Butyryl Cholinesterase Activity Microplate Assay Kit

Instructions for Use | Cat. No.: ARS6864

Detection and Quantification of Butyryl Cholinesterase (BChE)Activity in Serum, Plasma, Tissue extracts, Cell lysate, Cell culture media, Other biological fluids Samples.

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

1. Introduction

Butyryl Cholinesterase (BChE , EC 3.1.1.8) is sometimes referred to as pseudocholinesterase but it preferentially uses butyrylcholine and benzoylcholine as substrates. Butyrylcholinesterase is distinct from acetylcholinesterase and is found in mammalian blood plasma, liver, pancreas, intestinal mucosa and the white matter of the central nervous system. Butyrylcholinesterase has a molecular weight of 440,000.

It is a glycoprotein which has an optimum pH of 6.0 -8.0. Assay of butyrylcholinesterase activity is of diagnostic value in various liver diseases, malignancies, and pulmonary tuberculosis. The enzyme can also be used for the assay of organophosphorous compounds such as pesticides.

Butyryl Cholinesterase ActivityColorimetric Microplate Assay Kit provides a simple and sensitive method for monitoring Butyryl Cholinesterase activity in various samples. The enzyme catalysed reaction products p-nitrophenol can be measured at a colorimetric readout at 412 nm.

2. Kit Components

Component	Volume	Storage
96-Well Microplate	1 plate	-
Assay Buffer	30mlx 4	4 °C
Reaction Buffer	20 mL x 1	4 °C
Substrate	Powderx 1	4 °C, keep in dark
Dye Reagent	Powderx 1	4 °C, keep in dark
Standard	Powderx 1	4 °C
Positive Control	Powderx 1	-20 °C
Technical Manual	1 Manual	-

3. Materials Required But Not Provided

1. Microplate reader to read absorbance at 412 nm
2. Distilled water
3. Pipettor , multi-channel pipettor
4. Pipette tips
5. Mortar
6. Centrifuge
7. Timer
8. Ice

4. Reagent Preparation

Standard: Briefly centrifuge prior to opening. Dissolve in 1 mL distilled water to generate 20 mmol/ L of standardstock solution.Store at 4°C for 1 week or -20°C for 1-2months after reconstitution. Then dilute to5mmol/L standard top solution by adding 250 µL stock solution into 750 µL distilled water. Perform 2-fold serial dilutions of the top standard solution using distilled water to make the standard curve. The concentration of standard curve could be 5/2.5/1.25/0.625/0.312/0.156/0.078mmol/L.

Substrate: Briefly centrifuge prior to opening. Add 1 mL distilled water to dissolve before use. Keep in dark and store at 4°C for 4-5 days or -20°C for 1 month after reconstitution.

Dye Reagent: Briefly centrifuge prior to opening. Add 1 mL ethanol to dissolve before use; store at 4°C for 1 week after reconstitution. Keep in dark and store at 4°C for 1-2 weeks or -20°C for 2 -3months after reconstitution.

Positive Control: Briefly centrifuge prior to opening. Add 1 mL Assay Bufferto dissolve before use; store at -80°C for 1 month after reconstitution.

Note: Divide into small aliquots to avoid repeated freeze-thaw cycles.

5. Sample Preparation

1. For cell and bacteria samples Collect cell or bacteria into centrifuge tube, discard the supernatant after centrifugation, add 1 mL Assay Buffer for 5 × 10^6 cell or bacteria, sonicate (with power 20%, sonicate 3s, interval 10s ,repeat 30 times); centrifuged at 1 0,000g 4°C for 2 0 minutes , take the supernatant into a new centrifuge tube and keep it on ice for detection.

2. For tissue samples Weigh out 0.1 g tissue, homogenize with 1 mL Assay Buffer on ice, centrifuged at 10,000g 4°C for 2 0 minutes , take the supernatant into a new centrifuge tube and keep it on ice for detection.

3. For liquid samples Detect directly.

6. Assay Procedure

Add following reagents in to the microplate :

Reagent*	Sample**	Control	Standard	Blank	Positive Control
Reaction Buffer	170µl	170µl	170µl	170µl	170µl
Substrate	10 µl	10 µl	--	--	10 µl
Sample	10 µl	--	--	--	--
Distilled water	--	10 µl	10 µl	20 µl	--
Standard	--	--	10 µl	--	--
Positive Control	--	--	--	--	10 µl
Dye Reagent	10 µl	10 µl	10 µl	10 µl	10 µl
Mix, wait for 2 minutes, record absorbance measured at 412 nm.					

Note:

*Reagents must be added sequentially and should not be premixed prior to addition.

**For unknown samples, we recommend doing a pilot experiment & testing several doses to ensure the readings are within the standard curve range. If the enzyme activity is lower, please add more samples into the reaction system; or increase the reaction time; if the enzyme activity is higher, please dilute the sample, or decrease the reaction time.

7. Calculation

Unit Definition:

One unit of Butyryl Cholinesterase activity is defined as the enzyme hydrolyze 1μmol of butyrylthiocholine iodide per minute at 25°C and pH 7.4 .

1. According to the slope of the standard curve

$$\text{Activity} = \frac{(OD_{Sample} - OD_{Control}) - \text{Intercept}}{\text{Slope} \times T} \times n \ (U/mL)$$

2. According to one point of the standard OD value and concentration

2.1 According to the protein concentration of sample

$$\text{Activity} = \frac{(C_{Standard} \times V_{Standard}) \times (OD_{Sample} - OD_{Control})}{(OD_{Standard} - OD_{Blank}) \times V_{Sample} \times C_{Protein} \times T} \ (U/mg)$$

2.2According to the quantity of cells or bacteria

$$\text{Activity} = \frac{(C_{Standard} \times V_{Standard}) \times (OD_{Sample} - OD_{Control})}{(OD_{Standard} - OD_{Blank}) \times N \times (V_{Sample} / V_{Assay}) \times T} \ (U/10^4)$$

2.3 According to the weight of sample

$$\text{Activity} = \frac{(C_{Standard} \times V_{Standard}) \times (OD_{Sample} - OD_{Control})}{(OD_{Standard} - OD_{Blank}) \times W \times (V_{Sample} / V_{Assay}) \times T} \ (U/g)$$

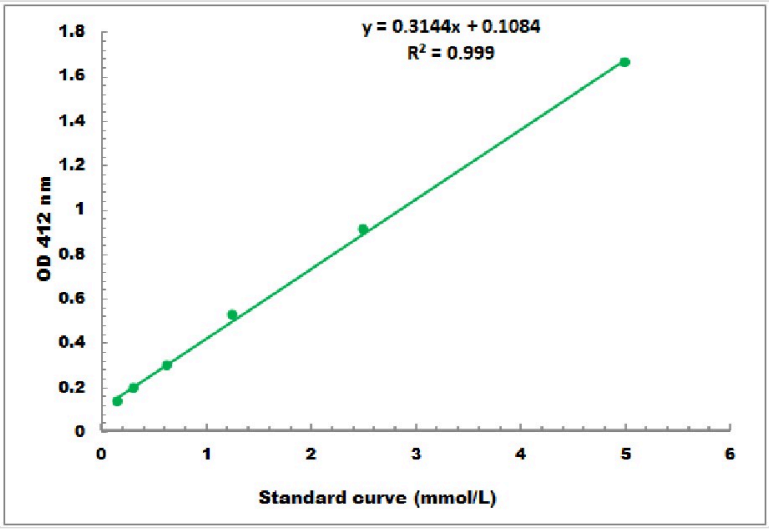
2.4 According to the volume of sample

$$\text{Activity} = \frac{(C_{Standard} \times V_{Standard}) \times (OD_{Sample} - OD_{Control})}{(OD_{Standard} - OD_{Blank}) \times V_{Sample} \times T} \ (U/mL)$$

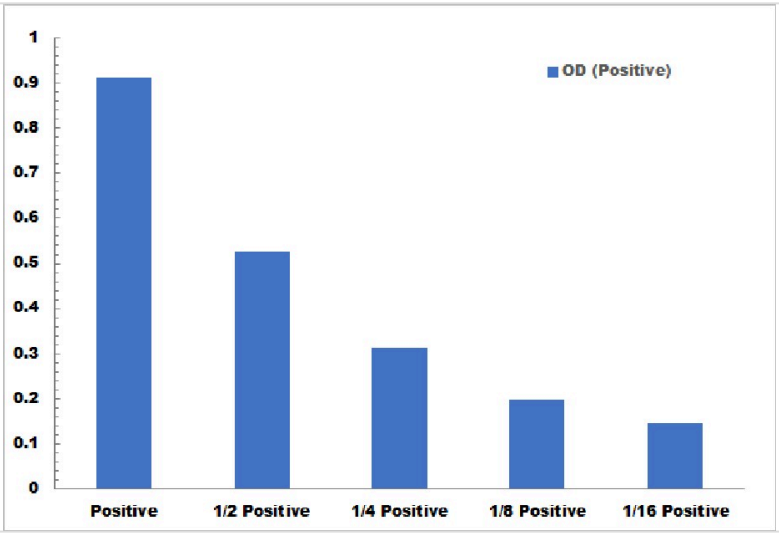
Item	Definition
Slope	the absorbance slope of standard curve
n	the dilution factor
$C_{Protein}$	the protein concentration of sample, mg/mL
W	the weight of total sample , g
N	the quantity of total cell or bacteria sample, 10^4
$C_{Standard}$	the concentration of standard, mmol/L = μmol/mL
$V_{Standard}$	the volume of standard in assay procedure, mL
V_{Sample}	the volume of sample in assay procedure, mL
V_{Assay}	the volume of Assay Buffer in sample preparation , mL
T	the reaction time , minute

8. Typical Data

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



Detection Range : 0.1 mmol/L - 5 mmol/L



Positive Control reaction in 96 -well plate assay with decreasing the concentration