abcam

Product datasheet

Mouse Osteoprotegerin ELISA Kit ab100733

7 References 2 图像

概述

产品名称 小鼠Osteoprotegerin ELISA试剂盒

检测方法 Colorimetric

样**品**类型 Cell culture supernatant, Serum, Plasma

检测类型 Sandwich (quantitative)

灵敏度 < 1 pg/ml

范围 1.65 pg/ml - 1200 pg/ml

回收率 87 %

特定样本回收率

样 品类型	平均%	范围
Cell culture supernatant	88.7	81% - 102%
Serum	91.7	82% - 104%
Plasma	81.9	65% - 89%

实验步骤 Multiple steps standard assay

种属反应性 与反应: Mouse

产品概述 Abcam's Osteoprotegerin Mouse ELISA (Enzyme-Linked Immunosorbent Assay) kit is an in vitro

enzyme-linked immunosorbent assay for the quantitative measurement of mouse Osteoprotegerin

in serum, plasma and cell culture supernatants.

This assay employs an antibody specific for mouse Osteoprotegerin coated on a 96-well plate. Standards and samples are pipetted into the wells and Osteoprotegerin present in a sample is bound to the wells by the immobilized antibody. The wells are washed and biotinylated anti-mouse Osteoprotegerin antibody is added. After washing away unbound biotinylated antibody, HRP-conjugated streptavidin is pipetted to the wells. The wells are again washed, a TMB substrate solution is added to the wells and color develops in proportion to the amount of Osteoprotegerin bound. The Stop Solution changes the color from blue to yellow, and the intensity of the color is

measured at 450 nm.

平台 Microplate

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存放说明

Store at -20°C. Please refer to protocols.

组件	1 x 96 tests
200X HRP-Streptavidin Concentrate	1 x 200µl
20X Wash Buffer	1 x 25ml
5X Assay Diluent B	1 x 15ml
Assay Diluent A	1 x 30ml
Biotinylated anti-mouse Osteoprotegerin	2 vials
Osteoprotegerin Microplate (12 x 8 wells)	1 unit
Recombinant Mouse Osteoprotegerin Standard (lyophilized)	2 vials
Stop Solution	1 x 8ml
TMB One-Step Substrate Reagent	1 x 12ml

功能 Acts as decoy receptor for RANKL and thereby neutralizes its function in osteoclastogenesis.

Inhibits the activation of osteoclasts and promotes osteoclast apoptosis in vitro. Bone

homeostasis seems to depend on the local RANKL/OPG ratio. May also play a role in preventing arterial calcification. May act as decoy receptor for TRAIL and protect against apoptosis. TRAIL

binding blocks the inhibition of osteoclastogenesis.

组织**特异性** Highly expressed in adult lung, heart, kidney, liver, spleen, thymus, prostate, ovary, small intestine,

thyroid, lymph node, trachea, adrenal gland, testis, and bone marrow. Detected at very low levels

in brain, placenta and skeletal muscle. Highly expressed in fetal kidney, liver and lung.

疾病相关 Defects in TNFRSF11B are the cause of juvenile Paget disease (JPD) [MIM:239000]; also known

as hyperostosis corticalis deformans juvenilis or hereditary hyperphosphatasia or chronic congenital idiopathic hyperphosphatasia. JPD is a rare autosomal recessive osteopathy that presents in infancy or early childhood. The disorder is characterized by rapidly remodeling woven bone, osteopenia, debilitating fractures, and deformities due to a markedly accelerated rate of

bone remodeling throughout the skeleton. Approximately 40 cases of JPD have been reported worldwide. Unless it is treated with drugs that block osteoclast-mediated skeletal resorption, the

disease can be fatal.

序列相似性 Contains 2 death domains.

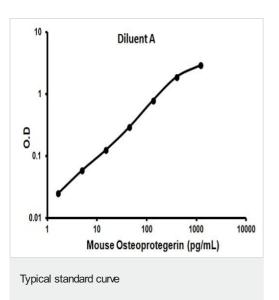
Contains 4 TNFR-Cys repeats.

翻译后修饰 N-glycosylated. Contains sialic acid residues.

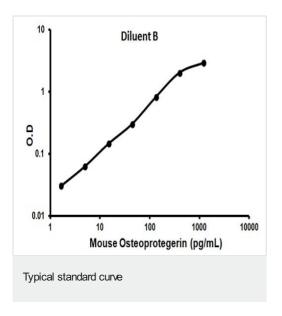
The N-terminus is blocked.

细胞定位 Secreted.

图片



Representative standard curve using ab100733



Representative standard curve using ab100733

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