

APA171Hu01 100µg
Active Meprin A Alpha (MEP1a)
Organism Species: *Homo sapiens* (Human)
Instruction manual

FOR RESEARCH USE ONLY
NOT FOR USE IN CLINICAL DIAGNOSTIC PROCEDURES

13th Edition (Revised in Aug, 2023)

[PROPERTIES]

Source: Prokaryotic expression.

Host: *E. coli*

Residues: His213~Val506

Tags: N-terminal His-tag

Purity: >95%

Endotoxin Level: <1.0EU per 1µg (determined by the LAL method).

Buffer Formulation: PBS, pH7.4, containing 0.01% Sarcosyl, 5%Trehalose .

Original Concentration: 200µg/mL

Applications: Activity Assays.

(May be suitable for use in other assays to be determined by the end user.)

Predicted isoelectric point: 6.4

Predicted Molecular Mass: 37.0kDa

Accurate Molecular Mass: 39kDa as determined by SDS-PAGE reducing conditions.

[USAGE]

Reconstitute in 10mM PBS (pH7.4) to a concentration of 0.1-1.0 mg/mL. Do not vortex.

[STORAGE AND STABILITY]

Storage: Avoid repeated freeze/thaw cycles.

Store at 2-8°C for one month.

Aliquot and store at -80°C for 12 months.

Stability Test: The thermal stability is described by the loss rate. The loss rate was determined by accelerated thermal degradation test, that is, incubate the protein at 37°C for 48h, and no obvious degradation and precipitation were observed. The loss rate is less than 5% within the expiration date under appropriate storage condition.

[SEQUENCE]

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                                HYQPFSTN KNASVPTITA KIPEFNSIIG
QRLDFAIDL ERLNRMYNCT THTLLDHCT FEKANICGMI QGTRDDTDWA HQDSAQAGEV
DHTLLGQCTG AGYFMQFSTS SGSAEEAALL ESRILYPERK QQCLQFFYKM TGSPSDRLVV
WVRDDSTGN VRKLVKVQTF QGDDDHNNKI AHVVLKEEQK FRYLFQGTKG DPQNSTGGIY
LDDITLTETP CPTGVWTVRN FSQVLENTSK GDKLQSPRFY NSEGYGFGVT LYPNSRESSG
YLRLAFHVCS GENDAILEWP VENRQV

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[ACTIVITY]

Meprin A alpha (MEP1a) is a metalloprotease belonging to the astacin family, primarily expressed in the kidney, intestine, and immune cells. It plays a crucial role in extracellular matrix remodeling, inflammation, and epithelial barrier function by cleaving substrates such as collagen, cytokines, and adhesion molecules. MEP1a is synthesized as a zymogen and activated through proteolytic processing. It forms heterodimers with Meprin A beta (MEP1b), which stabilizes the complex and modulates substrate specificity. To detect the activity of recombinant MEP1a, a functional ELISA assay was performed to evaluate the interaction between recombinant human IMEP1a and recombinant rat MEP1b. Briefly, MEP1a was diluted serially in PBS with 0.01% BSA (pH 7.4). Duplicate samples of 100 μ l were then transferred to MEP1b-coated microtiter wells and incubated for 1h at 37 °C. Wells were washed with PBST and incubated for 1h with anti-MEP1a pAb, then aspirated and washed 3 times. After incubation with HRP labelled secondary antibody for 1h at 37 °C, wells were aspirated and washed 5 times. With the addition of substrate solution, wells were incubated 15-25 minutes at 37°C. Finally, add 50 μ L stop solution to the wells and read at 450/630nm immediately. The binding activity of recombinant human IMEP1a and recombinant rat MEP1b was

shown in Figure 1, the EC50 for this effect is 0.048µg/mL.

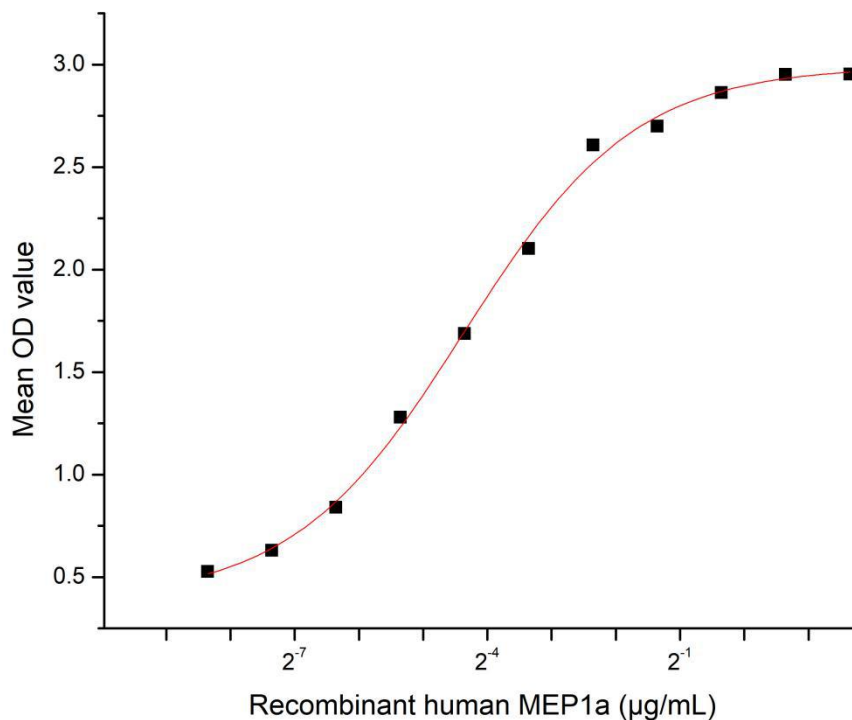


Figure 1. The binding activity of recombinant human IMEP1a and recombinant rat MEP1b

[IDENTIFICATION]

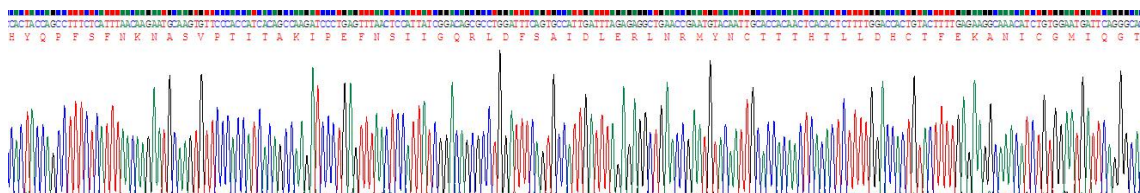


Figure 2. Gene Sequencing (extract)

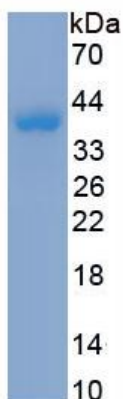


Figure 3. SDS-PAGE

Sample: Active recombinant MEP1a, Human

[IMPORTANT NOTE]

The kit is designed for research use only, we will not be responsible for any issue if the kit was used in clinical diagnostic or any other procedures.