

## Basic Information

Product Name	Anti-RXRA Antibody (Clone#5E7)
Gene Name	RXRA
Source	Mouse
Clonality	Monoclonal
Isotype	IgG1
Species Reactivity	human, mouse, rat
Tested Application	WB, FCM
Contents	500 ug/ml antibody with PBS, 0.02% NaN <sub>3</sub> , 1 mg/ml BSA and 50% glycerol.
Immunogen	E. coli-derived human RXRA recombinant protein (Position: A226-T462).
Concentration	500 ug/ml
Purification	protein G purified.
Observed MW	51-55 kDa
Dilution Ratios	Western blot (WB): 1:500-2000 Flow Cytometry (Fixed):1:50-200

## Storage

12 months from date of receipt, -20°C as supplied.

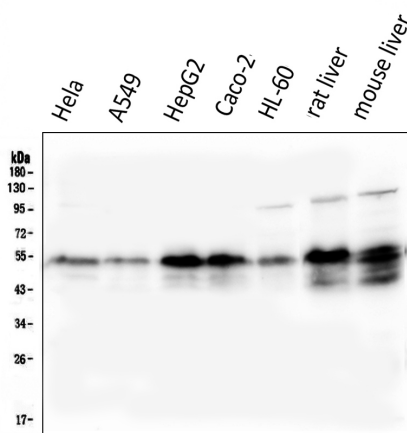
## Background Information

Retinoid X receptor alpha (RXR-alpha), also known as NR2B1 (nuclear receptor subfamily 2, group B, member 1) is a nuclear receptor that in humans is encoded by the RXRA gene. Retinoid X receptors (RXRs) and retinoic acid receptors (RARs) are nuclear receptors that mediate the biological effects of retinoids by their involvement in retinoic acid-mediated gene activation. These receptors function as transcription factors by binding as homodimers or heterodimers to specific sequences in the promoters of target genes. The protein encoded by this gene is a member of the steroid and thyroid hormone receptor superfamily of transcriptional regulators. Alternative splicing of this gene results in multiple transcript variants.

## Reference

Anti-RXRA Antibody (Clone#5E7)被引用在1文献中。

## Selected Validation Data



Western blot analysis of RXRA using anti-RXRA antibody (M01299). The sample well of each lane was loaded with 30 ug of sample under reducing conditions.

Lane 1: human HeLa whole cell lysates,

Lane 2: human A549 whole cell lysates,

Lane 3: human HepG2 whole cell lysates,

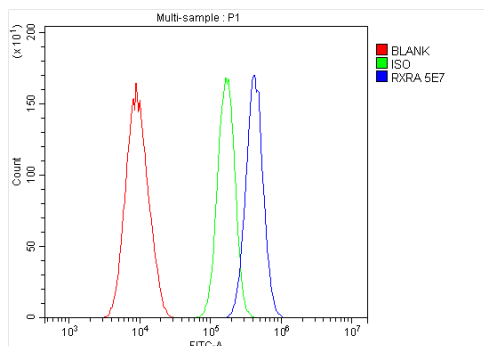
Lane 4: human CACO-2 whole cell lysates,

Lane 5: human HL-60 whole cell lysates,

Lane 6: Rat liver tissue lysates,

Lane 7: Mouse liver tissue lysates.

After electrophoresis, proteins were transferred to a membrane. Then the membrane was incubated with mouse anti-RXRA antigen affinity purified monoclonal antibody (M01299) at a dilution of 1:1000 and probed with a goat anti-mouse IgG-HRP secondary antibody (Catalog # BA1050). The signal is developed using ECL Plus Western Blotting Substrate (Catalog # AR1197). A specific band was detected for RXRA at approximately 51-55 kDa. The expected band size for RXRA is at 51 kDa.



Flow Cytometry analysis of A549 cells using anti-RXRA antibody (M01299).

Overlay histogram showing A549 cells stained with M01299 (Blue line). To facilitate intracellular staining, cells were fixed with 4% paraformaldehyde and permeabilized with permeabilization buffer. The cells were blocked with 10% normal goat serum. And then incubated with mouse anti-RXRA Antibody (M01299) at 1:100 dilution for 30 min at 20°C. DyLight®488 conjugated goat anti-mouse IgG (BA1126) was used as secondary antibody at 1:100 dilution for 30 minutes at 20°C. Isotype control antibody (Green line) was mouse IgG at 1:100 dilution used under the same conditions. Unlabelled sample without incubation with primary antibody and secondary antibody (Red line) was used as a blank control.