

Active Aspartate Aminotransferase (AST) Instruction Manual

SBPB214Hu01

Homo sapiens (Human)

Buffer Formulation	20mM Tris, 150mM NaCl, pH8.0, containing 0.01% SKL, 5% Trehalose.
Traits	Freeze-dried powder
Purity	> 97%
Isoelectric Point	6.6
Applications	Cell culture; Activity Assays.

ACTIVITY TEST

Calculation

$$\text{AST specific activity} = \frac{OD \times d}{t} / \text{amount of protein}$$

Where:

d - dilution factor

t - reaction time in minutes

The specific activity of recombinant human AST is 4.9 U/mg.

Aspartate transaminase(AST) or aspartate aminotransferase, also known as AspAT/ASAT/AAT or glutamic oxaloacetic transaminase, is a pyridoxal phosphate-dependent transaminase enzyme AST catalyzes the reversible transfer of an α -amino group between aspartate and glutamate and, as such, is an important enzyme in amino acid metabolism. AST is found in the liver, heart, skeletal muscle, kidneys, brain, and red blood cells. In this test, an amino group is transferred of from aspartate to α -ketoglutarate. The products of this reversible transamination reaction are oxaloacetate and glutamate. The oxaloacetic acid can be decomposed into pyruvate and carbon dioxide with the present of phenylamine citrate. The activity of aspartate transaminase can be measured by calculating the concentration of the pyruvate. The reaction was performed in adding 10 μ l different concentration recombinant AST(the blank tube add 10 μ l phosphate buffer) to 50 μ l mixture substrate containing 2mM 2-Ketoglutaric acid, 0.1M L-aspartic acid, in 0.2M phosphate buffer, pH7.4, incubate at 37 °C for 1h, then add 10 μ l phenylamine citrate and 50 μ l 2,4-dinitrophenylhydrazine continue incubate at 37 °C for 20min, stop the action by adding 500 μ l 0.4M NaOH, read the OD value at 520nm. Standard curve prepare by double dilute 2 μ M pyruvate with phosphate bufferr then add 10 μ l phenylamine citrate and 50 μ l 2,4-dinitrophenylhydrazine, incubate at 37 °C for

20min and record the OD value at 520nm. One unit of AST is the amount of enzyme that will generate 1 μ mole of pyruvate per minute at pH7.4 at 37 °C.

USAGE

Reconstitute in 20mM Tris, 150mM NaCl (pH8.0) to a concentration of 0.1-1.0 mg/mL. Do not vortex.

STORAGE

Avoid repeated freeze/thaw cycles. Store at 2-8°C for one month. Aliquot and store at -80°C for 12 months.

STABILITY

The thermal stability is described by the loss rate. The loss rate was determined by accelerated thermal degradation test, that is, incubate the protein at 37°C for 48h, and no obvious degradation and precipitation were observed. The loss rate is less than 5% within the expiration date under appropriate storage condition.

Image

CGGGATCCGAATTCATGGCACTCCGTCAGCTTTGCGAGGTTCCGAGGCCCGACCTGTCCTGGCTTTCAAGCTCAGCTGCGGACTTCAGGGAGGATCCGGACCCCGCAGGGTCAACCTGGGAGTGGGACATAATGCAAGGATGACTGCGCTCCCTGGGTTTTCGACGTGCTGAGAAATCGAGCG

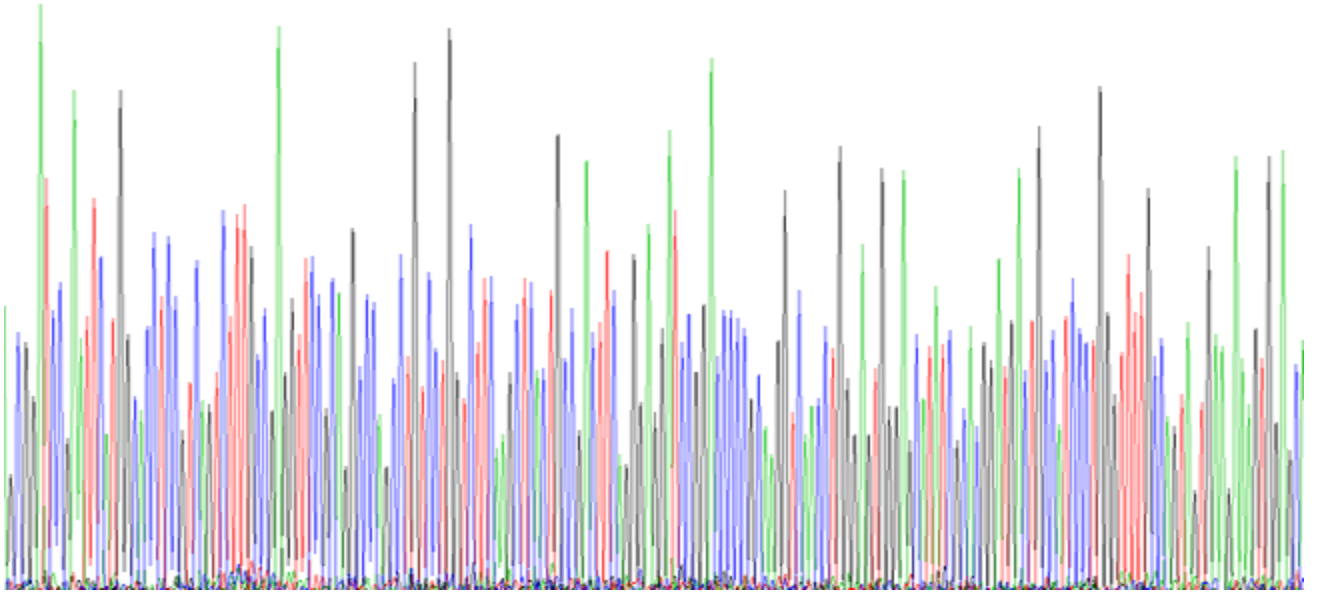


Figure. SDS-PAGE

[IMPORTANT NOTE]

The kit is designed for research use only, we will not be responsible for any issue if the kit was used in clinical diagnostic or any other procedures.