

ChloralMicroplate Assay Kit User Manual

Catalog # CAK1252

(Version 1.3A)

Detection and Quantification of Chloral Contentin Water and Other biological fluids Samples.

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



I. INTRODUCTION	2
II. KIT COMPONENTS	3
III. MATERIALS REQUIRED BUT NOT PROVIDED	
IV. SAMPLE PREPARATION	
V. ASSAY PROCEDURE	
VI. CALCULATION	
VII. TYPICAL DATA	
VIII. TECHNICAL SUPPORT	
IX NOTES.	7



I. INTRODUCTION

Chloral, also known as trichloroacetaldehyde or trichloroethanal, is the organic compound with the formula Cl₃CCHO. This aldehyde is a colourless oily liquid that is soluble in a wide range of solvents. It reacts with water to form chloral hydrate, a once widely used sedative and hypnotic substance.

Chloral Microplate Assay Kit provides a convenient tool for sensitive detection of Chloralconcentration in a variety of samples. The intensity of the product color, measured at 480 nm, is proportional to the Chloral concentration in the sample.



II.KIT COMPONENTS

Component	Volume	Storage
96-Well Microplate	1 plate	
Reaction Buffer	5 ml x 1	4 °C
Dye Reagent	Powderx 1	4 °C
Dye Reagent Diluent	5 ml x 1	4 °C
Standard	Powderx 1	4 °C
Plate Adhesive Strips	3 Strips	
Technical Manual	1 Manual	

Note:

Dye Reagent: add 5 ml Dye Reagent Diluent, mix before use.

Standard: add 1 ml distilled water to dissolve before use; then add 50μ l into 950μ l distilled water, mix, the concentration will be 1 mmol/L.

III. MATERIALS REQUIRED BUT NOT PROVIDED

- 1. Microplate reader to read absorbance at 480 nm
- 2. Distilled water
- 3. Pipettor, multi-channel pipettor
- 4. Pipette tips
- 5. Centrifuge
- 6. Timer
- 7. Convection oven



IV. SAMPLE PREPARATION

1. For liquid samples

Detect directly, or dilute with distilled water.



V. ASSAY PROCEDURE

Add following reagents into the microplate:

Reagent	Sample	Standard	Blank
Sample	100μΙ		
Standard		100μΙ	
Distilled water			100μΙ
Reaction Buffer	50 μΙ	50 μΙ	50 μΙ
Dye Reagent	50 μΙ	50 μΙ	50 μΙ

Mix, cover the plate adhesive strip, incubate at 90°C for 15minutes, recordabsorbance measured at 480 nm.

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 volume of sample

Chloral (
$$\mu$$
mol/mI)=(C_{Standard}×V_{Standard})×(OD_{Sample} - OD_{Blank}) / (OD_{Standard} - OD_{Blank}) / V_{Sample}

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

 $C_{Standard}$: the concentration of standard, 1 mmol/L = 1 μ mol/ml;

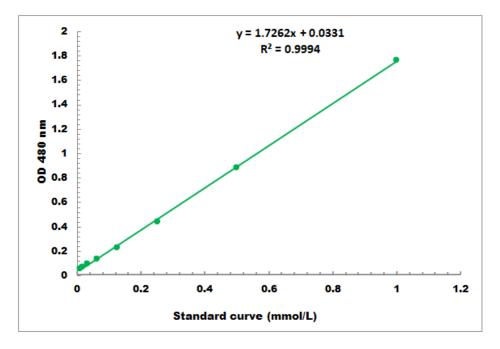
 $V_{Standard}$: the volume of standard, 0.1 ml;

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



VII. TYPICAL DATA

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



Detection Range: 0.01 mmol/L - 1 mmol/L

VIII. TECHNICAL SUPPORT

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

IX. NOTES