abcam

Product datasheet

Recombinant Mouse CTLA4 protein (Tagged) ab235718

画像数3

製品の詳細

製品名 Recombinant Mouse CTLA4 protein (Tagged)

精製度 > 90 % SDS-PAGE.

発現系 Escherichia coli

アクセッション番号 <u>P09793</u>

タンパク質長 Protein fragment

Animal free No

由来 Recombinant

生物種 Mouse

配列 EAIQVTQPSVVLASSHGVASFPCEYSPSHNTDEVRVTVLRQT

NDQMTEVC

ATTFTEKNTVGFLDYPFCSGTFNESRVNLTIQGLRAVDTGLY

LCKVELMY PPPYFVGMGNGTQIYVIDPEPCPDSD

予測される分子量 34 kDa including tags

領域 36 to 161

配列の追加情報 N-terminal 10xHis-SUMO-tagged and C-terminal Myc-tagged. Extracellular domain.

特性

Our Abpromise guarantee covers the use of ab235718 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

Mass Spectrometry

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. Store at -20°C or -80°C. Avoid freeze / thaw cycle.

pH: 7.2

Constituents: Tris buffer, 50% Glycerol (glycerin, glycerine)

関連情報

機能 Inhibitory receptor acting as a major negative regulator of T-cell responses. The affinity of CTLA4

for its natural B7 family ligands, CD80 and CD86, is considerably stronger than the affinity of their

cognate stimulatory coreceptor CD28.

組織特異性 Widely expressed with highest levels in lymphoid tissues. Detected in activated T-cells where

expression levels are 30- to 50-fold less than CD28, the stimulatory coreceptor, on the cell surface

following activation.

関連疾患 Genetic variation in CTLA4 influences susceptibility to systemic lupus erythematosus (SLE)

[MIM:152700]. SLE is a chronic, inflammatory and often febrile multisystemic disorder of connective tissue. It affects principally the skin, joints, kidneys and serosal membranes. SLE is

thought to represent a failure of the regulatory mechanisms of the autoimmune system.

Note=Genetic variations in CTLA4 may influence susceptibility to Graves disease, an

autoimmune disorder associated with overactivity of the thyroid gland and hyperthyroidism.

Genetic variation in CTLA4 is the cause of susceptibility to diabetes mellitus insulin-dependent

type 12 (IDDM12) [MIM:601388]. A multifactorial disorder of glucose homeostasis that is

characterized by susceptibility to ketoacidosis in the absence of insulin therapy. Clinical fetaures

are polydipsia, polyphagia and polyuria which result from hyperglycemia-induced osmotic diuresis and secondary thirst. These derangements result in long-term complications that affect the eyes,

kidneys, nerves, and blood vessels.

Genetic variation in CTLA4 is the cause of susceptibility to celiac disease type 3 (CELIAC3)

[MIM:609755]. It is a multifactorial disorder of the small intestine that is influenced by both environmental and genetic factors. It is characterized by malabsorption resulting from inflammatory injury to the mucosa of the small intestine after the ingestion of wheat gluten or

related rye and barley proteins. In its classic form, celiac disease is characterized in children by

malabsorption and failure to thrive.

配列類似性 Contains 1 lg-like V-type (immunoglobulin-like) domain.

翻訳後修飾 N-glycosylation is important for dimerization.

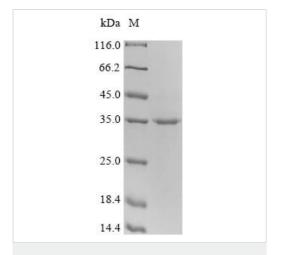
Phosphorylation at Tyr-201 prevents binding to the AP-2 adapter complex, blocks endocytosis,

and leads to retention of CTLA4 on the cell surface.

細胞内局在 Cell membrane. Exists primarily an intracellular antigen whose surface expression is tightly

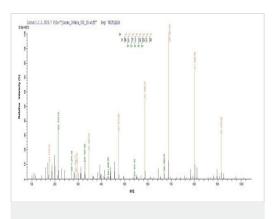
regulated by restricted trafficking to the cell surface and rapid internalisation and.

画像



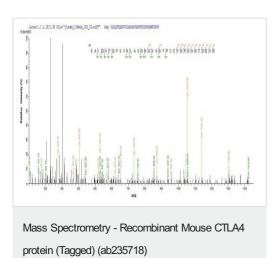
(Tris-Glycine gel) Discontinuous SDS-PAGE (reduced) with 5% enrichment gel and 15% separation gel using ab235718.





Mass Spectrometry - Recombinant Mouse CTLA4 protein (Tagged) (ab235718)

Based on the SEQUEST from database of E.coli host and target protein, the LC-MS/MS Analysis result of ab235718 could indicate that this peptide derived from E.coli-expressed Mus musculus (Mouse) Ctla4.



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