

# ELISA Kit for B-Lymphocyte Activation Antigen B7-2 (LAB7-2)

**Catalog No: tcue271** 

Available Sizes

**Size:** 96T

**Specifications** 

#### **Research Area:**

Signal transduction; CD & Adhesion molecule; Tumor immunity; Infection immunity; Immune molecule;

### **Species Reactivity:**

Homo sapiens (Human)

Sample Type:

Tissue homogenates, cell lysates, cell culture supernates and other biological fluids

#### Sensitivity:

The minimum detectable dose of this kit is typically less than 0.274ng/mL

#### **Detection Range:**

0.156-10ng/mL

## **Assay Time:** 3h

## **Detection Method:**

Enzyme-linked immunosorbent assay for Antigen Detection.

## **Tested Application:**

ELISA

## SwissProt:

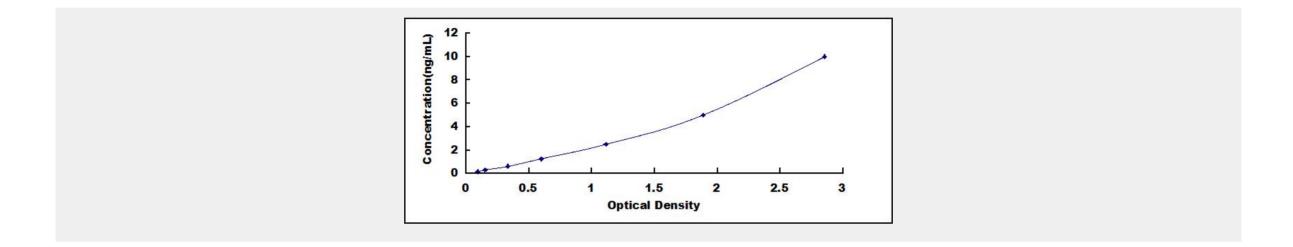
P42081

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## **Test Principle**

The test principle applied in this kit is Sandwich enzyme immunoassay. The microtiter plate provided in this kit has been pre-coated with an antibody specific to B-Lymphocyte Activation Antigen B7-2 (LAB7-2). Standards or samples are then added to the appropriate microtiter plate wells with a biotin-conjugated antibody specific to B-Lymphocyte Activation Antigen B7-2 (LAB7-2). Next, Avidin conjugated to Horseradish Peroxidase (HRP) is added to each microplate well and incubated. After TMB substrate solution is added, only those wells that contain B-Lymphocyte Activation Antigen B7-2 (LAB7-2), biotin-conjugated antibody and enzyme-conjugated Avidin will exhibit a change in color. The enzyme-substrate reaction is terminated by the addition of sulphuric acid solution and the color change is measured spectrophotometrically at a wavelength of 450nm ± 10nm. The concentration of B-Lymphocyte Activation Antigen B7-2 (LAB7-2) in the samples is then determined by comparing the O.D. of the samples to the standard curve.



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