

PPAR-γ Polyclonal Antibody

Catalog No: YT3836

Reactivity: Human; Mouse; Rat

Applications: IF;WB;IHC;ELISA

Target: PPAR-γ

Fields: >>PPAR signaling pathway;>>AMPK signaling pathway;>>Longevity regulating

pathway;>>Osteoclast differentiation;>>Thermogenesis;>>Non-alcoholic fatty liver disease;>>Huntington disease;>>Pathways in cancer;>>Transcriptional

misregulation in cancer;>>Thyroid cancer;>>Lipid and atherosclerosis

Gene Name: PPARG

Protein Name: Peroxisome proliferator-activated receptor gamma

P37231

P37238

Human Gene Id: 5468

Human Swiss Prot

No:

Mouse Gene Id: 19016

Mouse Swiss Prot

No:

Rat Gene ld: 25664

Rat Swiss Prot No: 088275

Immunogen: The antiserum was produced against synthesized peptide derived from human

PPAR-gamma. AA range:78-127

Specificity: PPAR-γ Polyclonal Antibody detects endogenous levels of PPAR-γ protein.

Formulation : Liquid in PBS containing 50% glycerol, 0.5% BSA and 0.02% sodium azide.

Source: Polyclonal, Rabbit, IgG

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Dilution: IF 1:50-200 WB 1:500 - 1:2000. IHC 1:100 - 1:300. ELISA: 1:10000. Not yet

tested in other applications.

Purification: The antibody was affinity-purified from rabbit antiserum by affinity-

chromatography using epitope-specific immunogen.

Concentration: 1 mg/ml

Storage Stability: -15°C to -25°C/1 year(Do not lower than -25°C)

Observed Band: 57kD

Cell Pathway : Protein_Acetylation

Background: peroxisome proliferator activated receptor gamma(PPARG) Homo sapiens This

gene encodes a member of the peroxisome proliferator-activated receptor (PPAR) subfamily of nuclear receptors. PPARs 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 PPAR-gamma. 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],

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

Subcellular Location : Nucleus. Cytoplasm. Redistributed from the nucleus to the cytosol through a MAP2K1/MEK1-dependent manner. NOCT enhances its nuclear translocation.

Expression : Highest expression in adipose tissue. Lower in skeletal muscle, spleen, heart

and liver. Also detectable in placenta, lung and ovary.

Tag: orthogonal

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Sort: 1

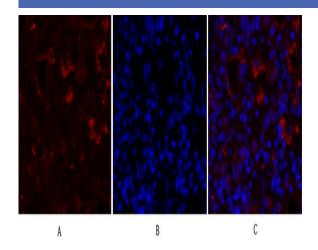
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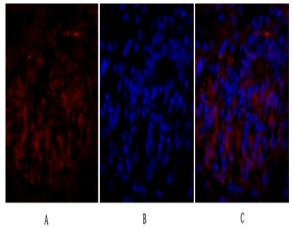
Host: Rabbit

Modifications: Unmodified

Products Images

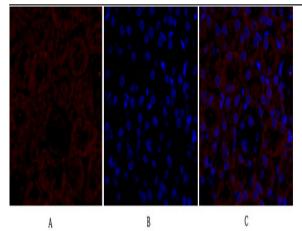


Immunofluorescence analysis of rat-lung tissue. 1,PPAR- γ Polyclonal Antibody(red) was diluted at 1:200(4°C,overnight). 2, Cy3 labled Secondary antibody was diluted at 1:300(room temperature, 50min).3, Picture B: DAPI(blue) 10min. Picture A:Target. Picture B: DAPI. Picture C: merge of A+B

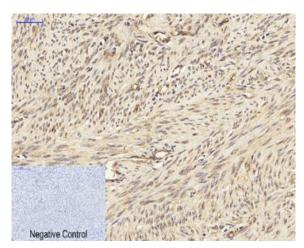


Immunofluorescence analysis of rat-kidney tissue. 1,PPAR-γ Polyclonal Antibody(red) was diluted at 1:200(4°C,overnight). 2, Cy3 labled Secondary antibody was diluted at 1:300(room temperature, 50min).3, Picture B: DAPI(blue) 10min. Picture A:Target. Picture B: DAPI. Picture C: merge of A+B

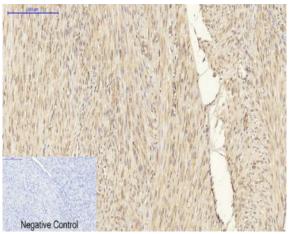
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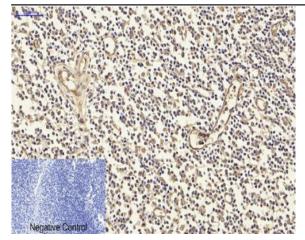
Immunofluorescence analysis of mouse-kidney tissue. 1,PPAR- γ Polyclonal Antibody(red) was diluted at 1:200(4°C,overnight). 2, Cy3 labled Secondary antibody was diluted at 1:300(room temperature, 50min).3, Picture B: DAPI(blue) 10min. Picture A:Target. Picture B: DAPI. Picture C: merge of A+B



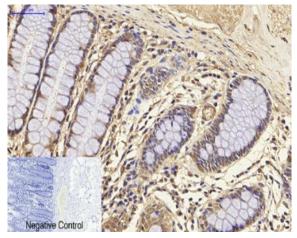
Immunohistochemical analysis of paraffin-embedded Humanuterus tissue. 1,PPAR-γ Polyclonal Antibody was diluted at 1:200(4°C,overnight). 2, Sodium citrate pH 6.0 was used for antibody retrieval(>98°C,20min). 3,Secondary antibody was diluted at 1:200(room tempeRature, 30min). Negative control was used by secondary antibody only.



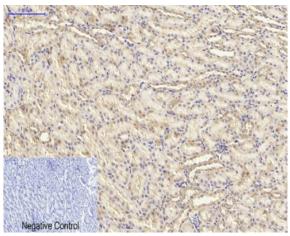
Immunohistochemical analysis of paraffin-embedded Humanuterus-cancer tissue. 1,PPAR- γ Polyclonal Antibody was diluted at 1:200(4°C,overnight). 2, Sodium citrate pH 6.0 was used for antibody retrieval(>98°C,20min). 3,Secondary antibody was diluted at 1:200(room tempeRature, 30min). Negative control was used by secondary antibody only.



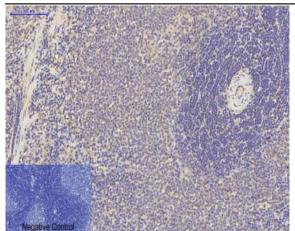
Immunohistochemical analysis of paraffin-embedded Human-Tonsil tissue. 1,PPAR- γ Polyclonal Antibody was diluted at 1:200(4°C,overnight). 2, Sodium citrate pH 6.0 was used for antibody retrieval(>98°C,20min). 3,Secondary antibody was diluted at 1:200(room tempeRature, 30min). Negative control was used by secondary antibody only.



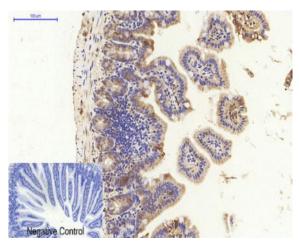
Immunohistochemical analysis of paraffin-embedded Human-colon tissue. 1,PPAR-γ Polyclonal Antibody was diluted at 1:200(4°C,overnight). 2, Sodium citrate pH 6.0 was used for antibody retrieval(>98°C,20min). 3,Secondary antibody was diluted at 1:200(room tempeRature, 30min). Negative control was used by secondary antibody only.



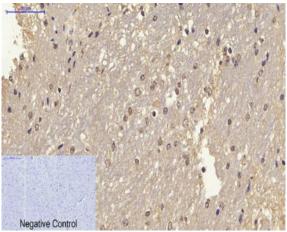
Immunohistochemical analysis of paraffin-embedded Rat-kidney tissue. 1,PPAR-γ Polyclonal Antibody was diluted at 1:200(4°C,overnight). 2, Sodium citrate pH 6.0 was used for antibody retrieval(>98°C,20min). 3,Secondary antibody was diluted at 1:200(room tempeRature, 30min). Negative control was used by secondary antibody only.



Immunohistochemical analysis of paraffin-embedded Rat-spleen tissue. 1,PPAR-γ Polyclonal Antibody was diluted at 1:200(4°C,overnight). 2, Sodium citrate pH 6.0 was used for antibody retrieval(>98°C,20min). 3,Secondary antibody was diluted at 1:200(room tempeRature, 30min). Negative control was used by secondary antibody only.

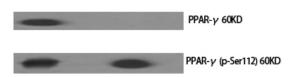


Immunohistochemical analysis of paraffin-embedded Mouse-colon tissue. 1,PPAR-γ Polyclonal Antibody was diluted at 1:200(4°C,overnight). 2, Sodium citrate pH 6.0 was used for antibody retrieval(>98°C,20min). 3,Secondary antibody was diluted at 1:200(room tempeRature, 30min). Negative control was used by secondary antibody only.

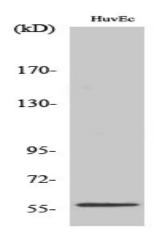


Immunohistochemical analysis of paraffin-embedded Mousebrain tissue. 1,PPAR-γ Polyclonal Antibody was diluted at 1:200(4°C,overnight). 2, Sodium citrate pH 6.0 was used for antibody retrieval(>98°C,20min). 3,Secondary antibody was diluted at 1:200(room tempeRature, 30min). Negative control was used by secondary antibody only.

Western Blot analysis of various cells using PPAR- $\!\gamma$ Polyclonal Antibody diluted at 1:1000



- + phospho-peptide
- + non-phospho-peptide
- + + Paclitaxel (1uM, 24hours)



Western Blot analysis of HuvEc cells using PPAR- γ Polyclonal Antibody diluted at 1:1000