

Anthocyanin Content of Plant Assay Kit

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

Operation Equipment: Spectrophotometer

Catalog Number: NA0761

Size: 50T/24S

Components:

Extract solution: Liquid 30 mL ×1. Storage at 4°C.

Reagent I: Liquid 30 mL×1. Storage at 4°C.

Reagent II: Liquid 30 mL×1. Storage at 4°C.

Product Description

Anthocyanin is a kind of edible natural pigment which is easily soluble in water and other solvents. Anthocyanins give plants a colorful color. It also has a variety of health functions. Therefore, it has a broad application prospect in natural edible pigments, health products and pharmaceutical industry.

According to the structure and properties of anthocyanins at different pH, the content of anthocyanins can be determined. The maximum absorption peak of anthocyanin was found at 530 nm when the pH is 1. When the pH is 4.5, anthocyanins are converted to colorless chalcone and there is no absorption peak at 530 nm. The content of anthocyanin can be calculated by measuring the absorbance values at 530 nm and 700 nm at different pH.

Reagents and Equipment Required but Not Provided.

Spectrophotometer, table centrifuge, water-bath, 1 mL glass cuvette, adjustable pipette, mortar/homogenizer and distilled water.

Procedure

I. Sample processing :

According to the tissue weight (g) : the volume of Extract solution (mL) is 1:5-10. (It is recommended that add 1 mL of Extract solution to 0.1 g tissue). Homogenate in ice bath, then transfer to EP tube. Dilute to 1 mL with the Extract solution. Cover tightly and extract at 60°C for 30 min. Several shocks during the period. Centrifuge at 12000 rpm for 10 minutes at room temperature. Take the supernatant for test.

II. Determination Procedure

1. Preheat the spectrophotometer for more than 30 minutes and set the counter to zero with distilled water.
2. Operation table

Reagent Name (μL)	Test tube 1	Test tube 2
Sample	100	100
Reagent I	900	-
Reagent II	-	900

Mix thoroughly. Measure the absorbance of Test tube1 and Test tube2 at 530 nm and 700 nm, respectively. The absorbance values of tube 1 at 530nm and 700nm are recorded as A1 and A1'. The absorbance values of tube 2 at 530nm and 700nm are recorded as A2 and A2'. $\Delta A=(A1-A1')-(A2-A2')$.

III. Calculation of Anthocyanins:

1) Calculate by sample weight

$$\text{Anthocyanins content } (\mu\text{mol/g fresh weight})=[\Delta A \div (\epsilon \times d) \times 10^3 \times F] \times V_E \div W = 0.037 \times \Delta A \times F \div W$$

2) Calculate by Sample protein concentration

$$\text{Anthocyanins content } (\mu\text{mol/mg prot})=[\Delta A \div (\epsilon \times d) \times 10^3 \times F] \times V_E \div (\text{Cpr} \times V_E) = 0.037 \times \Delta A \times F \div \text{Cpr}$$

F: Dilution Factor, 10;

d: Optical diameter of cuvette, 1 cm;

W: Sample weight, g;

ϵ : Molar extinction coefficient of chromoside, 2.69×10^4 mL/mmol/cm;

V_E : Extract volume, 1 mL;

10^3 : 1mmol= 10^3 μ mol;

Cpr: Sample protein concentration, mg/mL; (The protein concentration needs to be extracted separately by PBS and then determined).

Note:

1. If A1 is greater than 1, the dilution ratio can be increased appropriately. Ensure that the total volume is 1 mL. Such as, add 50 μ L of supernatant to 950 μ L of Reagent I (equivalent to 20 times of dilution). If A1 is less than 0.1, the dilution ratio can be reduced appropriately. Ensure that the total volume is 1 mL. Such as, add 500 μ L of supernatant to 500 μ L of Reagent I (equivalent to 2 times of dilution). Keep A1 in the range of 0.1-1. It can improve the detection sensitivity. Note that the volume ratio of supernatant and Reagent II should also be adjusted; when calculating, the actual dilution multiple should be substituted into the following formula.

2. Because the Extract solution will denature the protein, if use the protein concentration to calculate, it needs to be extracted separately with PBS and then measured.

Examples:

1. Add 0.1g prunus peel to 1mL extract solution and mix thoroughly, transfer to EP tube, seal with parafilm to avoid volatilization, immerse and extract at 60°C for 30 min., several shocks during the period. Dilute to 1 mL with the Extract solution, centrifuge at 12000 rpm for 10 minutes at room temperature, take the supernatant, follow the determination procedure to operate, with 96-well flat-bottom plates to calculate: $\Delta A=(A1-A1')-(A2-A2')=(0.249-0.101)-(0.133-0.065)=0.08$, according with mass of sample to calculate: Anthocyanins content ($\mu\text{mol/mg mass}$)= $0.037 \times \Delta A \times F \div W = 0.037 \times 0.08 \times 10 \div 0.1 = 0.296 \mu\text{mol/g mass}$.

Related Products:

NA0769/NA0528 Ceruloplasmin (CP) Assay Kit

NA0768/NA0527 Total antioxidant capacity (T-AOC) Assay Kit

NA0763/NA0522 Uric acid(UA) Assay Kit