pIRES Vector Information

**Description**

pIRES is a mammalian expression vector that allows high level expression of two genes of interest from the same bicistronic mRNA transcript. The vector contains the encephalomyocarditis virus (ECMV) internal ribosome entry site (IRES) flanked by two multiple cloning sites (MCS A and B), an arrangement that allows cap-independent translation of the gene cloned into MCS B (1–3). pIRES utilizes a partially disabled IRES sequence (1) that reduces the rate at which the gene cloned into MCS B is translated relative to that of MCS A.

Expression of the bicistronic transcript is driven by the constitutively active cytomegalovirus immediate early promoter ($P_{\text{CMV IE}}$), located upstream of MCS A. An intervening sequence (IVS) known to enhance the stability of mRNA (4) is located between $P_{\text{CMV IE}}$ and MCS A, and is efficiently spliced out following transcription. SV40 polyadenylation signals downstream of MCS B direct proper processing of the 3' end of the mRNA. Bacteriophage T7 and T3 promoters are located upstream of MCS A and downstream of MCS B, respectively.

pIRES includes a neomycin resistance gene (Neo') to aid in the selection of transfected cells. Neo' is expressed from the SV40 enhancer/promoter, and a synthetic polyadenylation signal directs proper processing of the 3' end of the Neo' mRNA. The SV40 origin allows for replication in mammalian cells expressing the SV40 T antigen. The vector also contains an ampicillin resistance gene (Amp'), and a ColE1 origin of replication for selection and propagation in E. coli, and an f1 origin for single-stranded DNA production.

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Use

Genes cloned into either MCS must contain ATG start codons. pIRES and derivatives can be introduced into mammalian cells by any standard transfection method. Transformed cells can be selected by growth in medium containing the antibiotic G418. Sense or antisense RNA can be transcribed from the T7 and T3 promoters, respectively.

Location of features

- **P<sub>CMV IE</sub>**
  - CMV IE enhancer: 1–659
  - CMV IE promoter: 669–750
- Intervening sequence (IVS): 890–1022
- T7 RNA polymerase promoter: 1067–1085
- Multiple cloning site A: 1085–1107
- IRES sequence: 1130–1725
- Multiple cloning site B: 1737–1763
- T3 RNA polymerase promoter: 1771–1792 (complementary)
- SV40 polyadenylation signal: 1802–2023
- f1 origin of replication: 2118–2573
- Neo<sup>+</sup> expression cassette: 2637–4004
  - SV40 enhancer/early promoter: 2637–3054
  - SV40 origin of replication: 2953–3018
  - Neo<sup>+</sup> structural gene: 3098–3892
    - Start codon (ATG): 3098–3100
    - Stop codon: 3890–3892
  - Synthetic polyadenylation signal: 3956–4004
- Ampicillin resistance (β-lactamase) gene: 4415–5275
  - Start codon (ATG): 4415–4417
  - Stop codon (TAA): 5273–5275

Propagation in *E. coli*

- Suitable host strains: DH5α, HB101, and other general purpose strains.
- Selectable marker: plasmid confers resistance to ampicillin (100 µg/ml) in *E. coli* hosts.
- *E. coli* replication origin: CoIE1
- Copy number: low

References


Note: 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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CMV Sequence: The CMV promoter is covered under U.S. Patent Nos. 5,168,062, and 5,385,839 assigned to the University of Iowa Research Foundation.

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