Urease (UE) Activity Assay Kit

Note: Take two or three different samples for prediction before test. Operation Equipment: Spectrophotometer/ microplate reader Catalog Number: NA0346 Size:100T/48S

Components:

Extract solution: 60ml×1 bottle, storage at 4°C.

Reagent I: powder×1 bottle, storage at 4°C. dissolve with 3ml of distilled water before use.

Reagent II: 10ml×1 bottle, storage at 4°C and protected from light.

Reagent III A: 0.4 ml×1 bottle, storage at 4°C.

Reagent III B: 1.6 ml×1 bottle, storage at 4°C. Add Reagent 3A to Reagent 3B, mix for use (name reagent III),

Reagent IV: 2 ml×1 bottle, storage at 4°C.

Standard: 1ml×1 bottle, storage at 4°C. 1mg/ml nitrogen standard solution.

Product Description:

Urease (UE) is widely distributed in the seeds of plants, also in the blood and urine of animals. Some microorganisms can also secrete urease. UE can hydrolyze urea to ammonia and carbonic acid, which plays a key role in urea transformation. The UE activity can be determined by calculating the content of NH3-N with indophenol blue colorimetry.

Reagents and Equipment Required but Not Provided:

Spectrophotometer/ microplate reader, micro glass cuvette/ 96 well flat-bottom plate, constant temperature water bath, low temperature centrifuge, balance, mortar/homogenizer.

Sample preparation:

- I. Bacteria or cells: Number of cells / bacteria (10⁴): volume of extract solution (mL) is 500-1000:1. Suggested 5 million with 1mL of extract solution. Splitting bacteria or cells with ultrasonication (ice bath, power 300W, work time 3s, interval 7s, repeat 3 min), centrifuge at 12000g and 4°C for 15min, supernatant on ice is used for test.
- II. Tissue: Mass (g): extraction volume (mL) is 1:5-10. Add 1 ml of extract solution into 0.1g of tissue, fully grinding on ice. centrifuge at 12000g and 4°C for 15min, supernatant on ice is used for test
- III. Serum/ plasma: Detect directly.

Procedure:

- 1. Preheat Spectrophotometer/ microplate reader for 30min, adjust the wavelength to 630 nm, set the counter to zero with distilled water.
- 2. Dilute 1mg/mL nitrogen standard solution to 2ug/mL with distilled water for use.

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|---|-----------------|--------------------|----------------|--------------------|
| Reagent name(ul) | Blank tube (A2) | Standard tube (A1) | Test tube (A3) | Contrast tube (A4) |
| Sample | - | - | 20 | 20 |
| Distilled water | - | - | | 40 |
| Reagent I | - | - | 40 | - |
| Reagent II | - | - | 80 | 80 |
| Mix thoroughly form mixture, react at 37° C for 1 hour, add the following in EP tube or 96 well plate. | | | | |
| Mixture | - | - | 80 | 80 |
| Distilled water | 80 | - | | |
| Standard | - | 80 | | |
| Reagent III | 16 | 16 | 16 | 16 |
| Reagent IV | 12 | 12 | 12 | 12 |
| Mix thoroughly, stand at RT for 20min. | | | | |
| Distilled water | 92 | 92 | 92 | 92 |
| Mix thoroughly, detect absorbance at 630nm, $\Delta A(\text{standard}) = \Delta A(S) = A1 - A2$, $\Delta A(\text{test}) = \Delta A(T) = A3 - A4$. | | | | |

3. Add the following reagents:

Calculation:

1. Liquid:

Unit definition: One unit of enzyme activity is defined as the amount of enzyme that catalyzes the production of 1ug of NH₃-N in per min every ml liquid.

 $UE(U/mL) = \Delta A(T) \div \Delta A(S) \times C \times V \div Vs \div T = 0.233 \times \Delta A(T) \div \Delta A(S)$

2. Tissue, bacteria or cell:

a. Protein concentration:

Unit definition: One unit of enzyme activity is defined as the amount of enzyme that catalyzes the production of 1ug of NH₃-N in per min every mg tissue protein.

UE (U/mg prot)= $\Delta A(T) \div \Delta A(S) \times C \times V \div (Cpr \times Vs) \div T = 0.233 \times \Delta A(T) \div \Delta A(S) \div Cpr$

b. Sample weight:

Unit definition: One unit of enzyme activity is defined as the amount of enzyme that catalyzes the production of 1ug of NH₃-N in per min every gram tissue.

 $UE(U/g) = \Delta A(T) \div \Delta A(S) \times C \times V \div (W \times Vs \div Ve) \div T = 0.233 \times \Delta A(T) \div \Delta A(S) \div W$

c. Density of bacteria or cell:

Unit definition: One unit of enzyme activity is defined as the amount of enzyme that catalyzes the production of 1ug of NH₃-N in per min every 1 million bacteria or cells.

UE (U/10⁶ cell)= $\Delta A(T) \div \Delta A(S) \times C \times V \div (N \div Vs \times Ve) \div T = 0.233 \times \Delta A(T) \div \Delta A(S) \div N$

C: standard concentration, 2ug/ mL Cpr: Sample concentration (mg/mL);

W: Sample weight(g);

Vs: Sample volume (mL), 0.02 mL;

V: Enzyme reaction volume, 0.14 mL;

Ve: Extraction volume, 1 mL;

T: Reaction time (min), 60 min.

N: cell or bacteria amount, 1 million.

Note:

Dilute the mixture or sample with distilled water before detecting if the ΔA >0.6.

Experimental Examples:

1. Take 0.1g of Vigna radiata and add 1mL extract for sample processing, take the supernatant and operate according to the measurement procedure, measure by the 96 well plate and calculate $\Delta A=A3-A4=0.152-0.092=0.047$, $\Delta As=A1-A2=0.241-0.041=0.200$, calculate the enzyme based on the sample weight:

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UE Activity (U/g weight) =0.233 \times \Delta A \div \Delta A \Rightarrow W = 0.233 \times 0.047 \div 0.2 \div 0.1 = 0.548 U/g weight.
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2. Take 0.1g of kidney and add 1mL extract for sample processing, take the supernatant and operate according to the measurement procedure, measure by the 96 well plate and calculate $\Delta A=A3-A4=0.215-0.162=0.053$, $\Delta s=A1-A2=0.241-0.041=0.200$, calculate the enzyme based on the sample weight:

UE Activity (U/g weight) = $0.233 \times \Delta A \div \Delta As \div W = 0.233 \times 0.053 \div 0.2 \div 0.1 = 0.6175$ U/g weight.

3 Take 20µL of turkey serum and directly follow the measurement procedure, and calculate $\Delta A = 0.145-0.097=0.048$, $\Delta As = A1-A2=0.241-0.041=0.200$, calculate the enzyme based on the sample volume UE Activity (U/mL= $0.233 \times \Delta A \div \Delta As = 0.233 \times 0.048 \div 0.2=0.0559$ U/mL_o

Related Products:

NA0865/NA0622 Nitrate Reductase (NR) Activity Assay Kit NA0754/NA0512 Glutaminase (GLS) Activity Assay Kit NA0753/NA0511 Glutamic Acid Dehydrogenase (GDH) Activity Assay Kit