

Albumin (Plant) Microplate Assay Kit User Manual

Catalog # CAK1197

(Version 1.2A)

Detection and Quantification of Albumin (Plant) Content in Tissue extracts, Powder Samples.

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



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I. INTRODUCTION

Albumin (Plant) Microplate Assay Kit is a sensitive assay for determining albumin content in plant samples. The color intensity, measured at 595 nm, is proportionate to albumin content in the sample.



II. KIT COMPONENTS

Component	Volume	Storage
96-Well Microplate	1 plate	
Assay Buffer	30 ml x 2	4 °C
Dye Reagent	20 ml x 1	4 °C
Standard	Powder x 1	-20 °C
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Note:

Standard: add 1 ml distilled water to dissolve before use, the concentration will be 2 mg/ml.

III. MATERIALS REQUIRED BUT NOT PROVIDED

- 1. Microplate reader to read absorbance at 595 nm
- 2. Distilled water
- 3. Pipettor, multi-channel pipettor
- 4. Pipette tips
- 5. Mortar
- 6. Ice
- 7. Centrifuge
- 8. Timer
- 9. Lab rotator



IV. SAMPLE PREPARATION

1. For tissue samples

Weigh out 0.05 g tissue, homogenize with 0.5 ml Assay Buffer on ice, transfer it to centrifuge tube and mix on a lab rotator for 30 minutes; centrifuged at $10000g\ 4\ ^{\circ}C$ for 10 minutes, take the supernatant into a new centrifuge tube and keep it on ice for detection.

2. For powder samples

Weigh out 0.05 g powder, add 0.5 ml Assay Buffer to dissolve, mix on a lab rotator for 30 minutes; centrifuged at 10000g 4 °C for 10 minutes, take the supernatant into a new centrifuge tube and keep it on ice for detection.



V. ASSAY PROCEDURE

Add following reagents into the microplate:

Reagent	Sample	Standard	Blank
Sample	10 μΙ		
Standard		10 μΙ	
Distilled water			10 μΙ
Dye Reagent	200 μΙ	200 μΙ	200 μΙ

Mix, wait for 2 minutes, measured at 595 nm and record the absorbance.

Note:

- 1) Perform 2-fold serial dilutions of the top standards to make the standard curve.
- 2) The concentrations can vary over a wide range depending on the different samples. For unknown samples, we recommend doing a pilot experiment & testing several doses to ensure the readings are within the standard curve range.
- 3) Reagents must be added step by step, can not be mixed and added together.



VI. CALCULATION

1. According to the weight of sample

Albumin (mg/g) =
$$(C_{Standard} \times V_{Standard}) \times (OD_{Sample} - OD_{Blank}) / (OD_{Standard} - OD_{Blank}) / (V_{Sample} \times W / V_{Assay})$$

= $(OD_{Sample} - OD_{Blank}) / (OD_{Standard} - OD_{Blank}) / W$

C_{Standard}: the standard concentration, 2 mg/ml;

 $V_{Standard}$: the volume of standard, 10 μ l = 0.01 ml;

 V_{Sample} : the volume of sample, 10 μ l = 0.01 ml;

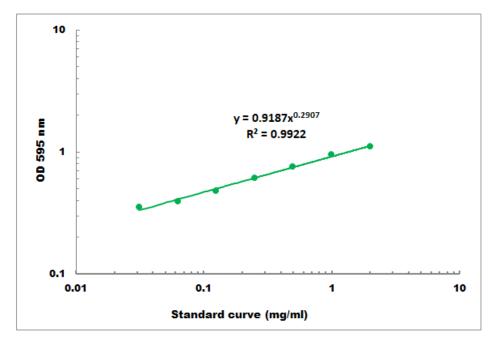
W: the weight of sample, g;

 V_{Assay} : the volume of Assay Buffer, 0.5 ml.



VII. TYPICAL DATA

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



Detection Range: 0.02 mg/ml - 2 mg/ml

VIII. TECHNICAL SUPPORT

For troubleshooting, information or assistance, please go online to www.cohesionbio.com or contact us at techsupport@cohesionbio.com

IX. NOTES