c4 Whole Cell Lysate: sc-364186



The Power to Question

BACKGROUND

Santa Cruz Biotechnology offers a variety of whole cell lysates for use in combination with our antibodies as Western Blotting controls. c4 Whole Cell Lysate is derived from the c4 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.

The c4 cell line lacks functional aryl hydrocarbon receptor nuclear translocator protein (ARNT), due to a point mutation (Gly 326 to Asp) in the ARNT gene. ARNT is directly involved in the regulation of xenobiotic metabolism (including chemical carcinogenesis), hypoxia and differentiation during embryogeneses.

REFERENCES

- Hankinson, O. 1980. Mutants of cultured hepatoma cells deficient in aryl hydrocarbon hydroxylase. In Hankinson, O. Microsomes, drug oxidations, and chemical carcinogenesis. New York: Academic Press, 1149-1152.
- Hoffman, E.C., Reyes, H., Chu, F.F., Sander, F., Conley, L.H., Brooks, B.A. and Hankinson, O. 1991. Cloning of a factor required for activity of the Ah (dioxin) receptor. Science 252: 954-958.
- Numayama-Tsuruta, K., Kobayashi, A., Sogawa, K. and Fujii-Kuriyama, Y. 1997. A point mutation responsible for defective function of the arylhydrocarbon-receptor nuclear translocator in mutant Hepa-1c1c7 cells. Eur. J. Biochem. 246: 486-495.

SOURCE

c4 Whole Cell Lysate is derived from the c4 cell line.

Organism: Mus musculus (mouse)

Strain: C57L/J
Tissue: Liver
Disease: Hepatoma
Cell Type: Epithelial
Growth Properties: Adherent

PRODUCT

Each vial contains 500 μg protein in 200 μl of an SDS-PAGE Western Blotting buffer, which consists of 100 μl RIPA Lysis Buffer and 100 μl Electrophoresis Buffer, 2X.

APPLICATIONS

c4 Whole Cell Lysate is provided as a Western Blotting positive control. Recommended use is 50 μg (20 $\mu l)$ per lane. Sample vial should be boiled once prior to use.

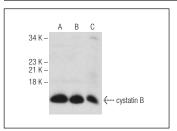
STORAGE

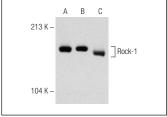
Store at -20° C; stable for one year from the date of shipment. Non-hazardous. No MSDS required. Minimize repeated freezing and thawing.

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 μg of total cellular protein in 100 μ l before adding an equal volume of Electrophoresis Sample Buffer, 2X (sc-24945). Final concentration of product is 500 μg total protein in a final volume of 200 μ l.

DATA





cystatin B (M-40): sc-66854. Western blot analysis of cystatin B expression in c4 (**A**) and EOC 20 (**B**) whole cell lysates and mouse lymph node tissue extract (**C**).

Rock-1 (G-6): sc-17794. Western blot analysis of Rock-1 expression in KNRK ($\bf A$) and c4 ($\bf B$) whole cell lysates and mouse liver tissue extract ($\bf C$).

RESEARCH USE

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

PROTOCOLS

See our web site at www.scbt.com or our catalog for detailed protocols and support products.

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