

APD228Hu01 100µg
Active Troponin C Type 2, Fast (TNNC2)
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: Thr2~Gln160

Tags: N-terminal His and GST Tag

Purity: >90%

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: 4.7

Predicted Molecular Mass: 47.9kDa

Accurate Molecular Mass: 48kDa 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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TDQQAARS YLSEEMIAEF KAAFDMFDAD GGGDISVKEL GTVMRLGQT  
PTKEELDAII EEVDEDESGT IDFEEFLVMM VRQMKEDAKG KSEEELAECF  
RIFDRNADGY IDPEELAEIF RASGEHVTDE EIESLMKDGD KNNDGRIDFD  
EFLKMMEGVQ
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[ACTIVITY]

Troponin C Type 2, Fast (TNNC2) is a protein predominantly expressed in fast - twitch skeletal muscles. It is a crucial component of the troponin complex. TNNC2 binds calcium ions, which initiates a conformational change in the troponin - tropomyosin complex. This change allows myosin to interact with actin, enabling muscle contraction. Mutations in the TNNC2 gene can lead to various skeletal muscle disorders affecting muscle function. It has been reported that the binding of TNNC2 and TNNT3 plays a key role in the process of rapid skeletal muscle contraction. Thus a functional ELISA assay was conducted to detect the interaction of recombinant human TNNC2 and recombinant human TNNT3.

Briefly, TNNC2 was diluted serially in PBS with 0.01% BSA (pH 7.4). Duplicate samples of 100 μ l were then transferred to TNNT3-coated microtiter wells and incubated for 1h at 37°C. Wells were washed with PBST and incubated for 1h with anti-TNNC2 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 TNNC2 and recombinant human TNNT3 was shown in Figure 1, the EC₅₀ for this effect is 0.003ug/mL.

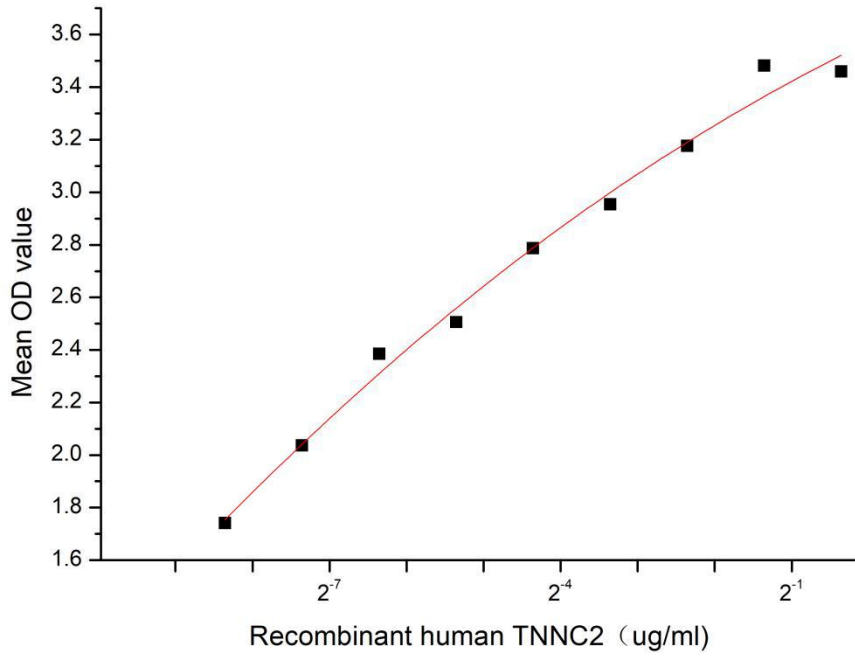


Figure 1. The binding activity of recombinant human TNNC2 and human TNNT3

