGCACCTGGCAGGGAGAAAAGGAAGAGGTACTGGC
311▶ I A Y Q E P A D D S S F S L S Q E V L R H L R Q E E K E E V T V G

BpuAI (1605)
BbsI (1605)

1601 AGCTTGAAGACCTAGCGGTGCCAGTACCTCACGATGTCCAAAGAGCTGAGCTCTCATCAGTGAATGGAAAAGCCCTCCCTCTCCGACGGATT
345▶ S L K T S A V P S T S T M S Q E P E L L I S G M E K P L P L R T D

MscI (1745)

1701 TCTCTGAGACCCAGGGTACCGAGGCCAGAGCTCCAGTGTAGCTGGCCAGACATGATAAGATACTTGTGAGGTTGGACAAACACAACAGTAATGC
378▶ F S •

HpaI (1877) **MfeI (1888)**

1801 AGTAAAAAAATGTTATTGTAAATTGTATGGCTATTGTTATTGTAACCATTAAAGCTGAATAAACAGTTAACACAACATGCAATTGATTCA

EcoRI (1973)

1901 TTTATGTTCAAGTTCAAGGGGAGGTGGAGGTTTAAAGCAAGTAAACCTCTACAAATGTGTTATGAAATTCAAATACAGCATAGCAAAAC

2001 TTTAACCTCAAATCAAGCTCTACTGAATCCTTCTGAGGGATGAATAAGGCATAGGCATAGGGCTGTTGCCATGTGATTAGCTGTTGAGC

2101 CTCACCTTCTTCATGGAGTTAACATAGTGTATTCTCAAGGTTGAACAGCTCTTCAATTCTTATGTTAAATGCACTGACCTCCACATT

SspI (2212) **SwaI (2226)**

2201 CCTTTTAGAAAAATTCAAGAACATTAAATACATCATTGCAATGAAATAATGTTTATTAGGCAGAACATCCAGATGCTCAAGGCCCTCATAA

2301 TATCCCCAGTTAGTTAGTGGACTAGGAACAAAGGAACCTTAATAGAAATTGGACAGCAAGAACGGCTAGCTTGTGCTGGTACTT

2401 GAGGGGATGAGTCCTCAATGGGGTTGACAGCTGCCATTCTCAATGAGCACAAAGCAGTCAGGAGCATAGTCAGAGATGAGCTCTGCAC
135◀ L P I L E E I T T K V L K G N M E I L V F C D P A Y D S I L E R C

BstXI (2516)

2501 ATGCCACAGGGTACCCCTGATGGATCTGTCACCTCATCAGAGTAGGGTGCCTGAGCAGCCACAATGGTCAAAGTCCTTGCCGTTGCTA
101◀ M G C P S V V R I S R D V E D S Y P H R V A V I T D F D K Q G N S V

StuI (2651)

2601 CAGCAGACCCAATGCAATGGCTCAGCACAGACAGTGACCCCTGCCAATGTAGGCTCAATGTGGACAGCAGAGATGATCTCCCAGTCAGGTTGCTA
68◀ A S G I A I A E A C V T V R G I Y A E I H V A S I I E G T K T R I

BpuAI (2797)
BbsI (2797)
XmnI (2793)

2701 GGCGCCCCGACATGGTCTTGTCTCATAGAGCATGGTATTCCTCAGTGGCACCTCACCAAGCTCCAGATCTGCTGAGAGATGTTGAAGGTC
35 A A G V H H K N D E Y L M T I K E T A V E V L E L D Q Q S I N F T
BspHI (2801)
2801 TTCAATGGCCCTCTATAGTGAATCTATGCCATATACTATGCCATATTGTCAAACAGCGTGGATGGCTCCAGCTTAC
1 K M ← ← Asel (2859)
2901 TGACGGTCACTAAACGAGCTCTGTTATAGACCTCCCACCGTACAGCCTACCGCCATTGCGTCAATGGGCGGAGTTGTTACGACATTTGAA

SpeI (3014)

3001 AGTCCCGTTGATTTACTAGTCAAAACAACCTCCATTGACGTCAATGGGTGGAGACTTGGAAATCCCGTGAGTCACACCGCTATCCACGCCATTGAT ← ←

SnaBI (3142)

3101 GTACTGCCAAACCGCATCATGGTAATAGCGATGACTAATACGTAGATGACTGCAAGTAGGAAAGTCCATAAGGTATGTACTGGCATAATGC

NdeI (3247)

3201 CAGGGGGCATTACCGTCATTGACGTCAATAGGGGGTACTTGCATATGATACTTGTACTGCCAAGTGGCAGTTACCGTAAATCTCCA

3301 CCCATTGACGTCAATGGAAAGTCCCTATTGGCTTACTATGGAACATACGTCAATTGACGTCAATGGCGGGGTCGGTGGCGGTAGCCAGGGG

PacI (3433)

SdaI (3425)

SbfI (3425)

PciI (3443)

BspLU1II (3443)

3401 GCCATTACCGTAAGTTATGTAACGCCCTGCAGGTTAATTAAAGAACATGTGAGCAAAGGCCAGCAAAGGCCAGGAACCGTAAAAGGCCGCTGCTGG ← ←

3501 CGTTTTCCATAGGCTCCGCCCCCTGACGAGCATCACAAAATCGACGCTCAAGTCAGAGGTGGCAGAACCGACAGGACTATAAGATACCAGGGTT

3601 TCCCCCTGGAAGCTCCCTCGCGCTCTCTGTTCCGACCCCTGCCCTTACCGGATACCTGTCCGCCCTTCCCTCGGAAGCGTGGCGCTTCTCAT

ApaLI (3757)

3701 AGCTCACGCTGTAGGTATCTCAGTTCGGTGTAGTCGTCGCTCCAAGCTGGCTGTGACGAACCCCCCGTTAGCCGACCGCTGCGCTTACCG

3801 GTAATATCGTCTTGAGTCCAACCCGTAAGACACGACTTATGCCACTGGCAGCAGCCACTGGTAACAGGATTAGCAGAGCGAGGTATGTAGCGGTG

3901 TACAGAGTTCTGAAGTGGGGCTAACTACGGCTACACTAGAACAGTATTGGTATCGCCTGCTGAAGCCAGTTACCTCGGAAAAGAGTT

4001 GGTAGCTCTGATCCGCAAACAAACACCACCGCTGGTAGCGGTGGTTTTGGCAAGCAGCAGATTACGCGCAGAAAAAAAGGATCTCAAGAACATC

EagI (4193)

PacI (4173) SwaI (4182) **NotI (4192)**

4101 CTTTGATCTTTCTACGGGTCTGACGCTCAGTGGAAACGAAACTCACGTTAAGGGATTTGGTCACTGGCTAGTTAATTAACATTAAATCAGCGCCGC

4201 AATAAAATATCTTATTTCTTACATCTGTGTTGGTTGGTGAATCTGAACATACGCTCTCCATCAAACAAACGAAACAAACAAAC

4301 TAGCAAAATAGGCTGTCCCCAGTGCAGTGCAAGTGCAGGTGCCAGAACATTCTATCGAA