### bsm-52220R

- DATASHEET -

## [ Primary Antibody ]

## PPAR gamma Recombinant Rabbit mAb

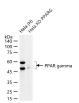


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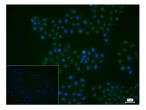
Host: Rabbit	t	Isotype: IgG	Applications: WB (1:500-2000)
Clonality: Recon	nbinant <b>C</b>	CloneNo.: 45G1	Flow-Cyt (1:50-100) ICC/IF (1:50-200)
GenelD: 5468	<b>GenelD:</b> 5468 <b>SWISS:</b> P37231		- ( , ,
Target: PPAR gamma			Reactivity: Human
Immunogen: A synthesized peptide derived from human PPAR gamma: 100-150.			
Purification: affinity purified by Protein A			
Concentration: 1mg/ml			Predicted MW.: <sup>57 kDa</sup>
<b>Storage:</b> 0.01M TBS (pH7.4) with 1% BSA, 0.02% Proclin300 and 50% Glycerol. Shipped at 4°C. Store at -20°C for one year. Avoid repeated freeze/thaw cycles.			Subcellular Location: <sup>Nucleus</sup>
activa	ted receptor (PPAR) subfan	the peroxisome proliferator- nily of nuclear receptors. PPARs X recentors (RXRs) and these	

form heterodimers with retinoid X receptors (RXRs) and these heterodimers regulate transcription of various genes. Three subtypes of PPARs are known: PPAR-alpha, PPAR-delta, and PPARgamma. The protein encoded by this gene is PPAR-gamma and is a regulator of adipocyte differentiation. Additionally, PPAR-gamma has been implicated in the pathology of numerous diseases including obesity, diabetes, atherosclerosis and cancer. Alternatively spliced transcript variants that encode different isoforms have been described. [provided by RefSeq, Jul 2008]

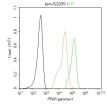
#### - VALIDATION IMAGES -



25 ug total protein per lane of various lysates (see on figure) probed with PPAR gamma monoclonal antibody, unconjugated (bsm-52220R) at 1:1000 dilution and 4°C overnight incubation. Followed by conjugated secondary antibody incubation at r.t. for 60 min.



4% Paraformaldehyde-fixed Hela (H) cell; Triton X-100 at r.t. for 20 min; Antibody incubation with (PPAR gamma) monoclonal Antibody. unconjugated (bsm-52220R) 1:100, 90 min at 37°C; followed by conjugated Goat Anti-Rabbit IgG antibody (green, bs-60295G-BF488) at 37°C for 90 min, DAPI (blue, C02-04002) was used to stain the cell nuclei. PBS instead of the primary antibody was used as the blank control.



The Hela (H) cells were fixed with 4% PFA (10 min at r.t.) and then permeabilized with 90% icecold methanol for 20 min at -20°C.the cells then were incubated in 5%BSA to block non-specific protein-protein interactions (30 min at r.t.). followed by secondary antibody incubation for 40 min at room temperature.Primary Antibody (green):Rabbit Anti-PPAR gamma antibody (bs-52220R,1:100); Isotype Control (orange): Rabbit IgG (bs-0295P). Blank control (black): PBS. Acquisition of 20,000 events was performed.

#### - SELECTED CITATIONS -

- [IF=5.4] Yang Liu. et al. Tuberostemonine may enhance the function of the SLC7A11/glutamate antiporter to restrain the ferroptosis to alleviate pulmonary fibrosis. J ETHNOPHARMACOL. 2023 Aug;:116983 Other ;. 37532076
- [IF=4.8] Guijiang Huang. et al. Osteoking inhibits apoptosis of BMSCs in osteoporotic rats via PI3K/AKT signaling pathway. J ETHNOPHARMACOL. 2024 Dec;:118961 WB ;Rat. 39653105

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