



Total Carbohydrate Microplate Assay Kit

Catalog # AS0156

Detection and Quantification of Total Carbohydrate Content in
Serum, Plasma, Tissue extracts, Cell lysate, Food, Juice, Beverage,
Other agricultural products Samples.

This instruction must be read in its entirety before using this product.

For research use only, Not for use in diagnostic procedures.

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I. INTRODUCTION

Carbohydrates are the most abundant biomolecules present in all living organisms. Carbohydrates have many functions, as structural components to the cell walls of bacteria and plants, or as energy storage in the form of starch and glycogen.

Carbohydrates are also a major component of the human diet.

Total Carbohydrate Microplate Assay Kit can be used for measuring carbohydrates in a variety of samples, including food and beverage products. These compounds react with the developer to generate a chromagen, which can be detected at 540 nm.

II. KIT COMPONENTS

Component	Volume	Storage
96-Well Microplate	1 plate	
Assay BufferI	30 mlx 2	4 °C
Assay BufferII	30 mlx 2	4 °C
Dye Reagent	10 mlx 1	4 °C
Standard (0.5 mg/ml)	1 mlx 1	4 °C
Technical Manual	1 Manual	

III. MATERIALS REQUIRED BUT NOT PROVIDED

1. Microplate reader to read absorbance at 540 nm
2. Distilled water
3. Pipettor
4. Pipette tips
5. Mortar
6. Centrifuge
7. Timer
8. Convection oven

IV. SAMPLE PREPARATION

1. For tissue samples

Weigh out 0.05g sample in a centrifuge tube, homogenize with 0.5 ml Assay Buffer I, put it in boiling water bath for 30 minutes, when cold, add 0.5 ml Assay Buffer II; centrifuged at 8,000g at room temperature for 10 minutes, take the supernatant into a new centrifuge tube for detection.

2. For liquid samples

Add 0.05 ml sample into a centrifuge tube, then add 0.5 ml Assay Buffer I, put it in boiling water bath for 30 minutes, when cold, add 0.5 ml Assay Buffer II; centrifuged at 8,000g at room temperature for 10 minutes, take the supernatant into a new centrifuge tube for detection.

V. ASSAY PROCEDURE

Add following reagents into the microplate:

Reagent	Sample	Standard	Blank
Sample	100 μ l	--	--
Standard	--	100 μ l	--
Distilled water	--	--	100 μ l
Dye Reagent	100 μ l	100 μ l	100 μ l

Mix, put the plate into the convection oven, 90°C for 10 minutes. When cold, record absorbance measured at 540 nm.

VI. CALCULATION

1. According to the volume of sample

$$\begin{aligned} \text{Total Carbohydrate (mg/ml)} &= (C_{\text{Standard}} \times V_{\text{Standard}}) \times (\text{OD}_{\text{Sample}} - \text{OD}_{\text{Blank}}) / (\text{OD}_{\text{Standard}} - \\ &\quad \text{OD}_{\text{Blank}}) / (V_{\text{Sample}} \times V / V_{\text{Assay}}) \times 0.9 \\ &= 0.45 \times (\text{OD}_{\text{Sample}} - \text{OD}_{\text{Blank}}) / (\text{OD}_{\text{Standard}} - \text{OD}_{\text{Blank}}) / V \end{aligned}$$

2. According to the weight of sample

$$\begin{aligned} \text{Total Carbohydrate (mg/g)} &= (C_{\text{Standard}} \times V_{\text{Standard}}) \times (\text{OD}_{\text{Sample}} - \text{OD}_{\text{Blank}}) / (\text{OD}_{\text{Standard}} - \\ &\quad \text{OD}_{\text{Blank}}) / (V_{\text{Sample}} \times W / V_{\text{Assay}}) \times 0.9 \\ &= 0.45 \times (\text{OD}_{\text{Sample}} - \text{OD}_{\text{Blank}}) / (\text{OD}_{\text{Standard}} - \text{OD}_{\text{Blank}}) / W \end{aligned}$$

C_{Standard} : the standard concentration, 0.5mg/ml;

C_{Protein} : the protein concentration, mg/ml;

W: the weight of sample, g;

V: the volume of sample, ml;

V_{Standard} : the volume of standard, 0.1 ml;

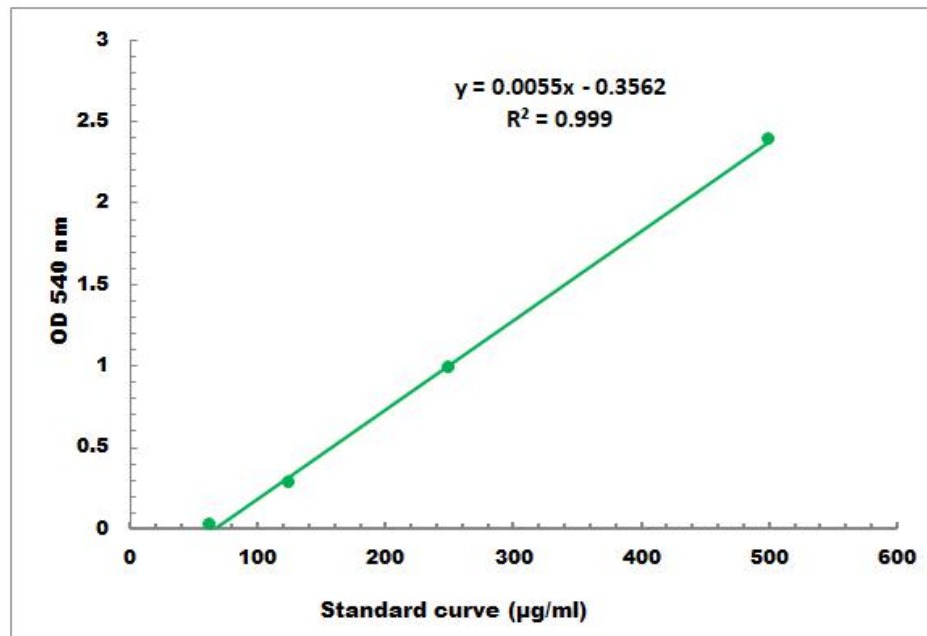
V_{Sample} : the volume of sample, 0.1 ml;

V_{Assay} : the volume of Assay BufferI and Assay BufferII, 1 ml;

0.9: conversion factor.

VII. TYPICAL DATA

The standard curve is for demonstration only. A standard curve must be run with each assay.



Detection Range: 50µg/ml - 500µg/ml

VIII. TECHNICAL SUPPORT

For troubleshooting, information or assistance, please go online to www.sabbiotech.cn or contact us at techcn@signalwayantibody.com

IX. NOTES