



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

Catalog No: tcue1589

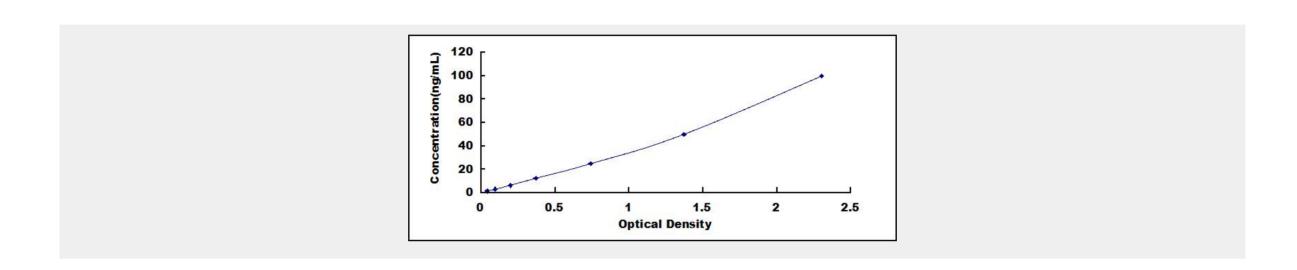
Avail	able Sizes
<b>Size:</b> 96T	
Spec	ifications
Research Ai Enzyme & Kir	rea: nase;Kidney biomarker;
<b>Species Rea</b> Rattus norve	
Sample Typ urine	e:
<b>Sensitivity:</b> The minimum	n detectable dose of this kit is typically less than 0.55ng/mL
<b>Detection R</b> 1.56-100ng/n	
<b>Assay Time</b> : 3h	:
<b>Detection M</b> Enzyme-linke	lethod: ed immunosorbent assay for Antigen Detection.
Tested Appl ELISA	lication:
<b>SwissProt:</b> Q8VIJ5	





## **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!