



# **ELISA Kit for N-Acetyl Beta-D-Glucosaminidase** (NAGase)

Catalog No: tcue1590



# **Available Sizes**

Size: 96T



# **Specifications**

#### **Research Area:**

Enzyme & Kinase; Kidney biomarker;

## **Species Reactivity:**

Mus musculus (Mouse)

## **Sample Type:**

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

# **Sensitivity:**

The minimum detectable dose of this kit is typically less than 1.28ng/mL

#### **Detection Range:**

3.12-200ng/mL

## **Assay Time:**

3h

#### **Detection Method:**

Enzyme-linked immunosorbent assay for Antigen Detection.

# **Tested Application:**

ELISA

#### **SwissProt:**

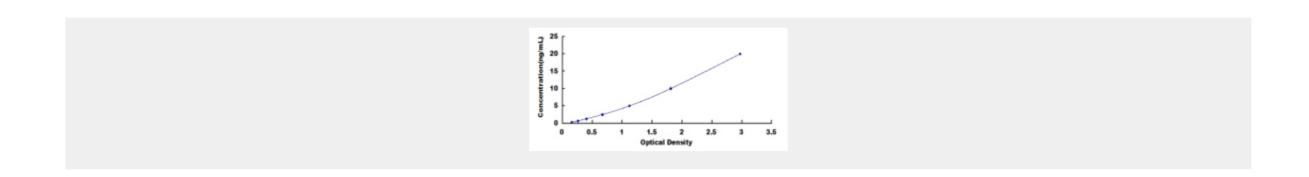
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# **Test Principle**

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 N-Acetyl Beta-D-Glucosaminidase (NAGase). Standards or samples are then added to the appropriate microtiter plate wells with a biotin-conjugated antibody specific to N-Acetyl Beta-D-Glucosaminidase (NAGase). 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 N-Acetyl Beta-D-Glucosaminidase (NAGase), 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  $\pm$  10nm. The concentration of N-Acetyl Beta-D-Glucosaminidase (NAGase) 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!