

**APP871Hu01 100µg**  
**Active MHC Class I Polypeptide Related Sequence A (MICA)**  
**Organism Species: *Homo sapiens* (Human)**  
***Instruction manual***

FOR RESEARCH USE ONLY  
NOT FOR USE IN CLINICAL DIAGNOSTIC PROCEDURES

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13th Edition (Revised in Aug, 2023)

## **[ PROPERTIES ]**

**Source:** Prokaryotic expression.

**Host:** *E. coli*

**Residues:** Glu24~Trp307

**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:** 6.8

**Predicted Molecular Mass:** 62.7kDa

**Accurate Molecular Mass:** 63kDa 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.

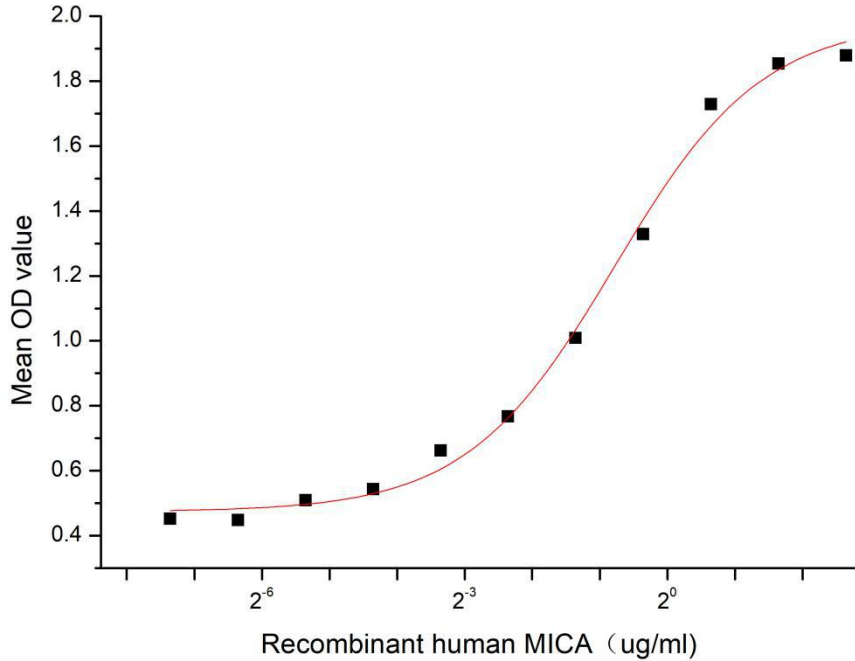
## **[ SEQUENCE ]**

EPHSLRYNLTVLSWDGVSQSGFLTEVHLDGQPFLRCDRQKCRAPQGQWAEDVLGNKTWDR  
ETRDLTGNGKDLRMTLAHIKDQKEGLHSLQEIRVCEIHEDNSTRSSQHFYYDGELFLSQNLETK  
EWTMPQSSRAQTLAMNVRNFLKEDAMKTKTHYHAMHADCLQELRRYLKSGVVLRRTVPPM  
VNVTRSEASEGNITVTCRASGFYPWNITLSWRQDGVSLSHDTQQWGDVLPDNGNGTYQTWVA  
TRICQGEEQRFCTYMEHSGNHSTHPVPSGKVLVLQSHW

## **[ ACTIVITY ]**

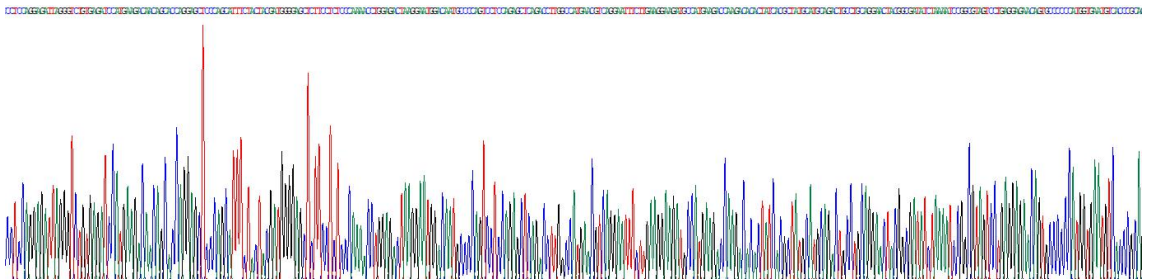
MHC Class I Polypeptide Related Sequence A (MICA) is a stress-induced cell surface protein. It belongs to the MHC class I-like molecules but has distinct features. MICA is highly expressed on cells under stress, such as those with DNA damage or infected cells. Structurally, it has a domain structure that allows for interactions with other molecules. MICA can function as a ligand and binds to Killer Cell Lectin Like Receptor Subfamily D, Member 1 (KLRD1, also known as NKG2D) on the surface of natural killer (NK) cells and some T cells. This binding activates immune cells and enables them to participate in the body's immune surveillance and defense responses. Thus a functional ELISA assay was conducted to detect the interaction of recombinant human MICA and recombinant human KLRD1. Briefly, MICA was diluted serially in PBS with 0.01% BSA (pH 7.4). Duplicate samples of 100  $\mu$ l were then transferred to KLRD1-coated microtiter wells and incubated for 1h at 37°C. Wells were washed with PBST and incubated for 1h with anti-MICA 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  $\mu$ L stop solution to the wells and read at 450/630nm immediately.

The binding activity of recombinant human MICA and recombinant human KLRD1 was shown in Figure 1, the EC50 for this effect is 0.582ug/mL.

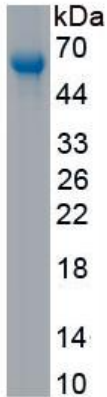


**Figure 1. The binding activity of recombinant human MICA and recombinant human KLRD1**

**[ IDENTIFICATION ]**



**Figure 2. Gene Sequencing (extract)**



**Figure 3. SDS-PAGE**

**Sample: Active recombinant MICA, Human**

**[ 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.