



Mannitol Microplate Assay Kit

User Manual

Catalog # ASK1165

Detection and Quantification of Mannitol Content in Serum, Urine,
Other biological fluids, Food, Beverage Samples.

For research use only. Not for diagnostic or therapeutic procedures.

Bioworld Technology, Inc. (USA)

Email: info@bioworld.com

Web: www.bioworld.com

Bioworld technology, co. Ltd. (China)

Email: info@biogot.com

Web: www.biogot.com



I. INTRODUCTION.....2

II. KIT COMPONENTS.....3

III. MATERIALS REQUIRED BUT NOT PROVIDED.....3

IV. SAMPLE PREPARATION.....4

V. ASSAY PROCEDURE.....5

VI. CALCULATION.....6

VII. TYPICAL DATA.....7



I. INTRODUCTION

Mannitol is a sugar alcohol used in dietary supplement, sweetener, intestinal permeability test for leaky gut, etc. It also serves as a coating for hard candies, dried fruits, and chewing gums due to its low ability to attract and hold water molecules. In addition, it is an osmoprotectant for plants and is used clinically in osmotherapy to reduce intracranial pressure.

Mannitol Microplate Assay Kit is designed to measure mannitol in various samples in 96-well microplate. The color intensity at 413 nm is directly proportional to mannitol concentration in the sample.



II. KIT COMPONENTS

Component	Volume	Storage
96-Well Microplate	1 plate	
Reaction Buffer	2.5 ml x 1	4 °C
Stop Solution	Powder x 1	4 °C
Dye Reagent	10 ml x 1	4 °C
Standard	Powder x 1	4 °C
Plate Adhesive Strips	3 Strips	
Technical Manual	1 Manual	

Note:

Standard: add 1 ml Distilled water to dissolve before use, then add 0.1 ml into 0.9 ml Distilled water, the concentration will be 200 µg/ml.

Dye Reagent: add 5 ml distilled water to dissolve before use.

III. MATERIALS REQUIRED BUT NOT PROVIDED

1. Microplate reader to read absorbance at 413 nm
2. Distilled water
3. Pipettor
4. Pipette tips
5. Distilled water



IV. SAMPLE PREPARATION

1. For serum, urine and other biological fluids samples

Samples can be assayed directly.

**V. ASSAY PROCEDURE**

Add following reagents in the microplate:

Reagent	Sample	Standard	Blank
Sample	25 μ l	--	--
Standard	--	25 μ l	--
Distilled water	--	--	25 μ l
Reaction Buffer	25 μ l	25 μ l	25 μ l
Mix, incubate at room temperature for 10 minutes.			
Stop Solution	50 μ l	50 μ l	50 μ l
Mix, incubate at room temperature for 5 minutes.			
Dye Reagent	100 μ l	100 μ l	100 μ l
Mix, cover the plate adhesive strip, put the plate into the oven, incubate at 53 °C for 15 minutes, then put it on ice immediately, record absorbance measured at 413 nm.			



VI. CALCULATION

1. According to the volume of sample

$$\begin{aligned} \text{Mannitol } (\mu\text{g/ml}) &= C_{\text{Standard}} \times V_{\text{Standard}} \times (\text{OD}_{\text{Sample}} - \text{OD}_{\text{Blank}}) / (\text{OD}_{\text{Standard}} - \text{OD}_{\text{Blank}}) / \\ & \quad V_{\text{Sample}} \\ &= 200 \times (\text{OD}_{\text{Sample}} - \text{OD}_{\text{Blank}}) / (\text{OD}_{\text{Standard}} - \text{OD}_{\text{Blank}}) \end{aligned}$$

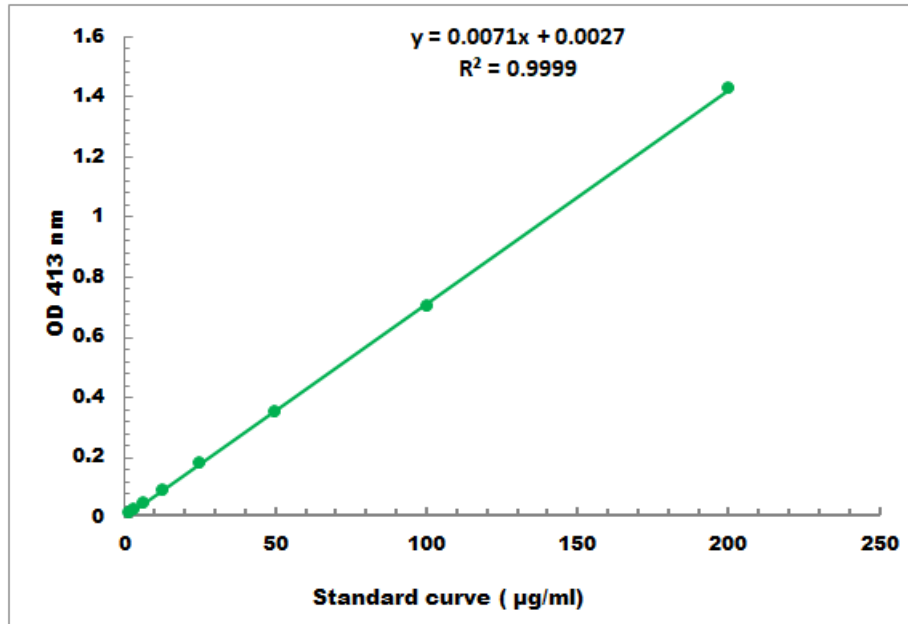
C_{Standard} : the standard concentration, 200 $\mu\text{g/ml}$;

V_{Standard} : the volume of standard, 0.025 ml;

V_{Sample} : the volume of sample, 0.025 ml.

VII. TYPICAL DATA

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



Detection Range: 2 µg/ml - 200 µg/ml