

Soil Alkaline Xylanase (S-BAX) Activity Assay Kit

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

Detection equipment: Spectrophotometer

Cat No: BC5740 **Size:** 50T/24S

Components:

Buffer Fluid: Liquid 25 mL×1, store at 2-8°C. **Reagent I:** Liquid 12 mL×1, store at 2-8°C. **Reagent II:** Liquid 16 mL×1, store at 2-8°C.

Standard: Powder×1, 10mg xylose. Before use, a standard solution of 100μ mol/mL was prepared by adding 667μ L distilled water and stored at 2-8°C for 8 weeks.

Product Description:

Soil alkaline xylanase (S-BAX), also known as soil alkaline hemicellulase, is mainly isolated from microorganisms with an optimal growth pH of 9-11.

In an alkaline environment, S-BAX catalyzes the degradation of xylan into reducing oligosaccharides and monosaccharides. Under boiling water bath conditions, it further undergoes a color reaction with 3,5-dinitrosalicylic acid, with a characteristic absorption peak at 540nm. The color depth of the reaction solution is directly proportional to the amount of reducing sugars produced by enzymatic hydrolysis. By measuring the rate of increase in absorbance of the reaction solution at 540nm, S-BAX activity can be calculated.

Required but not provided:

Spectrophotometer, 1mL glass cuvette, balance, desk centrifuge, water bath, 30-50 mesh sieve, distilled water.

Procedure:

I. Sample preparation(The sample size to be tested can be adjusted appropriately, and the specific proportion can be referred to in the literature)

Fresh soil samples are naturally air dried or air dried in a 37 °C oven, and sieved through a 30-50 mesh sieve.

II. Determination procedure

- 1. Preheat spectrophotometer for 30 minutes, adjust wavelength to 540 nm, set the counter to zero with distilled water.
- 2. Dilution of standard solution: Using distilled water to dilute the Standard into 1.5, 1.2, 1, 0.8, 0.6, 0.4µmol/mL of standard solution before measured.
- 3. Quasi-dilution table:

Num-	Predilution concentration	Standard volume	Volume of distilled	Diluted concentration
ber	(µmol/mL)	(µL)	water (µL)	(µmol/mL)



1	100	100	900	10
2	10	150	850	1.5
3	10	120	880	1.2
4	10	150	1350	50) 50°
5	1	400	100	0.8
6	1	300	200	0.6
7	1	200	300	0.4

Note: 300µL per tube is required in the experiment.

4. Sample determination (add the following reagents in EP tube in turn).

Reagent Name (µL)	Control tube (A _C)	Test tube (A _T)	Blank tube (A _B)	Standard tube (A _S)
Sample	0.1g	0.1g	-	OS CHENCES
Buffer Fluid	400	400	- /	SUFE"
Reagent I	105	200	-	(g) <u>-</u>
Mix well and place	ce in a 50°C water bath	for 2 hours.		
Immediately after, heat	in a boiling water bath	**************************************		
deactivate (be careful	not to let the lid burst of	1 Style Fig.	-	
water from entering an	nd altering the reaction	system). Cool to	0,500	©
(2,00,1	room temperature.		1010	
Reagent I	200	-	-	COLSTERN
At room temperature,	centrifuge at 12000g for	4	3 716.	
ta	ake the supernatant	-	-	
Supernatant	300	300	® -	-
Distilled water		-	300	-
Standard	<u>-</u>	-	18 JENO -	300
Reagent II	200	200	200	200
Mix well, accurately of	color in a boiling water	bath for 5 minutes	(be careful not to let	the lid burst open to
avoid wat	er entering and altering	the reaction syster	n), cool to room temp	perature.
Distilled water	500	500	500	500

 Distilled water
 500
 500
 500

After thorough mixing, take the supernatant and measure the absorbance at 540nm in a 1mL glass cuvette. Record the absorbance values as A_C , A_T , A_B , A_S , calculate $\Delta A_T = A_T - A_C$, $\Delta A_S = A_S - A_B$. The blank tube and standard curve only need to be measured 1-2 times. Each test tube should have a corresponding control tube.

III. Calculation

1. Drawing of standard curve:

The standard curve is established according to the concentration of the standard tube $(x, \mu mol/mL)$

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and the absorbance ΔA_S (y, ΔA_S). According to the standard curve, the ΔA_T (y, ΔA_T) is brought into the formula to calculate the sample concentration (x, μ mol/mL).

2. Calculation of soil S-BAX activity:

Enzyme activity definition: An enzyme activity unit of an alkaline xylanase is defined as the amount of enzyme required to produce 1 μ mol of reducing sugar per gram of soil per hour by decomposing xylan at 50°C and pH 9.0.

S-BAX activity (U/g soil) =
$$x \times V_{RV} \div W \div T \times F = 0.3 \times x \div W \times F$$

V_{RV}: Total reaction volume, 0.6mL;

W: Sample mass, g;

T: Reaction time, 2h;

F: Sample dilution ratio;

Note:

- 1. If ΔA_T is less than 0.01, the sample size can be appropriately increased or the reaction time can be extended by 50 °C before measurement; If the ΔA_T is greater than 1.5 or A_T is greater than 1.5, the supernatant can be diluted with distilled water before measurement, and attention should be paid to synchronously modifying the dilution factor in the calculation formula.
- 2. It is recommended to use a spiral tube to prevent the lid from bursting during the boiling water bath process and change the reaction system.

Experimental example:

Take 0.1g of wild mushroom soil and follow the measurement steps to measure it using a 1mL glass cuvette, calculate $\Delta A_T = A_T - A_C = 0.406 - 0.108 = 0.298$, bring in the standard curve y=0.5887x-0.1537 (R²=0.9982), calculate x=0.767, and calculate S-BAX activity based on sample mass:

S-BAX activity (U/g soil)= $0.3 \times x \div W \times F=2.301$ U/g soil.

Related Products:

BC5720/BC5725 Soil Acid Xylanase (S-ACX) Activity Assay Kit BC5730/BC5735 Soil Neutral Xylanase (S-NEX) Activity Assay Kit