abcam

Product datasheet

Human MMP13 ELISA Kit ab100605

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製品の概要

製品名 Human MMP13 ELISA Kit

検出方法 Colorimetric

再現性

サンプル	N	平均値	SD	CV%
Overall				< 10%

Inter-Assay(日差再現性)

Intra-Assay(同時再現性)

サンプル	N	平均値	SD	CV%	
Overall				< 12%	

サンプルの種類 Cell culture supernatant, Plasma

アッセイタイプ Sandwich (quantitative)

検出感度 < 6 pg/ml

検出範囲 8.23 pg/ml - 6000 pg/ml

添加回収試験 93 %

特定サンプルでの回収試験

サンプルの種類	平均 %	測定範囲
Cell culture supernatant	94.72	84% - 103%
Plasma	93.13	83% - 103%

ステップ Multiple steps standard assay

種交差性 交差種: Human

製品の概要 Abcam's MMP13 Human ELISA (Enzyme-Linked Immunosorbent Assay) kit is an *in vitro* enzyme linked immunosorbent assay for the quantitative measurement of Human MMP13 in plasma and

cell culture supernatants.

This assay employs an antibody specific for Human MMP13 coated on a 96-well plate. Standards and samples are pipetted into the wells and MMP13 present in a sample is bound to the wells by

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the immobilized antibody. The wells are washed and biotinylated anti-Human MMP13 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 MMP13 bound. The Stop Solution changes the color from blue to yellow, and the intensity of the color is measured at 450 nm.

The kit detects total MMP-13 levels.

We have not been able to detect endogenous Human MMP13 in normal serum with ab100605, only in serum spiked with Human MMP13.

特記事項

Optimisation may be required with urine samples.

試験プラットフォーム

Microplate

製品の特性

保存方法

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

内容	1 x 96 tests
20X Wash Buffer	1 x 25ml
5X Assay Diluent	1 x 15ml
700X HRP-Streptavidin Concentrate	1 x 200µl
Biotinylated anti-Human MMP13	2 vials
MMP13 Microplate (12 x 8 wells)	1 unit
Recombinant Human MMP13 Standard (lyophilized)	2 vials
Stop Solution	1 x 8ml
TMB One-Step Substrate Reagent	1 x 12ml

機能

組織特異性

関連疾患

Degrades collagen type I. Does not act on gelatin or casein. Could have a role in tumoral process. Seems to be specific to breast carcinomas.

Defects in MMP13 are the cause of spondyloepimetaphyseal dysplasia Missouri type (SEMD-MO) [MIM:602111]. A bone disease characterized by moderate to severe metaphyseal changes, mild epiphyseal involvement, rhizomelic shortening of the lower limbs with bowing of the femora and/or tibiae, coxa vara, genu varum and pear-shaped vertebrae in childhood. Epimetaphyseal changes improve with age.

Defects in MMP13 are the cause of metaphyseal anadysplasia type 1 (MANDP1) [MIM:602111]. Metaphyseal anadysplasia consists of an abnormal bone development characterized by severe skeletal changes that, in contrast with the progressive course of most other skeletal dysplasias, resolve spontaneously with age. Clinical characteristics are evident from the first months of life and include slight shortness of stature and a mild varus deformity of the legs. Patients attain a normal stature in adolescence and show improvement or complete resolution of varus deformity of the legs and rhizomelic micromelia.

配列類似性 Belongs to the peptidase M10A family.

Contains 4 hemopexin-like domains.

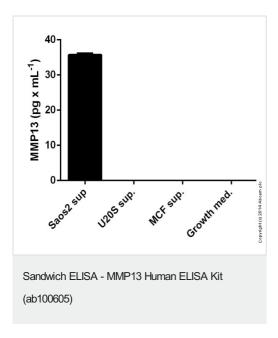
ドメイン The conserved cysteine present in the cysteine-switch motif binds the catalytic zinc ion, thus

inhibiting the enzyme. The dissociation of the cysteine from the zinc ion upon the activation-

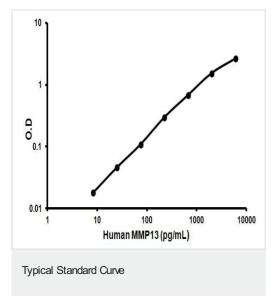
peptide release activates the enzyme.

細胞内局在 Secreted > extracellular space > extracellular matrix.

画像



MMP13 measured in cell supernatanses showing quantity (pg) per mL of tested sample



Representative standard curve using ab100605

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