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Anthocyanin Reductase Activity Assay Kit

Note: Take two or three different samples for prediction before test.

Operation Equipment: Spectrophotometer /Microplate Reader

Cat No: AK0351

Size:100T/48S

Components:

Extract solution: 50mL×1. Storage at 4℃, shake thoroughly before use.

Reagent I: 25mL×1. Storage at 4℃ .

Reagent II: Powder×2. Storage at 4℃, dissolve thoroughly with 1 ml of distilled water before use.

Reagent III: Powder×1. Storage at -20℃, dissolve thoroughly with 1 ml of distilled water and 1 ml of alcohol before use. It can be stored at -20℃ after dispensing to avoid repeated freezing and thawing.

Reagent IV: 1mL×1. Storage at 4℃ .

Product Description:

Anthocyanin reductase (ANR) is a key enzyme in the biosynthesis pathway of procyanidins, which converts anthocyanins into the cis-flavan-3-alcohol. It plays an important role in plants regulation. ANR converts cyanidin chloride to flavane-3-alcohol under the action of NADPH. The activity of ANR can be reflected by measuring the reduction rate of NADPH at 340nm.

Reagents and Equipment Required but Not Provided:

Spectrophotometer/ microplate reader, micro quartz cuvette/96 well flat-bottom UV plate, water bath, low temperature centrifuge, adjustable transferpettor, mortar/homogenizer, ice, alcohol and distilled water.

Sample preparation:

1. Tissue: Add 1 ml of extract solution into 0. 1g of tissue, fully grinding on ice. Centrifuge at 12000rpm and 4C for 15min, supernatant on ice is used for test.
2. Cells or microbial sample: collect cells or microbial sample to centrifuge and remove supernatant. Suggested 5 million with 1mL of extract solution, split bacteria and cells with ultrasonication (power 20%, work time 3s, interval 10s, for 30 times), 12000rpm 4C centrifuge at 12000rpm and 4C for 15min, supernatant on ice is used for test.

Procedure:

1. Preheat spectrophotometer/ microplate reader for 30min, adjust the wavelength to 340 nm, set the counter to zero with distilled water.
2. Add the following reagents:

Reagent name	Test tube (T)	Contrast tube (C)
Reagent I(μL)	170	170
Reagent II (μL)	10	10
Reagent III (μL)	5	5
Sample (μL)	10	-
Mix thoroughly at 37℃ for 30 min		

Reagent IV(μL)	5	5
Sample(μL)	-	10

Mix thoroughly, detect absorbance of test tube and contrast tube at 340nm, named A(T), A(C), ΔA=A(C)- A(T)=A2-A1.

Calculation:

ultra-micro quartz cuvette

1. Protein concentration:

Unit definition: One unit of enzyme activity is defined as the amount of enzyme catalyzes the decreasing of 1 nmol of NADPH per minutes every milligram tissue protein in the reaction system.

ANR (U/mg prot) = ΔA÷(ε×d)×10^9 ×Vrv÷(Vs×Cpr)÷T =107. 18×ΔA÷Cpr

2. Sample weight:

Unit definition: One unit of enzyme activity is defined as the amount of enzyme catalyzes the decreasing of 1 nmol of NADPH per min every gram tissue in the reaction system.

ANR (U/g) =ΔA÷(ε×d)×10^9 ×Vrv÷(W÷Vsv×Vs)÷T=107. 18×ΔA÷W

3. Cells or bacteria:

Unit definition: One unit of enzyme activity is defined as the amount of enzyme catalyzes the decreasing of 1 nmol of NADPH every 10^4 cells or bacteria in the r eaction system per min.

ANR (U/10^4cell) =ΔA÷(ε×d)×10^9 ×Vrv÷(500÷Vsv×Vs)÷T=0.2144×ΔA

Vrv: total reaction volume,0.0002 mL;

ε : NADPH molar extinction coefficient, 6.22×10^3L/mol/cm;

d: light path of cuvette, 1cm;

Vs: supernatant volume (mL), 0.01 mL;

Cpr: sample protein concentration (mg/mL);

T: Reaction time (min), 30 min;

W: Sample weight(g);

Vsv: Extraction volume, 1 mL;

500: 5 million cells.

10^9 : unit conversion coefficient, 1 mol = 10^9 nmol.

96 well UV plate

Change d=1cm to d=0.6cm in the formula.

Note:

1. Dilute react mixture with reagent 1 or decrease sample volume if ΔA>0.4 or A(C)>1(ΔA>0.2 or A(C)>1 with 96 well UV plate). Increase react time (45min or 60min) and sample volume if ΔA is too low.
2. After adding reagent 4, the determination should be completed within 15 minutes.
3. Detect sample concentrate separately.

Experimental Examples:

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1. Take 0. 1g of apple and add 1mL extract to homogenize and grind, take the supernatant and operate according to the measurement procedure, measure by the micro quartz cuvette and calculate ΔA=Ac- At=0.9333-0.8095=0. 1238 , calculate the enzyme based on the sample weight:

ANR Activity（U/g weight）= 107. 18×ΔA÷W=107. 18×0. 1238÷0. 1=132.69 U/g U/g weight。

Related Products:

AK0446/AK0445 Uric Acid(UA) Content Assay Kit
AK0450/AK0449 Plant Total Phenol Content Assay Kit
AK0452/AK0451 Plant Flavonoids Content Assay Kit