PRODUCT: pCMV-Myc & pCMV-HA Vector Set

CATALOG No. 631604

LOT NUMBER Specified on product label.

STORAGE CONDITIONS Store at -20°C

SHELF LIFE

1 year from date of receipt under proper storage conditions.

SHIPPING CONDITIONS

Dry ice (-70°C)

DESCRIPTION

A kit for expression, detection and coimmunoprecipitation of epitope-tagged proteins in mammalian cells. These vectors allow you to easily clone and express a gene of interest fused to an epitope tag. The antibodies to the tags allow detection and purification of the expressed tagged proteins.

PACKAGE CONTENTS

- 10 μg pCMV-Myc-N Vector (500 ng/μl)
- 10 μg pCMV-HA-N Vector (500 ng/μl)
- 500 µl c-Myc Monoclonal Antibody (200 µg/ml)
- 100 µl HA-Tag Polyclonal Antibody (IgG; 1 mg/ml)
- 50 μl pCMV Sequencing Primer (20 μM)

SUGGESTED ANTIBODY DILUTIONS FOR WESTERN BLOTS:

- c-Myc Monoclonal Antibody (1:100)
- HA-Tag Polyclonal Antibody (1:500)

PRIMER SEQUENCE

5'-GAT-CCG-GTA-CTA-GAG-GAA-CTG-AAA-AAC-3' (primer location indicated on Vector Information Packets)

ANTIBODY SOURCES

- c-Myc Monoclonal Antibody was raised in mice against a synthetic peptide (residues 408–439 of the human p62-c-Myc protein).
- HA-Tag Polyclonal Antibody was raised in rabbits against KLH-conjugated synthetic HA epitope (YPYDVPDYA).

PRODUCT USER MANUALS

User manuals for Clontech products are available for download at www.clontech.com/manuals.The following user manuals apply to this product:

- pCMV-Myc-N Vector Information
- pCMV-HA-N Vector Information



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QUALITY CONTROL DATA

See back of page.

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QUALITY CONTROL DATA

1. Plasmid identity and purity

The identity of each plasmid was verified by: a) electrophoresis on an agarose/EtBr gel after digestion with the indicated enzymes, and b) sequence analysis of the epitope tag region. The purity of each plasmid was checked by determining the A_{260}/A_{280} .

Vector	<u>Enzyme</u>	Fragment <u>Size (kb)</u>
pCMV-Myc-N	EcoR I Pst I	3.8 2.7 & 1.1
pCMV-HA-N	EcoR I Pst I	3.8 2.7 & 1.1

2. Antibodies

The c-Myc and HA-Tag antibodies were tested by Western blotting. Cell lysates containing c-Myc- or HA-tagged proteins were electrophoresed on SDS polyacrylamide gels, followed by transfer to PVDF membranes. The blots were probed with 2 μ g/ml c-Myc Monoclonal Antibody or 0.5 μ g/ml HA-Tag Polyclonal Antibody, followed by secondary antibody.



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