

Recombinant Human PPAR gamma protein ab53382

画像数 1

製品の詳細

製品名	Recombinant Human PPAR gamma protein
精製度	> 95 % SDS-PAGE.
発現系	Escherichia coli
タンパク質長	Full length protein
Animal free	No
由来	Recombinant
生物種	Human
配列	MTMVDTEMPFWPTNFGISSVDLSVMEDHSHSFDIKPFTTVDF SSISTPHY EDIPFTRTDPVVADYKYDLKQLQEYQSAIKVEPASPPYYSEKT QLYNKPHE EPSNSLMAIECRVCGDKASGFHYGVHACEGCKGFFRRTIRLK LIYDRCDL NCRIHKSRNKCQYCRFQKCLAVGMSHNAIRFGRMPQAEKEK LLAEISSD IDQLNPESADLRALAKHLYDSYIKSFPLTKAKARAILTGKTT DKSPFVIY DMNSLMMGEDKIKFKHITPLQEQSKEVAIRIFQGCQFRSVEA VQEITEYA KSIPGFVNLDLNDQVTLKYGVHEIIYTMLASLMNKDGVLI EGQGMTR EFLKSLRKPFGDFMEPKFEFAVKFNALELDDSDLAIFIAVII LSGDRPGL LNVKPIEDIQDNLLQALELQLKLNHPESQLFAKLLQKMTDL RQIVTEHV QLLQVIKKTETDMSLHPLLQEIYKDLY
予測される分子量	59 kDa

特性

Our **Abpromise guarantee** covers the use of **ab53382** in the following tested applications.

The application notes include recommended starting dilutions; optimal dilutions/concentrations should be determined by the end user.

アプリケーション SDS-PAGE

製品の状態

Liquid

前処理および保存

保存方法および安定性

Shipped at 4°C. Upon delivery aliquot and store at -20°C. Avoid freeze / thaw cycles.

pH: 8.00

Constituents: 0.242% Tris, 50% Glycerol

関連情報

機能

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.

組織特異性

Highest expression in adipose tissue. Lower in skeletal muscle, spleen, heart and liver. Also detectable in placenta, lung and ovary.

関連疾患

Note=Defects in PPARG can lead to type 2 insulin-resistant diabetes and hypertension. PPARG mutations may be associated with colon cancer.

Defects in PPARG may be associated with susceptibility to obesity (OBESITY) [MIM:601665]. It is a condition characterized by an increase of body weight beyond the limitation of skeletal and physical requirements, as the result of excessive accumulation of body fat.

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.

Genetic variations in PPARG can be associated with susceptibility to glioma type 1 (GLM1) [MIM:137800]. Gliomas are central nervous system neoplasms derived from glial cells and comprise astrocytomas, glioblastoma multiforme, oligodendrogliomas, and ependymomas.

Note=Polymorphic PPARG alleles have been found to be significantly over-represented among a cohort of American patients with sporadic glioblastoma multiforme suggesting a possible contribution to disease susceptibility.

配列類似性

Belongs to the nuclear hormone receptor family. NR1 subfamily.

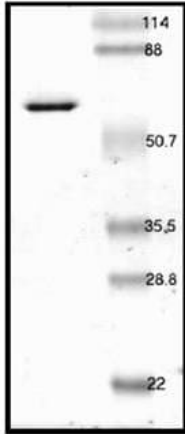
Contains 1 nuclear receptor DNA-binding domain.

細胞内局在

Nucleus.

画像

SDS Page analysis of ab53382



SDS-PAGE - Recombinant Human PPAR gamma protein (ab53382)

Please note: All products are "FOR RESEARCH USE ONLY. NOT FOR USE IN DIAGNOSTIC PROCEDURES"

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