

# Product datasheet

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# ARG83225 Human ACY1 / Aminoacylase 1 ELISA Kit

Package: 96 wells Store at: 4°C

### **Summary**

Product Description ARG83225 Human ACY1 / Aminoacylase 1 ELISA Kit is an Enzyme Immunoassay kit for the quantification

of Human ACY1 / Aminoacylase 1 in Serum, Plasma and Cell culture supernatants.

Tested Reactivity Hu

Tested Application ELISA

Specificity There is no detectable cross-reactivity with other relevant proteins.

Target Name ACY1 / Aminoacylase 1

Conjugation HRP

Conjugation Note Substrate: TMB and read at 450 nm.

Sensitivity 25 pg/ml

Detection Range 312 pg/ml - 20,000 pg/ml

Sample Type Serum, Plasma and Cell culture supernatants

Precision Intra-Assay CV: 5.0%

Inter-Assay CV: 4.6%

Alternate Names ACY-1; N-acyl-L-amino-acid amidohydrolase; ACY1D; EC 3.5.1.14; HEL-S-5; Aminoacylase-1

### **Application Instructions**

Assay Time ~ 5 hours

### **Properties**

Form 96 well

Storage instruction Store the kit at 2-8°C. Keep microplate wells sealed in a dry bag with desiccants. Do not expose test

reagents to heat, sun or strong light during storage and usage. Please refer to the product user manual

for detail temperatures of the components.

Note For laboratory research only, not for drug, diagnostic or other use.

#### Bioinformation

Gene Symbol ACY1

Gene Full Name aminoacylase 1

Background This gene encodes a cytosolic, homodimeric, zinc-binding enzyme that catalyzes the hydrolysis of

acylated L-amino acids to L-amino acids and an acyl group, and has been postulated to function in the catabolism and salvage of acylated amino acids. This gene is located on chromosome 3p21.1, a region reduced to homozygosity in small-cell lung cancer (SCLC), and its expression has been reported to be reduced or undetectable in SCLC cell lines and tumors. The amino acid sequence of human

aminoacylase-1 is highly homologous to the porcine counterpart, and this enzyme is the first member of a new family of zinc-binding enzymes. Mutations in this gene cause aminoacylase-1 deficiency, a

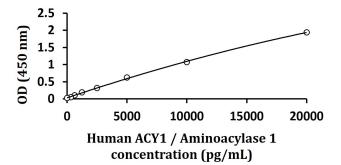
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metabolic disorder characterized by central nervous system defects and increased urinary excretion of N-acetylated amino acids. Alternative splicing of this gene results in multiple transcript variants. Readthrough transcription also exists between this gene and the upstream ABHD14A (abhydrolase domain containing 14A) gene, as represented in GeneID:100526760. A related pseudogene has been identified on chromosome 18. [provided by RefSeq, Nov 2010]

Function Involved in the hydrolysis of N-acylated or N-acetylated amino acids (except L-aspartate). [UniProt]

Cellular Localization Cytoplasm. [UniProt]

### **Images**



ARG83225 Human ACY1 / Aminoacylase 1 ELISA Kit standard curve image

ARG83225 Human ACY1 / Aminoacylase 1 ELISA Kit results of a typical standard run with optical density reading at 450 nm.