



# **ELISA Kit for Paraoxonase 1 (PON1)**

Catalog No: tcue3067



# **Available Sizes**

Size: 96T



# **Specifications**

#### **Research Area:**

Metabolic pathway; Endocrinology;

#### **Species Reactivity:**

Mus musculus (Mouse)

#### **Sample Type:**

serum, plasma, tissue homogenates, cell lysates, cell culture supernates and other biological fluids

## **Sensitivity:**

The minimum detectable dose of this kit is typically less than 0.51ng/mL

# **Detection Range:**

1.25-80ng/mL

#### **Assay Time:**

3h

# **Detection Method:**

Enzyme-linked immunosorbent assay for Antigen Detection.

# **Tested Application:**

**ELISA** 

#### **SwissProt:**

P52430

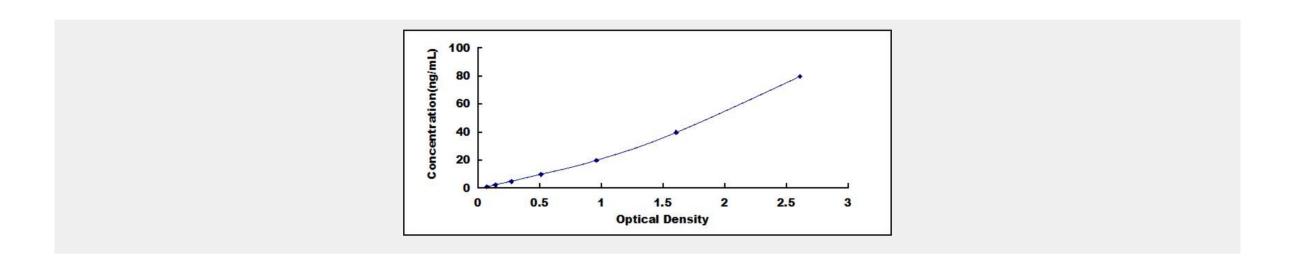
# **Test Principle**

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 Paraoxonase 1 (PON1). Standards or samples are then added to the appropriate microtiter plate wells with a biotin-conjugated antibody specific to Paraoxonase 1 (PON1). 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 Paraoxonase 1 (PON1), 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  $450 \text{nm} \pm 10 \text{nm}$ . The concentration of Paraoxonase 1 (PON1) in the samples is then determined by comparing the O.D. of the samples to the standard curve.



All products are for RESEARCH USE ONLY. Not for diagnostic & therapeutic purposes!