

Recombinant Woodchuck CTLA4 protein (Tagged) ab271482

画像数 1

製品の詳細

製品名	Recombinant Woodchuck CTLA4 protein (Tagged)
精製度	>= 90 % SDS-PAGE.
発現系	HEK 293 cells
アクセッション番号	<u>Q9JLV3</u>
タンパク質長	Protein fragment
Animal free	No
由来	Recombinant
生物種	Woodchuck
予測される分子量	42 kDa
領域	36 to 162
タグ	Avi tag N-Terminus , Fc tag N-Terminus
配列の追加情報	Fc region of Human IgG1.

特性

Our **Abpromise guarantee** covers the use of **ab271482** 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

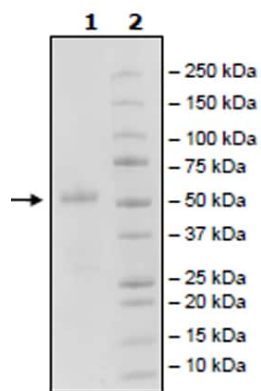
前処理および保存

保存方法および安定性	Shipped on Dry Ice. Store at -80°C. Avoid freeze / thaw cycle. pH: 7.40 Constituents: 0.13% Sodium phosphate, 0.64% Sodium chloride, 0.02% Potassium chloride, 20% Glycerol (glycerin, glycerine)
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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	<p>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.</p> <p>Note=Genetic variations in CTLA4 may influence susceptibility to Graves disease, an autoimmune disorder associated with overactivity of the thyroid gland and hyperthyroidism.</p> <p>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 features 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.</p> <p>Genetic variation in CTLA4 is the cause of susceptibility to celiac disease type 3 (CELAC3) [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.</p>
配列類似性	Contains 1 Ig-like V-type (immunoglobulin-like) domain.
翻訳後修飾	<p>N-glycosylation is important for dimerization.</p> <p>Phosphorylation at Tyr-201 prevents binding to the AP-2 adapter complex, blocks endocytosis, and leads to retention of CTLA4 on the cell surface.</p>
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



SDS-PAGE analysis of 4 µg ab271482.

This protein runs at a higher MW by SDS-PAGE due to glycosylation.

SDS-PAGE - Recombinant Woodchuck CTLA4 protein (Tagged) (ab271482)

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