

# LADMAC Whole Cell Lysate: sc-364189

## BACKGROUND

Santa Cruz Biotechnology offers a variety of whole cell lysates for use in combination with our antibodies as Western Blotting controls. LADMAC Whole Cell Lysate is derived from the LADMAC cell line using a procedure that ensures protein integrity and lot-to-lot reproducibility. All lysates are tested by Western Blotting to assure that each one contains the expected concentration and assortment of proteins. Numerous antibodies directed against a wide array of mammalian proteins are used to test each lysate.

LADMAC is a transformed cell line derived by transfecting mouse bone marrow cells highly enriched for macrophage progenitors with cloned human cellular Myc-homologous sequences covalently attached to pBR325 (pR Myc). The cell line has monocyte-like morphology; contains nonspecific esterase; is phagocytic for latex beads; secretes lysozyme, and bears the Mac-1 antigen. A minority of cells are Fc receptor positive and an appreciable number of cells are complement receptor 1 positive. The cells are tumorigenic in  $\nu^+$ ,  $\nu^+$  mice but not in syngenic mice. The cells are not phagocytic for antibody or complement-coated particles and do not constitutively secrete Interleukin-1. LADMAC cells secrete the growth factor colony stimulating factor 1 (CSF-1).

## REFERENCES

1. Sklar, M.D., Tereba, A., Chen, B.D. and Walker, W.S. 1985. Transformation of mouse bone marrow cells by transfection with a human oncogene related to c-myc is associated with the endogenous production of macrophage colony stimulating factor 1. *J. Cell. Physiol.* 125: 403-412.
2. Olivas, E., Chen, B.B. and Walker, W.S. 1995. Use of the Pannell-Milstein roller bottle apparatus to produce high concentrations of the CSF-1, the mouse macrophage growth factor. *J. Immunol. Methods* 182: 73-79.

## SOURCE

LADMAC Whole Cell Lysate is derived from the LADMAC cell line.

Organism: *Mus musculus* (mouse)  
 Strain: C<sub>3</sub>H  
 Tissue: Bone marrow  
 Cell Type: Lymphoblast  
 Growth Properties: Suspension, with some loosely adherent cells

## PRODUCT

Each vial contains 500  $\mu$ g protein in 200  $\mu$ l of an SDS-PAGE Western Blotting buffer, which consists of 100  $\mu$ l RIPA Lysis Buffer and 100  $\mu$ l Electrophoresis Buffer, 2X.

## APPLICATIONS

LADMAC Whole Cell Lysate is provided as a Western Blotting positive control. Recommended use is 50  $\mu$ g (20  $\mu$ l) per lane. Sample vial should be boiled once prior to use.

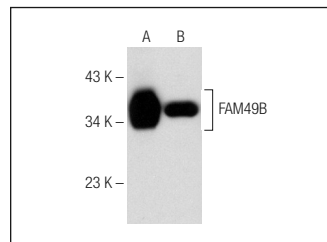
## STORAGE

Store at -20° C; stable for one year from the date of shipment. Non-hazardous. No MSDS required. Minimize repeated freezing and thawing.

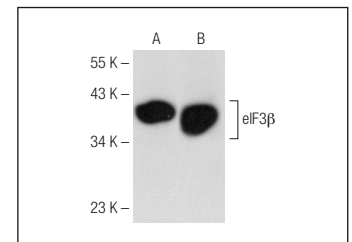
## PREPARATION METHOD

Cells are cultured with appropriate media conditions and allowed to reach a confluency of 75%. Cells are lysed using the RIPA Lysis Buffer System (sc-24948). The BCA Protein Assay Kit (sc-202389) is used to determine the total protein concentration. The lysate is adjusted to contain 500  $\mu$ g of total cellular protein in 100  $\mu$ l before adding an equal volume of Electrophoresis Sample Buffer, 2X (sc-24945). Final concentration of product is 500  $\mu$ g total protein in a final volume of 200  $\mu$ l.

## DATA



FAM49B (N-13): sc-87723. Western blot analysis of FAM49B expression in LADMAC (A) and JAR (B) whole cell lysates.



eIF3 $\beta$  (A-8): sc-374155. Western blot analysis of eIF3 $\beta$  expression in HeLa (A) and LADMAC (B) whole cell lysates.

## RESEARCH USE

For research use only, not for use in diagnostic procedures.

## PROTOCOLS

See our web site at [www.scbt.com](http://www.scbt.com) or our catalog for detailed protocols and support products.