abcam

Product datasheet

HRP Anti-CTLA4 antibody [KT50] ab106490

1 图像

概述

产品名称 HRP Anti-CTLA4抗体[KT50]

描述 HRP小鼠单克隆抗体[KT50] to CTLA4

宿主 Mouse 偶联物 HRP

经测试应用 适用于: Sandwich ELISA

种属反应性 与反应: Human

免疫原 Recombinant Human CTLA4

常规说明

The Life Science industry has been in the grips of a reproducibility crisis for a number of years.

Abcam is leading the way in addressing this with our range of recombinant monoclonal antibodies and knockout edited cell lines for gold-standard validation. Please check that this product meets

your needs before purchasing.

If you have any questions, special requirements or concerns, please send us an inquiry and/or contact our Support team ahead of purchase. Recommended alternatives for this product can be

found below, along with publications, customer reviews and Q&As

性能

形式 Liquid

存放说明 Shipped at 4°C. Store at +4°C.

存储溶液 Preservative: 0.01% Thimerosal (merthiolate)

Constituent: PBS

纯**度** Protein A purified

 克隆
 单克隆

 克隆编号
 KT50

 同种型
 IgG2a

应用

The Abpromise guarantee Abpromise™承诺保证使用ab106490于以下的经测试应用

"应用说明"部分下显示的仅为推荐的起始稀释度;实际最佳的稀释度/浓度应由使用者检定。

应 用	Ab评论	说 明
Sandwich ELISA		1/1000. Can be paired for Sandwich ELISA with <u>Mouse</u> monoclonal [KT56] to CTLA4 (ab110650).

靶标

功能 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.

组织特异性 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.

疾病相关 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.

序列相似性 Contains 1 lg-like V-type (immunoglobulin-like) domain.

翻译后修饰 N-glycosylation is important for dimerization.

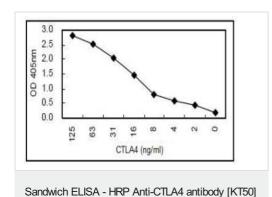
Phosphorylation at Tyr-201 prevents binding to the AP-2 adapter complex, blocks endocytosis,

and leads to retention of CTLA4 on the cell surface.

细胞定位 Cell membrane. Exists primarily an intracellular antigen whose surface expression is tightly

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

图片



ab106490 at 1/1000 dilution detecting CTLA4 by sandwich ELISA. The capture antibody is a mouse monoclonal (lgG1) raised against Human CTLA4.

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

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