#### APA431Hu01 10µg Active Granzyme M (GZMM) **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: Ile26~Ala257 Tags: N-terminal His-tag **Purity: >92% Endotoxin Level:** <1.0EU per 1µg (determined by the LAL method). Buffer Formulation: 20mM Tris, 150mM NaCl, pH8.0, containing 1mM EDTA and 5% trehalose. Applications: Cell culture; Activity Assays. (May be suitable for use in other assays to be determined by the end user.) Predicted isoelectric point: 10.3 Predicted Molecular Mass: 26.3kDa Accurate Molecular Mass: 26kDa as determined by SDS-PAGE reducing conditions.

# [USAGE]

Reconstitute in 20mM Tris, 150mM NaCl (pH8.0) 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.

Aliguot 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]

IIGGR EVIPHSRPYM ASLQRNGSHL CGGVLVHPKW VLTAAHCLAQ RMAQLRLVLG LHTLDSPGLT FHIKAAIQHP RYKPVPALEN DLALLQLDGK VKPSRTIRPL ALPSKRQVVA AGTRCSMAGW GLTHQGGRLS RVLRELDLQV LDTRMCNNSR FWNGSLSPSM VCLAADSKDQ APCKGDSGGP LVCGKGRVLA RVLSFSSRVC TDIFKPPVAT AVAPYVSWIR KVTGRSA

## [ACTIVITY]

GZMM (Granzyme M) is one of the neutral serine proteases, which is specifically expressed by NK cells and mediates a novel major and perforin-dependent cell death pathway. Granzyme M has been proven to targets a-Tubulin and disorganizes the microtubule network, besides, Ezrin has also been identified as a substrate of GZMM. Human granzyme M is synthesized as a precursor (264 residues) with a signal peptide (residues 1-23), a propeptide (residues 24-25) and a mature chain (residues 26-257). The purified recombinant human Granzyme M consists of residues 26 to 257 which activity was measured by its ability to cleaves a thioester substrate Z-Lys-SBzI+HCI. The reaction was performed in 0.05 M Tris, 0.15 M NaCl, 0.01% Triton X-100, pH 8.0 (assay buffer), initiated by addition 50  $\mu$ L of various concentrations of GZMM (diluted by assay buffer) to 50  $\mu$ L of 1.2 mM substrate and DTNB mixture. The final well serves as a negative control with no GZMM, replace with 50  $\mu$ L assay buffer. Incubated at 25 °C for 5min, then read at a wavelength of 405 nm. The specific activity of recombinant human Granzyme M is >80 pmol/min/µg.

Specific Activity (pmol/min/ug)=

Adjusted Vmax\* (OD/min) x well volume (L) x 1012 pmol/mol

ext. coeff\*\* (M-1cm-1) x path corr.\*\*\* (cm) x amount of enzyme (ug)

\*Adjusted for Substrate Blank

\*\*Using the extinction coefficient 13800 M<sup>-1</sup>cm<sup>-1</sup>

\*\*\*Using the path correction 0.320 cm

## [IDENTIFICATION]



Figure 1. Gene Sequencing (extract)



Figure 2. SDS-PAGE

Sample: Active recombinant GZMM, Human



Figure 3. Western Blot

Sample: Recombinant GZMM, Human;

Antibody: Rabbit Anti-Human GZMM Ab (PAA431Hu01)

## [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.