

EPA097Hu61 100µg
Eukaryotic Matrix Metalloproteinase 1 (MMP1)
Organism Species: *Homo sapiens (Human)*
Instruction manual

FOR RESEARCH USE ONLY
NOT FOR USE IN CLINICAL DIAGNOSTIC PROCEDURES

12th Edition (Revised in Aug, 2016)

[PROPERTIES]

Source: Eukaryotic expression

Host: CHO Cell

Residues: Phe20~Asn469

Tags: N-terminal His Tag

Subcellular Location: Extracellular matrix

Purity: > 95%

Traits: Freeze-dried powder

Buffer formulation: PBS, pH7.4, containing 0.01% SKL, 1mM DTT, 5% Trehalose and Proclin300.

Original Concentration: 200µg/mL

Applications: Positive Control; Immunogen; SDS-PAGE; WB.

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

Predicted isoelectric point: 6.9

Predicted Molecular Mass: 53.5kDa

Accurate Molecular Mass: 58&62kDa as determined by SDS-PAGE reducing conditions.

Phenomenon explanation:

The possible reasons that the actual band size differs from the predicted are as follows:

1. Splice variants: Alternative splicing may create different sized proteins from the same gene.
2. Relative charge: The composition of amino acids may affects the charge of the protein.
3. Post-translational modification: Phosphorylation, glycosylation, methylation etc.
4. Post-translation cleavage: Many proteins are synthesized as pro-proteins, and then cleaved to give the active form.
5. Polymerization of the target protein: Dimerization, multimerization etc.

[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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                F PATLETQEQD VDLVQKYLEK YYNLKNDGRQ
VEKRRNSGPV VEKLVQMGEF FGLKVTGKPD AETLKVMKQP RCGVPDVAQF
VLTEGNPRWE QTHLTYRIEN YTPDLPRADV DHAIEKAFQL WSNVTPLTFT
KVSEGOADIM ISFVRGDHRD NSPFDGPGGN LAHAFQPGPG IGGDAHFDDE
ERWTNNFREY NLHRVAAHEL GHSLGLSHST DIGALMYPSTY TFSGDVQLAQ
DDIDGIQAIY GRSQNPVQPI GPQTPKACDS KLTFDAITTI RGEVMFFKDR
FYMRTNPFYP EVELNFISVF WPQLPngleA AYEFADRDEV RFFKGNKYWA
VQGQNVLHGY PKDIYSSFGF PRTVKHIDAA LSEENTGKTY FKVANKYWRY
DEYKRSDMPG YPKMIAHDFP GIGHKVDAVF MKDGFFYFFH GTRQYKFDPK
TKRILTQKA  NSWFNCRKN
  
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[IDENTIFICATION]

TTCTGCGGCTCTCGGACGAGGAGAGCTCTGACTTGTCGAAATCTCTGAAATCTCAACCTCGAGATGTCGGGCGCATGGAACCGGAGCAATGTCGCGCCAGCTGCTGAAATGAGCCAAATTCAGCATTTCTTCCGCTGAAATGCTCTGGAGACGATCTGAAACCCCTGAGCGATGAGCCGCTCGATGCTGCTGCTG
 F PATLETQEQD VDLVQKYLEK YYNLKNDGRQ VEKRRNSGPV VEKLVQMGEF FGLKVTGKPD AETLKVMKQP RCGVPDVAQF

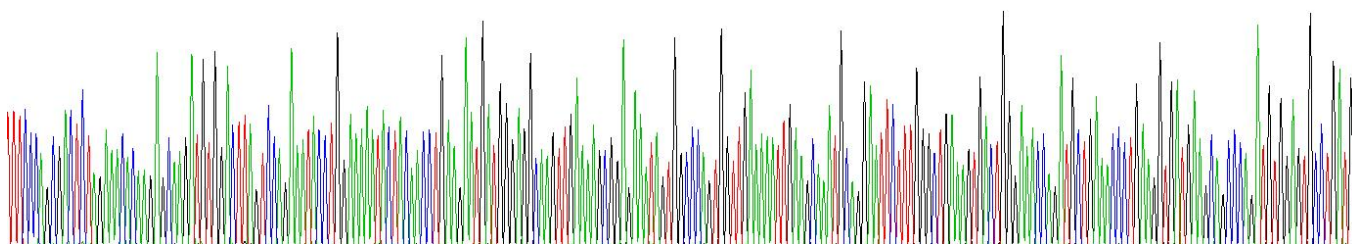


Figure. Gene Sequencing (Extract)

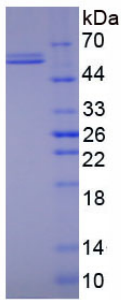


Figure. SDS-PAGE

[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.