abcam

Product datasheet

HRP Anti-STAT1 alpha antibody [EPYR2154] ab193891



ייבע RabMAb

画像数3

製品の概要

製品名 HRP Anti-STAT1 alpha antibody [EPYR2154]

製品の詳細 HRP Rabbit monoclonal [EPYR2154] to STAT1 alpha

由来種 Rabbit HRP 標識

アプリケーション **適用あり:** WB

種交差性 交差種: Mouse, Human

免疫原 Synthetic peptide. This information is proprietary to Abcam and/or its suppliers.

ポジティブ・コントロール WB: HeLa, HEK293, NIH-3T3 and A431 cell lysates.

特記事項 Our RabMAb® technology is a patented hybridoma-based technology for making rabbit

monoclonal antibodies. For details on our patents, please refer to RabMAb® patents.

製品の特性

製品の状態 Liquid

保存方法 Shipped at 4°C. Store at +4°C short term (1-2 weeks). Store at -20°C. Avoid freeze / thaw cycle.

Store In the Dark.

pH: 7.40 バッファー

Preservative: 0.1% Proclin 300 Solution

Constituents: 30% Glycerol (glycerin, glycerine), 1% BSA, PBS

精製度 Protein A purified

ポリ/モノ モノクローナル クローン名 **EPYR2154**

アイソタイプ ΙgG

アプリケーション

Abpromise保証は、次のテスト済みアプリケーションにおけるab193891の使用に適用されます The Abpromise guarantee アプリケーションノートには、推奨の開始希釈率がありますが、適切な希釈率につきましてはご検討ください。

アプリケーション	Abreviews	特記事項
WB		1/5000. Detects a band of approximately 91 kDa (predicted molecular weight: 87 kDa).

ターゲット情報

機能

Signal transducer and activator of transcription that mediates signaling by interferons (IFNs). Following type I IFN (IFN-alpha and IFN-beta) binding to cell surface receptors, Jak kinases (TYK2 and JAK1) are activated, leading to tyrosine phosphorylation of STAT1 and STAT2. The phosphorylated STATs dimerize, associate with ISGF3G/IRF-9 to form a complex termed ISGF3 transcription factor, that enters the nucleus. ISGF3 binds to the IFN stimulated response element (ISRE) to activate the transcription of interferon stimulated genes, which drive the cell in an antiviral state. In response to type II IFN (IFN-gamma), STAT1 is tyrosine- and serine-phosphorylated. It then forms a homodimer termed IFN-gamma-activated factor (GAF), migrates into the nucleus and binds to the IFN gamma activated sequence (GAS) to drive the expression of the target genes, inducing a cellular antiviral state.

関連疾患

Note=STAT1 deficiency results in impaired immune response leading to severe mycobacterial and viral diseases. In the case of complete deficiency, patients can die of viral disease. Defects in STAT1 are a cause of mendelian susceptibility to mycobacterial disease (MSMD) [MIM:209950]; also known as familial disseminated atypical mycobacterial infection. This rare condition confers predisposition to illness caused by moderately virulent mycobacterial species, such as Bacillus Calmette-Guerin (BCG) vaccine and environmental non-tuberculous mycobacteria, and by the more virulent Mycobacterium tuberculosis. Other microorganisms rarely cause severe clinical disease in individuals with susceptibility to mycobacterial infections, with the exception of Salmonella which infects less than 50% of these individuals. The pathogenic mechanism underlying MSMD is the impairment of interferon-gamma mediated immunity whose severity determines the clinical outcome. Some patients die of overwhelming mycobacterial disease with lepromatous-like lesions in early childhood, whereas others develop, later in life, disseminated but curable infections with tuberculoid granulomas. MSMD is a genetically heterogeneous disease with autosomal recessive, autosomal dominant or X-linked inheritance.

配列類似性

Belongs to the transcription factor STAT family.

Contains 1 SH2 domain.

翻訳後修飾

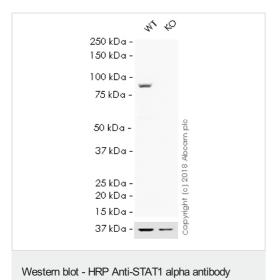
Phosphorylated on tyrosine and serine residues in response to IFN-alpha, IFN-gamma, PDGF and EGF. Phosphorylation on Tyr-701 (lacking in beta form) by JAK promotes dimerization and subsequent translocation to the nucleus. Phosphorylation on Ser-727 by several kinases including MAPK14, ERK1/2 and CAMKII on IFN-gamma stimulation, regulates STAT1 transcriptional activity. Phosphorylation on Ser-727 promotes sumoylation though increasing interaction with PIAS. Phosphorylation on Ser-727 by PKCdelta induces apoptosis in response to DNA-damaging agents.

Sumoylated by SUMO1, SUMO2 and SUMO3. Sumoylation is enhanced by IFN-gamma-induced phosphorylation on Ser-727, and by interaction with PIAS proteins. Enhances the transactivation activity.

ISGylated.

細胞内局在

Cytoplasm. Nucleus. Translocated into the nucleus in response to IFN-gamma-induced tyrosine phosphorylation and dimerization.



[EPYR2154] (ab193891)

All lanes : HRP Anti-STAT1 alpha antibody [EPYR2154] (ab193891) at 1/2000 dilution

Lane 1: Wild-type HAP1 whole cell lysate

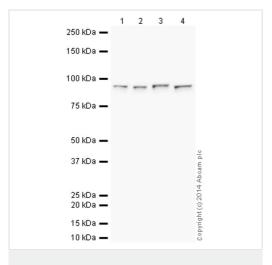
Lane 2: STAT1 knockout HAP1 whole cell lysate

Lysates/proteins at 20 µg per lane.

Predicted band size: 87 kDa **Observed band size:** 91 kDa

Exposure time: 2 minutes

ab193891 was shown to specifically react with STAT1 alpha in wild-type HAP1 cells as signal was lost in STAT1 knockout cells. Wild-type and STAT1 knockout samples were subjected to SDS-PAGE. Ab193891 and ab184095 (Mouse monoclonal [mAbcam 9484] to GAPDH - Loading Control (Alexa Fluor® 680) loading control) were incubated overnight at 4°C at 1/2000 dilution and 1/20000 dilution respectively. The loading control was imaged using the Licor Odyssey CLx prior to blots being developed with ECL technique.



Western blot - HRP Anti-STAT1 alpha antibody [EPYR2154] (ab193891)

All lanes : HRP Anti-STAT1 alpha antibody [EPYR2154] (ab193891) at 1/5000 dilution

Lane 1: HeLa whole cell lysate (ab150035) at 10 µg

Lane 2: HEK293 (Human embryonic kidney cell line) Whole Cell Lysate at 10 μg

Lane 3 : NIH 3T3 (Mouse embryonic fibroblast cell line) Whole Cell Lysate at 20 μg

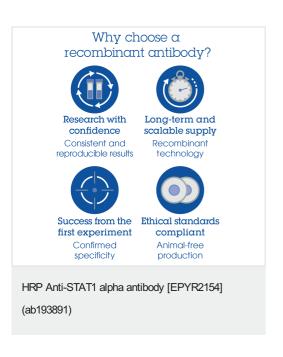
Lane 4: A431 (Human epithelial carcinoma cell line) Whole Cell Lysate at 20 µg

Developed using the ECL technique.

Performed under reducing conditions.

Predicted band size: 87 kDa Observed band size: 91 kDa Exposure time: 2 minutes

This blot was produced using a 4-12% Bis-tris gel under the MOPS buffer system. The gel was run at 200V for 50 minutes before being transferred onto a Nitrocellulose membrane at 30V for 70 minutes. The membrane was then blocked for an hour using 3% milk before being incubated with ab193891 overnight at 4°C. Antibody binding was visualised using ECL development solution **ab133406**.



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