

Active Peptidylglycine Alpha Amidating Monooxygenase (PAM)

Catalog No.: TP09174 50µg

Sequence Information

Species: Human Gene ID:5066

Swiss Prot:P19021 Synonyms:PAL; PHM; Peptidylamidoglycolate

lyase; Peptidyl-Alpha-hydroxyglycine

Alpha-amidating Lyase; Peptidylglycine

Alpha-Hydroxylating Monooxygenase

Residues:Phe21~Cys288

FRSPLSVFKRFKETTRPFSNECLGTTRPVVPIDSSDFALDIRMPGVTPKQSDTY

FCMSMRIPVDEEAFVIDFKPRASMDTVHHMLLFGCNMPSSTGSYWFCDEGTCTD

KANILYAWARNAPPTRLPKGVGFRVGGETGSKYFVLQVHYGDISAFRDNNKDCS

GVSLHLTRLPQPLIAGMYLMMSVDTVIPAGEKVVNSDISCHYKNYPMHVFAYRV

HTHHLGKVVSGYRVRNGQWTLIGRQSPQLPQAFYPVGHPVDVSFGDLLAARC

Product Information

Source: Recombinant expression.

Host: E.coli

Tags: N-terminal His-Tag

Subcellular Location: Secreted

Purity: >90%

Traits: Freeze-dried powder

Buffer formulation: 20mM Tris, 150mM NaCl, pH8.0, containing 0.05%sarcosyland 5%

trehalose

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: 8.8

Predicted Molecular Mass: 33.7kDa

Accurate Molecular Mass: 34kDa as determined by SDS-PAGE reducing conditions.

[USAGE]

Reconstitute in ddH₂O to a concentration of 0.1-0.5 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.

[ACTIVITY]

Peptidyl-glycine alpha-amidating monooxygenase (PAM) is an enzymethat isrequired for the biosynthesis of many signaling peptides. This enzymemainlyincludes two domains with distinct catalytic activities, a peptidylglycinealpha-hydroxylating monooxygenase (PHM) domain and a peptidyl-alpha-hydroxyglycine alpha-amidating lyase (PAL) domain. These catalytic domainswork sequentially to catalyze neuroendocrine peptides to active alpha-amidatedproducts. Besides, Glucosidase Alpha, Acid (GaA) has been identifiedasaninteractor of PAM, thus a binding ELISA assay was conducted to detect theinteraction of recombinant human PAM and recombinant human GaA. Briefly, PAM were diluted serially in PBS, with 0.01% BSA (pH 7.4). Duplicate samplesof100uL were then transferred to GaA-coated microtiter wells and incubatedfor 2hat 37 °C. Wells were washed with PBST and incubated for 1h with anti-PAMpAb, then aspirated and washed 3 times. After incubation with HRPlabelledsecondary antibody, wells were aspirated and washed 3 times. With theadditionof substrate solution, wells were incubated 15-25 minutes at 37 °C . Finally, add50µL stop solution to the wells and read at 450nm immediately. Thebindingactivity of PAM and GaA.was shown in Figure 1, and this effect was inadosedependent manner.

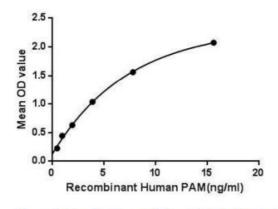


Figure 1. The binding activity of PAM with GaA.

[IDENTIFICATION]



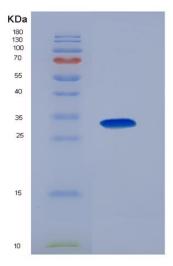


Figure 1. SDS-PAGE

[IMPORTANT NOTE]

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