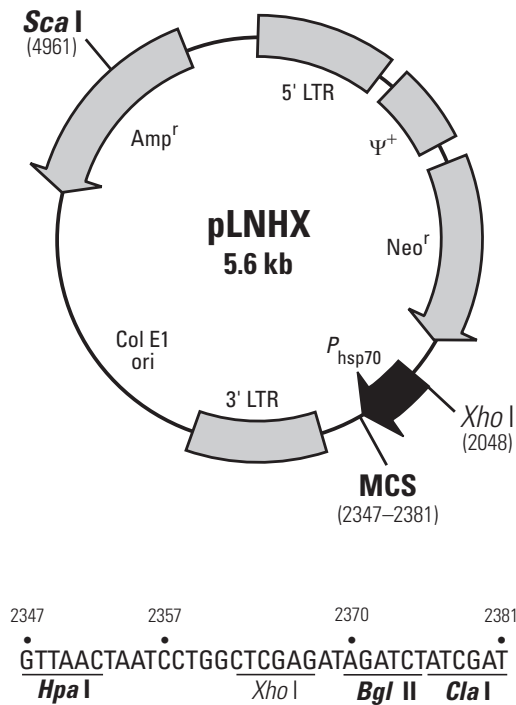


pLNHX Retroviral Vector Information

PT3344-5

GenBank Accession No.: Submission in progress

Sold as part of Catalog No. 631512



Restriction Map and Multiple Cloning Site (MCS) of pLNHX Retroviral Vector. Unique restriction sites are in bold.

Description

pLNHX contains elements derived from Moloney murine leukemia virus (MoMuLV) and Moloney murine sarcoma virus (MoMuSV), and is designed for retroviral gene delivery and expression (1–4). Upon transfection into a packaging cell line, pLNHX can transiently express, or integrate and stably express, a transcript containing the viral packaging signal Ψ^+ , the neomycin selection marker (Neo^r), and a target gene. The 5' viral LTR in this vector contains promoter/enhancer sequences that control expression of the Neo^r gene, which allows antibiotic selection in eukaryotic cells. A target gene can be cloned into the multiple cloning site downstream of the *Drosophila* heat shock protein 70 (hsp70) promoter (P_{hsp70}). pLNHX also includes the Col E1 origin of replication and *E. coli* Amp^r gene for propagation and antibiotic selection in bacteria.

Use

As part of the Pantropic Retroviral Expression System (Cat. No. 631512), pLNHX can be co-transfected with pVSV-G into the GP-293 Packaging Cell Line (5) to produce infectious, replication-*incompetent* retrovirus. pLNHX does not contain the structural genes necessary for viral particle formation and replication. The genes encoding the viral *gag* and *pol* proteins are stably integrated into GP-293. Because the VSV-G envelope protein is toxic, this protein is expressed transiently from pVSV-G (5). Although the virus can infect target cell lines and transmit a target gene, it cannot replicate because the target cell lines lack the viral structural genes. By using the minimal viral sequences and separately introducing the structural genes into the packaging cell line, the chance of producing replication-competent virus due to recombination events is minimized. Alternatively, virus can be produced by transfecting pLNHX into either RetroPack PT67 (Cat. No. 631510) or EcoPack-293 cells (#C3200-1). Note, these packaging cell lines stably express the MoMuLV envelope and thus, do not require pVSV-G.

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Clontech

United States/Canada
800.662.2566

Asia Pacific
+1.650.919.7300

Europe
+33.(0)1.3904.6880

Japan
+81.(0)77.543.6116

Clontech Laboratories, Inc.
A Takara Bio Company
1290 Terra Bella Ave.
Mountain View, CA 94043
Technical Support (US)
E-mail: tech@clontech.com
www.clontech.com

Location of Features

- 5' MoMuSV LTR: 1–588
- Ψ^+ (packaging signal): 657–988
- Neomycin resistance gene (Neo^r):
Start codon: 1067–1069; stop codon: 1859–1861
- *Drosophila* hsp70 promoter (P_{hsp70}): 2047–2330
- Multiple Cloning Site (MCS): 2347–2381
- 3' MoMuLV LTR: 2518–3111
- Col E1 origin of replication:
Site of replication initiation: 3647
- Ampicillin resistance gene (β -lactamase):
Start codon: 5267–5265; stop codon: 4409–4407

Propagation in *E. coli*

- Suitable host strains: DH5 α , HB101, and other general purpose strains.
- Selectable marker: plasmid confers resistance to ampicillin (100 $\mu\text{g/ml}$) to *E. coli* hosts.
- *E. coli* replication origin: Col E1
- Copy number: low

References

1. Coffin, J. M. & Varmus, H. E., Eds. (1996) *Retroviruses* (Cold Spring Harbor Laboratory Press, NY).
2. Ausubel, F. M., et al., Eds. (1995) *Current Protocols in Molecular Biology* (John Wiley & Sons, Inc., NY).
3. Miller, A. D. & Rosman, G. J. (1989) *BioTechniques* 7:980–990.
4. Matsubara, et al. (1996) *Proc. Natl. Acad. Sci. USA* 93:6181–6185.
5. Burns, J. C., et al. (1993) *Proc. Natl. Acad. Sci. USA* 90:8033–8037.

Notes: The viral supernatants produced by this retroviral vector could, depending on your cloned insert, contain potentially hazardous recombinant virus. Due caution must be exercised in the production and handling of recombinant retrovirus. Appropriate NIH, regional, and institutional guidelines apply.

The attached sequence file has been compiled from information in the sequence databases, published literature, and other sources, together with partial sequences obtained by Clontech. This vector has not been completely sequenced

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