# abcam

#### Product datasheet

## MMP8 Inhibitor Screening Assay Kit (Colorimetric) ab139452

1 References 4 图像

概述

产品名称 MMP8抑制剂Screening Assay试剂盒(Colorimetric)

检**测方法** Colorimetric

样品类型 Inhibitor compounds

检测类型 Enzyme activity

产品概述 Abcam MMP8 Inhibitor Screening Assay Kit (Colorimetric) (ab139452) is a complete assay

system designed to screen MMP8 inhibitors using a thiopeptide as a chromogenic substrate (Ac-PLG-[2-mercapto-4-methyl-pentanoyl]-LG-OC<sub>2</sub>H<sub>5</sub>). The MMP cleavage site peptide bond is replaced by a thioester bond in the thiopeptide. Hydrolysis of this bond by an MMP produces a sulfhydryl group, which reacts with DTNB [5,5'-dithiobis(2-nitrobenzoic acid), Ellman's reagent] to form 2-nitro-5-thiobenzoic acid, which can be detected by its absorbance at 412 nm (e=13,600 M<sup>-1</sup>).

<sup>1</sup>cm<sup>-1</sup> at pH 6.0 and above). The assays are performed in a convenient 96-well microplate format.

This kit is useful to screen inhibitors of MMP8, a potential therapeutic target. The MMP inhibitor

NNGH is also included as a prototypic control inhibitor.

Thiol inhibitors should not be used with this kit, as they may interfere with the colorimetric assay.

平台 Microplate reader

性能

说明

**存放说明** Please refer to protocols.

组 <b>件</b>	1 x 96 tests
96-well Clear Microplate (1/2 Volume)	1 unit
Colorimetric Assay Buffer	1 x 20ml
MMP Inhibitor	1 x 50µl
MMP Substrate	1 x 50µl
MMP8 Enzyme (Human, Recombinant)	1 x 22µl

组织特异性 Neutrophils.

序列相似性 Belongs to the peptidase M10A family.

Contains 4 hemopexin-like domains.

结构域 The conserved cysteine present in the cysteine-switch motif binds the catalytic zinc ion, thus

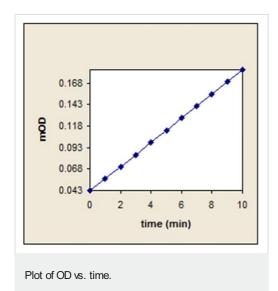
inhibiting the enzyme. The dissociation of the cysteine from the zinc ion upon the activation-

peptide release activates the enzyme.

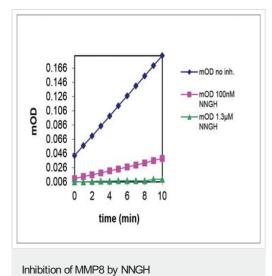
细胞定位 Cytoplasmic granule. Secreted > extracellular space > extracellular matrix. Stored in intracellular

granules.

#### 图片

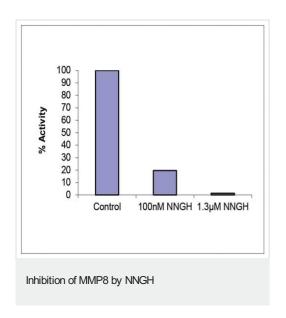


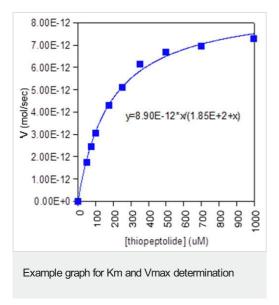
Slope = V = 1.39E-02 OD/min



control slope = 1.39E-02 OD/min inhibitor slope (100nM) = 2.73E-03 OD/min

inhibitor % activity remaining = (2.73E-03/1.39E-02) x 100 = 19.6%





 $Km = 185 \mu M$ Vmax = 8.90 pmol/sec

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