

# ELISA Kit for Paraoxonase 1 (PON1)

# **Catalog No: tcue2399**

**Available Sizes** 

#### **Size:** 96T

Specifications

## **Research Area:** Metabolic pathway;Endocrinology;

**Species Reactivity:** Rattus norvegicus (Rat)

#### 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 0.066ng/mL

#### **Detection Range:**

0.156-10ng/mL

# Assay Time:

3h

## **Detection Method:**

Enzyme-linked immunosorbent assay for Antigen Detection.

Tested Application:	
ELISA	
SwissProt:	

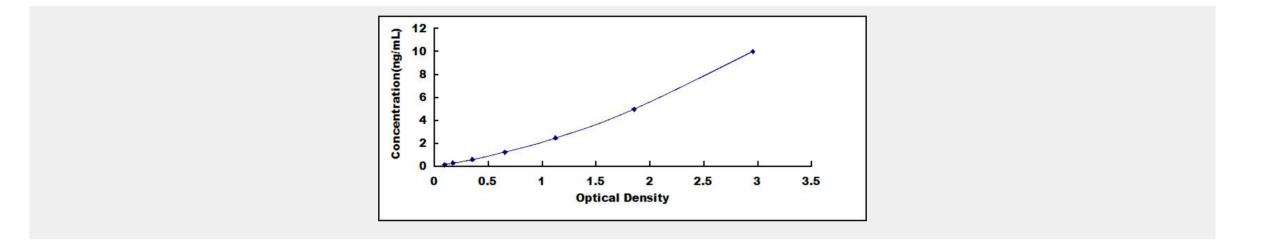
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# **Test Principle**

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 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 450nm  $\pm 10$ 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.



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