

Datasheet for ABIN5657143

NAT2 ELISA Kit



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| Quantity: | 96 tests |
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| Target: | NAT2 |
| Reactivity: | Rat |
| Method Type: | Sandwich ELISA |
| Detection Range: | 0.625 ng/mL - 40 ng/mL |
| Minimum Detection Limit: | 0.625 ng/mL |
| Application: | ELISA |

Product Details

| Sample Type: | Tissue Homogenate |
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| Analytical Method: | Quantitative |
| Detection Method: | Colorimetric |
| Specificity: | This assay has high sensitivity and excellent specificity for detection of N-Acetyltransferase 2 (NAT2). No significant cross-reactivity or interference between N-Acetyltransferase 2 (NAT2) and analogues was observed. |
| Sensitivity: | 0.218 ng/mL |

Target Details

| Target: | NAT2 | |
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| Alternative Name: | N-Acetyltransferase 2 (NAT2 Products) | |

Target Details Background: Gene Name: N-Acetyltransferase 2 Gene Aliases: AAC2, PNAT, Arylamine N-Acetyltransferase, Arylamide acetylase 2, Polymorphic arylamine N-acetyltransferase Gene ID: 116632 UniProt: P50298 **Application Details** Comment: The stability of kit is determined by the loss rate of activity. The loss rate of this kit is less than 5 % within the expiration date under appropriate storage condition. To minimize extra influence on the performance, operation procedures and lab conditions, especially room temperature, air humidity, incubator temperature should be strictly controlled. It is also strongly suggested that the whole assay is performed by the same operator from the beginning to the end. Assay Time: 3 h Pre-coated Plate: Protocol: 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-Acetyltransferase 2 (NAT2). Standards or samples are then added to the appropriate microtiter plate wells with a biotin-conjugated antibody specific to N-Acetyltransferase 2 (NAT2). 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-Acetyltransferase 2 (NAT2), biotinconjugated antibody and enzyme-conjugated Avidin will exhibit a change in color. The enzymesubstrate 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 N-Acetyltransferase 2 (NAT2) in the samples is then determined by comparing the O.D. of the samples to the standard curve. Assay Precision: Intra-assay Precision (Precision within an assay): 3 samples with low, middle and high level N-

Intra-assay Precision (Precision within an assay): 3 samples with low, middle and high level N-Acetyltransferase 2 (NAT2) were tested 20 times on one plate, respectively
Inter-assay Precision (Precision between assays): 3 samples with low, middle and high level N-Acetyltransferase 2 (NAT2) were tested on 3 different plates, 8 replicates in each plate. CV(%) = SD/meanX100
Intra-Assay: CV<10%
Inter-Assay: CV<12%

Restrictions:

For Research Use only

Handling

| Handling Advice: | The Stop Solution is acidic. Do not allow to contact skin or eyes. Calibrators, controls and specimen samples should be assayed in duplicate. Once the procedure has been started, all steps should be completed without interruption. |
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| Storage: | 4 °C,-20 °C |
| Storage Comment: | -20°C. Bring all reagents to room temperature before beginning test. The kit may be stored at 4°C for immediate use within two days upon arrival. Reseal any unused strips with desiccant pack. Minimize freeze/thaw cycles. |
| Expiry Date: | 4-8 months |