Fructose-1, 6-diphosphate (FDP) Activity Assay Kit

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

Operation Equipment: Spectrophotometer

Cat No: BC2240 Size:50T/24S

Components:

Extract solution I: Liquid 30 mL×1. Store at 4°C. Extract solution II: Liquid 5 mL×1. Store at 4°C.

Reagent I: 20 mL×1. Store at 4°C.

Reagent II: 303 μ L×1. Store at 4°C. Dissolve with 0.697 mL of distilled water before use. Unused reagent can store at 4°C for one week.

Reagent III: 15 mL×1. Store at 4°C.

Reagent IV: 40 mL×1. Store at 4°C.

Standard: Powder×1. Store at 4°C. Dissolve with 1.176 mL of distilled water before use to form 50

μmol/mL FDP standard solution

Product Description:

Fructose-1,6-diphosphate (FDP) is an important intermediate product in the glycolysis process. It can regulate a variety of enzymes, improve cell energy metabolism, increase energy utilization, anti-arrhythmia and anti-tissue peroxidation. FDP is widely used in clinical medicine.

Aldolase catalyzes the cleavage of fructose 1,6-diphosphate. The product reacts with 2,4-dinitrophenylhydrazine in acid medium to form 2,4-dinitrophenylhydrazone, which is dark red in alkaline solution and has a characteristic absorption peak at 540 nm.

Reagents and Equipment Required but Not Provided:

spectrophotometer, desk centrifuge, adjustable transferpettor, water bath /incubator, 1 mL glass cuvette, mortar / homogenizer, Ultrasonic crusher, ice and distilled water.

Procedure:

I. Sample preparation:

1) Tissue

According to the tissue weight (g): the volume of the extract (mL) is $1:5 \sim 10$. Suggest adding 1 mL of Extract solution I to 0.1 g of tissue, fully homogenize on ice bath. Centrifuge at $12000 \times g$ for 10 minutes at 4°C. Take 0.8mL supernatant and 0.16mL Extract solution II to mix well, centrifuge at $12000 \times g$ for 10 minutes at 4°C. Then take supernatant for test.

2) Bacteria or cells

According to the Bacteria or cells (10⁴): the volume of the extract (mL) is 500~1000:1. Suggest add 1mL

of Extract solution to 5 million of bacteria or cells. Use ultrasonication to splitting bacteria and cells (placed on ice, ultrasonic power 300W, working time 3 seconds, interval 7 seconds, total time for 3 min). Centrifuge at 12000 ×g for 10 minutes at 4°C. Take 0.8mL supernatant and 0.16mL Extract solution II to mix well, centrifuge at 12000 ×g for 10 minutes at 4°C. Then take supernatant for test.

3) Liquid:

Add 1mL Extract solution I to 100μ L liquid sample, centrifuge at $12000 \times g$ for 10 minutes at 4°C. Take 0.8mL supernatant and 0.16mL Extract solution II to mix well, centrifuge at $12000 \times g$ for 10 minutes at 4°C. Then take supernatant for test.

Determination procedure:

- 1. Preheat spectrophotometer for 30 minutes, adjust the wavelength to 540 nm, set zero with distilled water.
- 2. $50 \mu mol/ml$ fructose-1,6-diphosphate standard solution is diluted to 1.5625, 0.78125, 0.39, 0.2 and 0.1 $\mu mol/ml$ standard solution with distilled water.

3. Sampling table:

Reagent name (µL)	Control tube (A _C)	Test tube (A _T)	Blank tube (A _B)	Standard tube (As)
Sample	100	100	-	-
Distilled water	-	-	100	-
Standard solution			-	100
Reagent I	220	200	220	200
Reagent II	-	20	-	20
Mix well, react accurately at 37°C for 2 h				
Reagent III	200	200	200	200
Mix well, react accurately at 37°C for 20 min				
Reagent IV	500	500	500	500
Mix well, react accurately at 37°C for 10 min				

The absorbance value at 540 nm is measured in 1 ml glass cuvette and recorded as A_C , A_T , A_B , A_S , respectively. Calculate $\Delta A = A_T - A_C$, $\Delta A_S = A_S - A_B$. The blank tube only needs to be tested 1-2 times.

II. Calculation:

1. According to concentration of standard solution and ΔA_S to create the standard curve, take standard solution as X-axis, ΔA_S as Y-axis. Take ΔA into the equation to obtain x (μ mol/ml).

2. Calculation:

(1) sample weight

FDP (mg/g fresh weight) =
$$x \times (V_{su} + V_{exII}) \times M \div (W \times V_{su} \div V_{exI}) = 408x \div W$$

(2) The number of bacteria or cells

FDP (mg/10⁴ cell) =
$$x \times (V_{su} + V_{exII}) \times M \div (cell \text{ amount } \times V_{su} \div V_{exI}) = 408x \div cell \text{ amount}$$

(3) Liquid:

FDP (
$$\mu$$
mol/mL) = $x \times (V_{su} + V_{exII}) \div (V_L \times V_{su} \div (V_{exI} + V_L)) = 13.2x$

 V_{su} : Supernatant volume of extraction, 0.8mL

 V_{exII} : Extract solution II volume, 0.16mL

M: Molecular weight of fructose-1,6-diphosphate,340

V_{exI}: Extract solution I volume, 1mL

W: sample weight, g

Cell amount: 10 thousand cells as unit V_L : liquid sample volume, 0.1mL.

Note:

1. If ΔA >0.5, please dilute the sample with water to appropriate concentration, multiply dilute times in the formula.

Related Products:

BC2270/BC2275 Fructose-bisphosphate aldolase(FBA) Activity Assay Kit BC2250/BC2255 Phosphoglycerate Kinase(PGK) Activity Assay Kit