Vector Map



pLVX-Hom-Mem1



pLVX-Hom-Mem1 vector map and multiple cloning site. To create a fusion of your protein of interest and the DmrB protein, clone your gene of interest in either the 5' or 3' MCS, in-frame with the DmrB coding sequence (see Notes).

Vector Map

pLVX-Hom-Mem1

Cat. No. 635072

Notes

When cloning your gene of interest into the pLVX-Hom-Mem1 vector, make certain that the N-myr-signal is located on the N-terminus of the DmrB fusion protein. The N-myr-signal is NOT functional if localized in the middle or at the C-terminus of a protein of interest. Your gene must either be cloned into the 3'MCS, or if cloned into the 5' MCS, which is located upstream of the N-myr signal, you must include a separate N-myr signal sequence on the forward primer used to amplify your gene of interest.

Location of Features

- 5' LTR (5' long terminal repeat): 1–635
- ψ (packaging signal): 685–822
- RRE (Rev-response element): 1303–1536
- cPPT/CTS (central polypurine tract/central termination sequence): 2028–2151
- P_{CMV} (human cytomegalovirus promoter): 2185–2787
- 5'MCS (5' multiple cloning site): 2803–2820
- N-myr signal (amino-terminal myristoylation signal): 2821–2862
- DmrB (DmrB fusion protein): 2869–3189, 3196–3516
- 3'MCS (3' multiple cloning site): 3518–3531
- IRES (Internal ribosome entry site): 3542–4117
- Puro^r (puromycin resistance gene; puromycin acetyltransferase): 4150–4749
- WPRE(woodchuck hepatitis virus posttranscriptional regulatory element): 4763–5354
- 3' LTR (3' long terminal repeat): 5558 –6194
- pUC ori (pUC origin of replication): 6664–7334(complementary)
- Amp^r (ampicillin resistance gene): 7479–8475 (complementary)

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