

PRAM-1 (h): 293T Lysate: sc-114643

BACKGROUND

Complete remission of acute promyelocytic leukemia can be achieved by treating patients with retinoic acid, and PML-RAR α (promyelocytic leukemia-retinoic acid receptor α fusion protein) plays a major role in mediating retinoic acid effects in leukemia cells. The retinoic acid-induced gene, PRAM-1 (PML-RAR α target gene encoding an adaptor molecule 1) encodes an adaptor protein which is expressed and modulated during normal human myelopoiesis. PRAM-1 expression is hindered by expression of PML-RAR α . The 718 amino acid PRAM-1 protein contains 8 N-terminal proline-rich repeats and several proline residues that are clustered as type I or type II SH3 recognition motifs. PRAM-1 demonstrates expression in hematopoietic tissues and lung.

REFERENCES

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2. Online Mendelian Inheritance in Man, OMIM™. 2001. Johns Hopkins University, Baltimore, MD. MIM Number: 606466. World Wide Web URL: <http://www.ncbi.nlm.nih.gov/omim/>
3. Clemens, R.A., et al. 2004. PRAM-1 is required for optimal integrin-dependent neutrophil function. *Mol. Cell. Biol.* 24: 10923-10932.
4. Denis, F.M., et al. 2005. PRAM-1 potentiates arsenic trioxide-induced JNK activation. *J. Biol. Chem.* 280: 9043-9048.
5. Heuer, K., et al. 2006. Lipid-binding HSH3 domains in immune cell adapter proteins. *J. Mol. Biol.* 361: 94-104.
6. Susic, D., et al. 2006. Cardiovascular effects of nonproteolytic activation of prorenin. *Hypertension* 48: E113.
7. Ghaffari, S.H., et al. 2006. Real-time PCR analysis of PML-RAR α in newly diagnosed acute promyelocytic leukaemia patients treated with arsenic trioxide as a front-line therapy. *Ann. Oncol.* 17: 1553-1559.
8. Asou, N., et al. 2007. A randomized study with or without intensified maintenance chemotherapy in patients with acute promyelocytic leukemia who have become negative for PML-RAR α transcript after consolidation therapy: the Japan Adult Leukemia Study Group (JALSG) APL97 study. *Blood* 110: 59-66.
9. Kitareewan, S., et al. 2007. Lysosomes and trivalent arsenic treatment in acute promyelocytic leukemia. *J. Natl. Cancer Inst.* 99: 41-52.

CHROMOSOMAL LOCATION

Genetic locus: PRAM1 (human) mapping to 19p13.2.

PRODUCT

PRAM-1 (h): 293T Lysate represents a lysate of human PRAM-1 transfected 293T cells and is provided as 100 μ g protein in 200 μ l SDS-PAGE buffer.

STORAGE

Store at -20° C. Repeated freezing and thawing should be minimized. Sample vial should be boiled once prior to use. Non-hazardous. No MSDS required.

APPLICATIONS

PRAM-1 (h): 293T Lysate is suitable as a Western Blotting positive control for human reactive PRAM-1 antibodies. Recommended use: 10-20 μ l per lane.

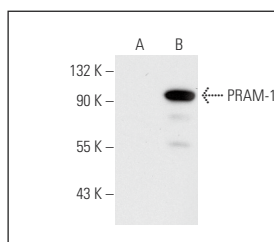
Control 293T Lysate: sc-117752 is available as a Western Blotting negative control lysate derived from non-transfected 293T cells.

PRAM-1 (H-10): sc-166395 is recommended as a positive control antibody for Western Blot analysis of enhanced human PRAM-1 expression in PRAM-1 transfected 293T cells (starting dilution 1:100, dilution range 1:100-1:1,000).

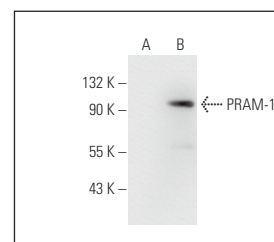
RECOMMENDED SUPPORT REAGENTS

To ensure optimal results, the following support reagents are recommended:
1) Western Blotting: use m-IgG κ BP-HRP: sc-516102 or m-IgG κ BP-HRP (Cruz Marker): sc-516102-CM (dilution range: 1:1000-1:10000), Cruz Marker™ Molecular Weight Standards: sc-2035, UltraCruz® Blocking Reagent: sc-516214 and Western Blotting Luminol Reagent: sc-2048.

DATA



PRAM-1 (H-10): sc-166395. Western blot analysis of PRAM-1 expression in non-transfected: sc-117752 (A) and human PRAM-1 transfected: sc-114643 (B) 293T whole cell lysates.



PRAM-1 (D-11): sc-166267. Western blot analysis of PRAM-1 expression in non-transfected: sc-117752 (A) and human PRAM-1 transfected: sc-114643 (B) 293T whole cell lysates.

RESEARCH USE

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

PROTOCOLS

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