

EZMeta™ Glycogen Colorimetric Assay Kit

Cat #: D-AKC1340

Size: 96T

Storage: Stored at 4°C for 12 months, protected from light.

Product Information

Detection range: 0.003125-0.25 mg/mL

Sensitivity: 0.003125 mg/mL

Applicable samples: Animal Tissues, Bacteria, Cells

Assay Principle

Glycogen is a polymer polysaccharide composed of glucose. It is one of the main storage forms of glucose. It is mainly stored in liver and muscle as reserve energy, which is called liver glycogen and muscle glycogen, respectively. Liver glycogen can regulate the concentration of blood glucose. When blood glucose rises, glycogen can be synthesized in the liver. When blood glucose decreases, liver glucose principle decomposes into glucose to supplement blood glucose. Therefore, liver glycogen is very important to maintain the relative balance of blood glucose. Muscle glycogen is the storage form of glucose in muscle. When strenuous exercise consumes a large amount of blood glucose, muscle glycogen cannot be decomposed directly into blood glucose. It was decomposed to produce lactic acid and circulate to the liver with blood. It is transformed into liver glycogen and glucose by gluconeogenesis. EZMeta™ Glycogen Colorimetric Assay Kit can detect glycogen concentration in animal tissues, bacteria and cells. The principle of kit is strong alkaline extract glycogen. under strong acidic conditions, blue compounds with anthrone chromogenic agent with characteristic absorption peak at 620 nm. In a certain concentration range, glycogen concentration is linearly related to 620 nm absorbance. According to the standard curve, the glycogen concentration in the sample can be calculated.

Materials Supplied and Storage Conditions

Kit components	Size (96T)	Storage conditions
Extraction Buffer	100 mL	4°C
Chromogen	1	4°C, protected from light
Standard	1 mL	4°C

Materials Required but Not Supplied

- Microplate reader or visible spectrophotometer capable of measuring absorbance at 620 nm
- Refrigerated centrifuge, water bath
- 96-well plate or microglass cuvette, precision pipettes, disposable pipette tips, EP tube
- Deionized water, concentrated sulfuric acid

Reagent Preparation

Extraction Buffer: Ready to use as supplied. Equilibrate to room temperature before use. Store at 4°C.

Chromogen: Add 6 mL deionized water to dissolve the powder, then slowly pour into 24 mL concentrated sulfuric acid, fully dissolve and mix. 4°C avoid light for one week.

Standard: 1 mg/mL. Store at 4°C.

Standard curve setting: Dilute 1 mg/mL standard with deionized water to 0.25, 0.1, 0.05, 0.025, 0.0125, 0.00625, 0.003125 mg/mL standard solution as shown in the table below.

NUM.	Volume of Standard	Deionized Water (μL)	Concentration (mg/mL)
Std.1	50 μL 1 mg/mL	150	0.25
Std.2	160 μL of Std.1	240	0.1
Std.3	200 μL of Std.2	200	0.05
Std.4	200 μL of Std.3	200	0.025
Std.5	200 μL of Std.4	200	0.0125

Std.6	200 μ L of Std.5	200	0.00625
Std.7	200 μ L of Std.6	200	0.003125

Sample Preparation

Note: Fresh samples are recommended. If the experiment is not carried out immediately, the samples can be stored at **-80°C for 1 month.**

1. Tissue samples: Weigh 0.1 g tissue, put it in 10 mL test tube, add 0.75 mL Extraction Buffer, boil 20 min (cover tightly to prevent water loss). shake test tube every 5 min, mix well. When the tissue is dissolved, dilute with deionized water to 5 mL, centrifuge at 8,000 g for 10 min at 25°C. Use supernatant for assay.

2. Bacteria or Cells: collect 5 million bacteria or cells into EP tubes, wash with PBS, add 0.75 mL Extraction Buffer, Ultrasonic lysis of bacteria or cells (200 W, work 3 s, intermittent 10 s, work 30 times). Transferred to 10 mL tube, boil 20 min (cover tightly to prevent water loss), shake the tube every 5 min, mix well. Dilute with deionized water to 5 mL, mix well, 8,000 g, 25°C centrifuge 10 min. Use supernatant for assay.

Assay Procedure

1. Preheat the microplate reader or visible spectrophotometer for more than 30 min, and adjust the wavelength to 620 nm. Visible spectrophotometer was returned to zero with deionized water.

2. Add the following reagents respectively into each EP tube:

Reagent	Blank Tube (μ L)	Standard Tube (μ L)	Test Tube (μ L)
Sample	0	0	60
Standard	0	60	0
Deionized Water	60	0	0
Chromogen	240	240	240

3. Mix well, incubate in 95°C 10 min (Cover tightly to prevent moisture evaporation), add 200 μ L of the reaction mix to 96-well plate or microglass cuvette. Measure absorbance at OD620 nm and record as A_{Blank} , $A_{Standard}$, A_{Test} . Calculate

$$\Delta A_{Test} = A_{Test} - A_{Blank}, \Delta A_{Standard} = A_{Standard} - A_{Blank}.$$

Note: Extraction Buffer is corrosive, Chromogen is toxic. Please take protective measures when operating. In order to guarantee the accuracy of experimental results, need to do a pre-experiment with 2-3 samples. If ΔA_{Test} is less than 0.001, increase the sample quantity appropriately. If ΔA_{Test} is greater than 1.5, the sample can be appropriately diluted with deionized water, the calculated result multiplied by the dilution factor, or decrease the sample quantity appropriately.

Data Analysis

Note: We provide you with calculation formulae, including the derivation process and final formula. The two are exactly equal. It is suggested that the concise calculation formula in bold is final formula.

1. Drawing of standard curve:

With the concentration of the standard solution as the y axis and the $\Delta A_{\text{Standard}}$ as the x axis, draw the standard curve.

Substitute the ΔA_{Test} into the equation to obtain the y value (mg/mL).

2. Calculated by sample quality

$$\text{Glycogen (mg/g)} = 1.11 \times (y \times V_{\text{Sample}}) \div (W \times V_{\text{Sample}} \div V_{\text{Total sample}}) \times n = \mathbf{5.55 \times y \div W \times n}$$

3. Calculated by protein concentration

$$\text{Glycogen (mg/mg prot)} = 1.11 \times (y \times V_{\text{Sample}}) \div (V_{\text{Sample}} \times \text{Cpr}) \times n = \mathbf{1.11 \times y \div \text{Cpr} \times n}$$

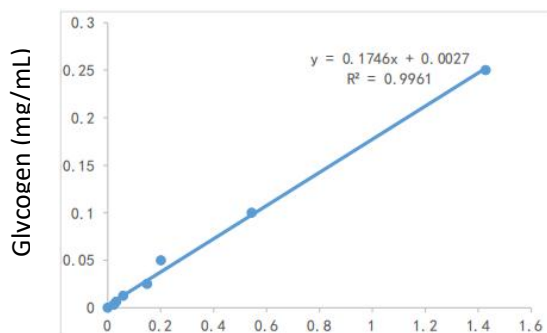
4. Calculated by the number of bacteria or cells

$$\text{Glycogen (mg/10}^4\text{)} = 1.11 \times (y \times V_{\text{Sample}}) \div (\text{the number of bacteria or cells} \times V_{\text{Sample}} \div V_{\text{Total sample}}) \times n = \mathbf{5.55 \times y \div \text{the number of bacteria or cells} \times n}$$

Where: 1.11: 100 μg glucose with anthrone reagent color equivalent to 111 μg glycogen. V_{Sample} : the sample volume to be tested, 0.06 mL. W: sample mass, g. $V_{\text{Total Sample}}$: sample volume, 5 mL. n: the sample dilution factor. Cpr: Sample protein concentration, mg/mL. The number of bacteria or cells: 10^4 as the unit, ten thousand.

Typical Data

Typical standard curve:



OD620

Fig.1. Standard curve of glycogen, data provided for demonstration purposes only. A new standard curve must be generated for each assay.

Disclaimer

The reagent is only used in the field of scientific research, not suitable for clinical diagnosis or other purposes.