PPAR-γ Mouse mAb

CatalogNo: YM1082   **Orthogonal Validated** ⊙

Key Features

Host Species
• Mouse

Reactivity
• Human,Mouse,Rat,Bovine,Dog,Goat,Pig,Rabbit,sheep

Applications
• WB,IF

MW
• 58kD
(Calculated)

Recommended Dilution Ratios

WB 1:1000-1:2000
IF 1:100-1:500
Not yet tested in other applications.

Storage

Storage*	-15°C to -25°C/1 year(Do not lower than -25°C)
Formulation	Liquid in PBS containing 50% glycerol, 0.5% BSA and 0.02% sodium azide.

Basic Information

Clonality	Monoclonal

Immunogen Information

Immunogen	Purified recombinant human PPAR-γ (C-terminus) protein fragments expressed in E.coli.
Specificity	PPAR-γ Monoclonal Antibody detects endogenous levels of PPAR-γ protein.

Target Information

Gene name	PPARG		

Protein Name	Peroxisome proliferator-activated receptor gamma		

Organism	Gene ID	UniProt ID
Human	5468;	P37231;
Mouse	19016;	P37238;
Rat	25664;	O88275;

Cellular Localization	Nucleus. Cytoplasm. Redistributed from the nucleus to the cytosol through a MAP2K1/MEK1-dependent manner. NOCT enhances its nuclear translocation.
Tissue specificity	Highest expression in adipose tissue. Lower in skeletal muscle, spleen, heart and liver. Also detectable in placenta, lung and ovary.
Function	Alternative products:Additional isoforms seem to exist,Disease:Defects in PPARG are the cause of familial partial lipodystrophy type 3 (FPLD3) [MIM:604367]. Familial partial lipodystrophies (FPLD) are a heterogeneous group of genetic disorders characterized by marked loss of subcutaneous (sc) fat from the extremities. Affected individuals show an increased preponderance of insulin resistance, diabetes mellitus and dyslipidemia.,Disease:Defects in PPARG can lead to type 2 insulin-resistant diabetes and hyptertension.,Disease:Defects in PPARG may be associated with colon cancer.,Disease:Defects in PPARG may be associated with susceptibility to obesity [MIM:601665].,Disease:Variation in PPARG is associated with carotid intimal medial thickness 1 (CIMT1) [MIM:609338]. CIMT is a measure of atherosclerosis that is independently associated with traditional atherosclerotic cardiovascular disease risk factors and coronary atherosclerotic burden. 35 to 45% of the variability in multivariable-adjusted CIMT is explained by genetic factors.,Function:Receptor that binds peroxisome proliferators such as hypolipidemic drugs and fatty acids. Once activated by a ligand, the receptor binds to a promoter element in the gene for acyl-CoA oxidase and activates its transcription. It therefore controls the peroxisomal beta-oxidation pathway of fatty acids. Key regulator of adipocyte differentiation and glucose homeostasis.,online information:Peroxisome proliferator-activated receptor entry,online information:The Singapore human mutation and polymorphism database,polymorphism:Genetic variation in PPARG may influence body mass index (BMI) [MIM:606641]. BMI reflects the amount of fat, lean mass, and body build.,similarity:Belongs to the nuclear hormone receptor family.,similarity:Belongs to the nuclear hormone receptor family. NR1 subfamily.,similarity:Contains 1 nuclear receptor DNA-binding domain.,subunit:Forms a heterodimer with the retinoic acid receptor RXRA called adipocyte-specific transcription factor ARF6. Interacts with NCOA6 coactivator, leading to a strong increase in transcription of target genes. Interacts with coactivator PPARBP, leading to a mild increase in transcription of target genes. Interacts with FAM120B (By similarity). Interacts with NOCA7 in a ligand-inducible manner. Interacts with NCOA1 LXXLL motifs. Interacts with TGFB1I1. Interacts with DNTTIP2.,tissue specificity:Highest expression in adipose tissue. Lower in skeletal muscle, spleen, heart and liver. Also detectable in placenta, lung and ovary.,

Validation Data
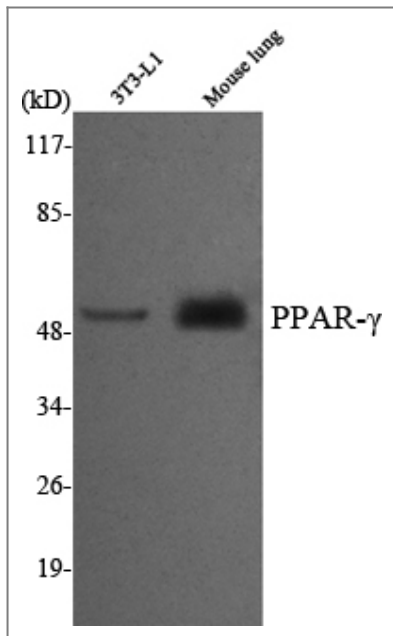
Western Blot analysis using PPAR-γ Monoclonal Antibody against 3T3-L1, mouse lung cell lysate.

| Contact information

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Please scan the QR code to access additional product information:
PPAR-γ Mouse mAb

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Antibody | ELISA Kits | Protein | Reagents