#### APA567Ra01 10µg Active S100 Calcium Binding Protein B (S100B) Organism Species: Rattus norvegicus (Rat) Instruction manual

#### FOR RESEARCH USE ONLY NOT FOR USE IN CLINICAL DIAGNOSTIC PROCEDURES

1st Edition (Apr, 2016)

### [PROPERTIES]

Source: Prokaryotic expression. Host: *E. coli* Residues: MSELEKAEF+Met1~Glu92 Tags: N-terminal His-tag Purity: >95% Endotoxin Level: <1.0EU per 1mL (determined by the LAL method). Buffer Formulation: 10mM PBS, pH7.4, containing 5% trehalose. Original Concentration: 800µg/mL Applications: Cell culture; Activity Assays; In vivo assays. (May be suitable for use in other assays to be determined by the end user.) Predicted isoelectric point: 4.5 Predicted Molecular Mass: 13.0kDa Accurate Molecular Mass: 12kDa as determined by SDS-PAGE reducing conditions.

## [ <u>USAGE</u> ]

Reconstitute in  $ddH_2O$  to a concentration of 0-0.1 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.

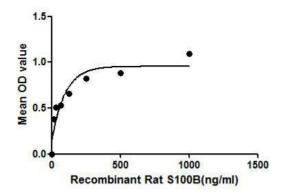
**Stability Test:** The thermal stability is described by the loss rate. The loss rate was determined by accelerated thermal degradation test, that is, incubate the protein at 37°C for 48h, and no obvious degradation and precipitation were observed. The loss rate is less than 5% within the expiration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

# [ <u>SEQUENCE</u> ]

MSELEKAEF MSELEKAMVA LIDVFHQYSG REGDKHKLKK SELKELINNE SHFLEEIKE QEVVDKVMET LDEDGDGECD FQEFMAFVSM VTTACHEFFE HE

## [ACTIVITY]

Protein S100B is a member of the S100 family. S100 proteins are EF-hand calcium-binding proteins and involved in the regulation of a number of cellular processes such as cell cycle progression and differentiation. Experimental results suggest that the receptor for advanced glycation end products (RAGE) plays important roles in mediating S100 protein-induced cellular signaling. Besides, rat RAGE shares similarities with human RAGE in amino acids sequence with the identity of 80.0%. Thus a binding ELISA assay was conducted to detect the interaction of recombinant rat S100B and recombinant human RAGE. Briefly, S100B were diluted serially in PBS, with 0.01%BSA (pH 7.4). Duplicate samples of 100uL were then transferred to RAGE-coated microtiter wells and incubated for 2h at 37°C. Wells were washed with PBST and incubated for 1h with anti-S100B mAb. then aspirated and washed 3 times. After incubation with HRP labelled secondary antibody, wells were aspirated and washed 3 times. With the addition of substrate solution, wells were incubated 15-25 minutes at 37°C. Finally, add 50µL stop solution to the wells and read at 450nm immediately. The binding activity of of S100B and RAGE was shown in Figure 1, and this effect was in a dose dependent manner.





#### [IDENTIFICATION]

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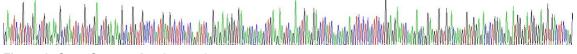
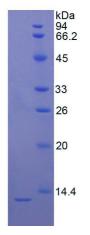
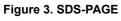


Figure 2. Gene Sequencing (extract)





Sample: Active recombinant S100B, Rat

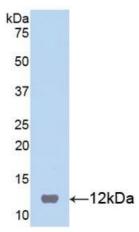


Figure 4. Western Blot Sample: Recombinant S100B, Rat; Antibody: Rabbit Anti-Rat S100B Ab (PAA567Ra01)

# [<u>IMPORTANT NOTE</u>]

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.