RARα (C-1): sc-515796



The Power to Ouestin

BACKGROUND

Retinoids (RA) are metabolites of vitamin A (retinol) that are important signaling molecules during vertebrate development and tissue differentiation. RAs activate the retinoic acid receptor (RAR) and retinoid X receptor (RXR) nuclear transcription factor families. Most retinoid forms activate RAR family members, whereas RXR family members are activated by 9-cis-RA only. RAR family members, which include RARα, RARβ and RARγ, have a high affinity for all transretinoic acids and belong to the same class of nuclear transcription factors as thyroid hormone receptors, vitamin D₃ receptor and ecdysone receptor. RAR isoforms are expressed in distinct patterns throughout development and in the mature organism. The human RAR α gene maps to chromosome 17q21.2 and is implicated in the chromosomal translocation associated with acute promyelocytic leukemia (APL-M3). Specifically, the RARlphagene is fused with the promyelocytic leukemia (PML) gene, which encodes the fusion protein PML/RAR α . The PML/RAR α fusion protein inhibits PMLdependent apoptotic pathways and halts myeloid differentiation at the promyelocytic stage.

CHROMOSOMAL LOCATION

Genetic locus: RARA (human) mapping to 17q21.2; Rara (mouse) mapping to 11 D.

SOURCE

RAR α (C-1) is a mouse monoclonal antibody raised against amino acids 63-362 mapping within an internal region of RAR α of human origin.

PRODUCT

Each vial contains 200 $\mu g \ lgG_1$ kappa light chain in 1.0 ml of PBS with < 0.1% sodium azide and 0.1% gelatin.

RAR α (C-1) is available conjugated to agarose (sc-515796 AC), 500 μ g/0.25 ml agarose in 1 ml, for IP; to HRP (sc-515796 HRP), 200 μ g/ml, for WB, IHC(P) and ELISA; to either phycoerythrin (sc-515796 PE), fluorescein (sc-515796 FITC), Alexa Fluor® 488 (sc-515796 AF488), Alexa Fluor® 546 (sc-515796 AF546), Alexa Fluor® 594 (sc-515796 AF594) or Alexa Fluor® 647 (sc-515796 AF647), 200 μ g/ml, for WB (RGB), IF, IHC(P) and FCM; and to either Alexa Fluor® 680 (sc-515796 AF680) or Alexa Fluor® 790 (sc-515796 AF790), 200 μ g/ml, for Near-Infrared (NIR) WB, IF and FCM.

APPLICATIONS

RAR α (C-1) is recommended for detection of RAR α of mouse, rat and human origin by Western Blotting (starting dilution 1:100, dilution range 1:100-1:1000), immunoprecipitation [1-2 μ g per 100-500 μ g of total protein (1 ml of cell lysate)], immunofluorescence (starting dilution 1:50, dilution range 1:50-1:500) and solid phase ELISA (starting dilution 1:30, dilution range 1:30-1:3000).

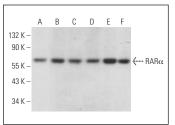
Suitable for use as control antibody for RAR α siRNA (h): sc-29465, RAR α siRNA (m): sc-36393, RAR α shRNA Plasmid (h): sc-29465-SH, RAR α shRNA Plasmid (m): sc-36393-SH, RAR α shRNA (h) Lentiviral Particles: sc-29465-V and RAR α shRNA (m) Lentiviral Particles: sc-36393-V.

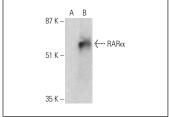
Molecular Weight of RARα: 52 kDa.

STORAGE

Store at 4° C, **DO NOT FREEZE**. Stable for one year from the date of shipment. Non-hazardous. No MSDS required.

DATA





RAR α (C-1): sc-515796. Western blot analysis of RAR α expression in HeLa (A), K-562 (B), HL-60 (C), RAW 264.7 (D), NIH/3T3 (E) and F9 (F) whole cell lysates.

RAR α (C-1): sc-515796. Western blot analysis of RAR α expression in non-transfected: sc-117752 (**A**) and mouse RAR α transfected: sc-125890 (**B**) 293T whole cell Ivsates

SELECT PRODUCT CITATIONS

- Vanderhoeven, F., et al. 2018. Synergistic antitumor activity by combining trastuzumab with retinoic acid in HER2 positive human breast cancer cells. Oncotarget 9: 26527-26542.
- Hai, Y., et al. 2019. Realgar transforming solution-induced differentiation of NB4 cell by the degradation of PML/RARα partially through the ubiquitinproteasome pathway. Arch. Pharm. Res. 42: 684-694.
- Danzl, K., et al. 2019. Early inhibition of endothelial retinoid uptake upon myocardial infarction restores cardiac function and prevents cell, tissue, and animal death. J. Mol. Cell. Cardiol. 126: 105-117.
- 4. Apaya, M.K., et al. 2020. Deregulating the CYP2C19/epoxy-eicosatrienoic acid-associated FABP4/FABP5 signaling network as a therapeutic approach for metastatic triple-negative breast cancer. Cancers 12: 199.
- 5. Zhang, T., et al. 2020. CUDC-101 overcomes arsenic trioxide resistance via caspase-dependent promyelocytic leukemia-retinoic acid receptor α degradation in acute promyelocytic leukemia. Anticancer Drugs 31: 158-168.
- Emde, B., et al. 2020. Microfluidic-based detection of AML-specific biomarkers using the example of promyelocyte leukemia. Int. J. Mol. Sci. 21: 8942.
- Tang, D., et al. 2021. Pontin functions as a transcriptional co-activator for retinoic acid-induced HOX gene expression. J. Mol. Biol. 433: 166928.
- 8. Kamada, S., et al. 2021. Functional inhibition of cancer stemness-related protein DPP4 rescues tyrosine kinase inhibitor resistance in renal cell carcinoma. Oncogene 40: 3899-3913.

RESEARCH USE

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

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