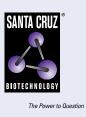
# SANTA CRUZ BIOTECHNOLOGY, INC.

# PPARγ<sub>2</sub> (A-1): sc-166731



## BACKGROUND

Peroxisome proliferator-activated receptors (PPARs) are members of the nuclear hormone receptor subfamily of transcription factors. PPARs form heterodimers with retinoid X receptors (RXRs). These heterodimers regulate transcription of genes involved in Insulin action, adipocyte differentiation, lipid metabolism and inflammation. PPARy is implicated in numerous diseases including obesity, diabetes, atherosclerosis and cancer. PPARy activators include prostanoids, fatty acids, thiazolidinediones and N-(2-benzoylphenyl) tyrosine analogues. A key component in adipocyte differentiation and fatspecific gene expression, PPARy may modulate macrophage functions such as proinflammatory activities, and stimulate oxidized low-den-sity lipoprotein (x-LDL) uptake. A Pro12Ala polymorphism of the PPARy<sub>2</sub> gene has been reported to reduce transactivation activity *in vitro*. This substitution may affect the immune response to ox-LDL and be associated with type 2 diabetes. In addition, the Pro12Ala variant of the PPARy<sub>2</sub> gene maybe correlated with abdominal obesity in type 2 diabetes.

## REFERENCES

- 1. Brun, R.P., et al. 1996. Differential activation of adipogenesis by multiple PPAR isoforms. Genes Dev. 10: 974-984.
- Mansen, A., et al. 1996. Expression of the peroxisome proliferator-activated receptor (PPAR) in the mouse colonic mucosa. Biochem. Biophys. Res. Commun. 222: 844-851.

## CHROMOSOMAL LOCATION

Genetic locus: PPARG (human) mapping to 3p25.2.

### SOURCE

 $PPAR_{Y_2}$  (A-1) is a mouse monoclonal antibody specific for an epitope mapping between amino acids 1-30 at the N-terminus of  $PPAR_{Y_2}$  of human origin.

## PRODUCT

Each vial contains 200  $\mu$ g lgG<sub>2a</sub> kappa light chain in 1.0 ml of PBS with < 0.1% sodium azide and 0.1% gelatin. Also available as TransCruz reagent for Gel Supershift and ChIP applications, sc-166731 X, 200  $\mu$ g/0.1 ml.

PPAR<sub>Y2</sub> (A-1) is available conjugated to agarose (sc-166731 AC), 500  $\mu$ g/ 0.25 ml agarose in 1 ml, for IP; to HRP (sc-166731 HRP), 200  $\mu$ g/ml, for WB, IHC(P) and ELISA; to either phycoerythrin (sc-166731 PE), fluorescein (sc-166731 FITC), Alexa Fluor<sup>®</sup> 488 (sc-166731 AF488), Alexa Fluor<sup>®</sup> 546 (sc-166731 AF546), Alexa Fluor<sup>®</sup> 594 (sc-166731 AF594) or Alexa Fluor<sup>®</sup> 647 (sc-166731 AF647), 200  $\mu$ g/ml, for WB (RGB), IF, IHC(P) and FCM; and to either Alexa Fluor<sup>®</sup> 680 (sc-166731 AF680) or Alexa Fluor<sup>®</sup> 790 (sc-166731 AF790), 200  $\mu$ g/ml, for Near-Infrared (NIR) WB, IF and FCM.

Blocking peptide available for competition studies, sc-166731 P, (100  $\mu$ g peptide in 0.5 ml PBS containing < 0.1% sodium azide and 0.2% stabilizer protein).

Alexa Fluor® is a trademark of Molecular Probes, Inc., Oregon, USA

#### **RESEARCH USE**

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

### **APPLICATIONS**

PPAR<sub>Y2</sub> (A-1) is recommended for detection of PPAR<sub>Y2</sub> of human origin by Western Blotting (starting dilution 1:100, dilution range 1:100-1:1000), immunoprecipitation [1-2  $\mu$ g per 100-500  $\mu$ g of total protein (1 ml of cell lysate)], immunofluorescence (starting dilution 1:50, dilution range 1:50-1:500), immunohistochemistry (including paraffin-embedded sections) (starting dilution 1:50, dilution range 1:50-1:500) and solid phase ELISA (starting dilution 1:30, dilution range 1:30-1:3000); non cross-reactive with PPAR<sub>Y1</sub>.

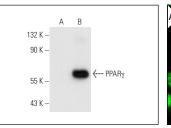
Suitable for use as control antibody for PPAR $_{\gamma_2}$  siRNA (h): sc-29455, PPAR $_{\gamma_2}$  shRNA Plasmid (h): sc-29455-SH and PPAR $_{\gamma_2}$  shRNA (h) Lentiviral Particles: sc-29455-V.

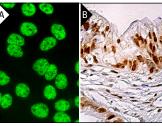
 $\text{PPAR}_{\gamma_2}$  (A-1) X TransCruz antibody is recommended for Gel Supershift and ChIP applications.

Molecular Weight of PPAR $\gamma_2$ : 60 kDa.

Positive Controls: U-937 cell lysate: sc-2239.

#### DATA





 $\begin{array}{l} \text{PPAR}_{\gamma_2} \text{ (A-1): sc-166731. Western blot analysis of} \\ \text{PPAR}_{\gamma} \text{ expression in non-transfected: sc-117752 (A)} \\ \text{and mouse PPAR}_{\gamma} \text{ transfected: sc-122729 (B) 293T} \\ \text{whole cell lysates.} \end{array}$ 

 $\begin{array}{l} PPAR_{Y_2} \ (A-1): \ sc-166731. \ Immunofluorescence \ staining \ of \ formalin-fixed \ Hep \ G2 \ cells \ showing \ nuclear \ localization \ (A). \ Immunoper \ cxide \ staining \ of \ formalin \ fixed, \ paraffin-embedded \ human \ gall \ bladder \ tissue \ showing \ nuclear \ staining \ of \ glandular \ cells \ (B). \end{array}$ 

## SELECT PRODUCT CITATIONS

- Rodriguez, R., et al. 2013. Expression of FUS-CHOP fusion protein in immortalized/transformed human mesenchymal stem cells drives mixoid liposarcoma formation. Stem Cells 31: 2061-2072.
- 2. Li, F., et al. 2021. Upregulated PPARG2 facilitates interaction with demethylated AKAP12 gene promoter and suppresses proliferation in prostate cancer. Cell Death Dis. 12: 528.
- Wang, Y., et al. 2022. AdipoRon exerts opposing effects on Insulin sensitivity via fibroblast growth factor FGF21-mediated time-dependent mechanisms. J. Biol. Chem. 298: 101641.
- Wu, M.C., et al. 2023. Early committed polarization of intracellular tension in response to cell shape determines the osteogenic differentiation of mesenchymal stromal cells. Acta Biomater. 163: 287-301.

#### **STORAGE**

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