Phospholipase D Assay Kit

Catalog Number KA1636
100 assays
Version: 04

Intended for research use only
# Table of Contents

**Introduction** .................................................................................................................. 3

- Intended Use .................................................................................................................. 3

- Background .................................................................................................................... 3

- Principle of the Assay .................................................................................................... 3

**General Information** ..................................................................................................... 4

- Materials Supplied ....................................................................................................... 4

- Storage Instruction ........................................................................................................ 4

- Materials Required but Not Supplied .......................................................................... 4

- Precautions for Use ...................................................................................................... 4

**Assay Protocol** ............................................................................................................... 5

- Assay Procedure ........................................................................................................ 5

**Data Analysis** ............................................................................................................... 6

- Calculation of Results .................................................................................................. 6

**Resources** ..................................................................................................................... 7

- References .................................................................................................................... 7
Introduction

Intended Use

Applications:

- Direct Assays: phospholipase D in biological samples.
- Drug Discovery/Pharmacology: effects of drugs on phospholipase D metabolism.

Features:

- Sensitive. Use 10 μL samples. Detection range: colorimetric assay 0.06 - 10 U/L, fluorimetric assay 0.04 - 1 U/L.
- Simple and High-throughput: the assay involves addition of a single working reagent and can be readily adapted to high-throughput assays for drug screening.

Background

PHOSPHOLIPASE D (PLD) catalyses the hydrolysis of the phosphodiester bond of glycerophospholipids to generate phosphatidic acid and a free headgroup. Abnormalities in PLD expression have been associated with human cancers.

Principle of the Assay

Phospholipase D Assay Kit provides a simple high-throughput assay for measuring PLD activity. In this assay, PLD hydrolyzes phosphatidylcholine to choline which is determined using choline oxidase and a H₂O₂ specific dye. The optical density of the pink colored product at 570 nm or fluorescence intensity (530/585 nm) is directly proportional to the PLD activity in the sample.
General Information

Materials Supplied

List of component

<table>
<thead>
<tr>
<th>Component</th>
<th>Amount</th>
</tr>
</thead>
<tbody>
<tr>
<td>Assay Buffer</td>
<td>10 mL</td>
</tr>
<tr>
<td>Dye Reagent</td>
<td>120 μL</td>
</tr>
<tr>
<td>Enzyme Mix (Dried)</td>
<td>1 vial</td>
</tr>
<tr>
<td>Substrate</td>
<td>1.5 mL</td>
</tr>
<tr>
<td>Calibrator</td>
<td>400 μL</td>
</tr>
</tbody>
</table>

Storage Instruction

Store all components at -20°C. Shelf life of six months after receipt.

Materials Required but Not Supplied

Pipetting devices, centrifuge tubes, clear flat-bottom uncoated 96-well plates, optical density plate reader; black flat-bottom uncoated 96-well plates, fluorescence plate reader.

Precautions for Use

Reagents are for research use only. Normal precautions for laboratory reagents should be exercised while using the reagents.
Assay Protocol

Assay Procedure

✓ Colorimetric assay

Liquid samples can be assayed directly. Solid samples should be homogenized in a suitable enzyme buffer prior to assay.

*Note: SH-containing reagents (e.g. β-mercaptoethanol, dithiothreitol, > 5 μM), sodium azide, EDTA, and sodium dodecyl sulfate are known to interfere in this assay and should be avoided in sample preparation. If a sample is known to contain choline, it should be removed by dialysis or membrane filtration.*

1. Equilibrate all components to room temperature. Briefly centrifuge the tubes before opening. Keep thawed tubes on ice during assay. Reconstitute Enzyme Mix with 120 μL Assay Buffer. Reconstituted Enzyme Mix is stable for 1 month when stored at -20°C. *Note: a yellow precipitate may form after thawing reconstituted Enzyme Mix. If a precipitate forms, pellet it by centrifuging for 2 min at 14000 rpm and use the clear supernatant.*

2. Calibrator: mix 33 μL Calibrator with 187 μL dH₂O (final 300 μM choline). Dilute calibrator in dH₂O as follows.

<table>
<thead>
<tr>
<th>No</th>
<th>300 μM Premix + H₂O</th>
<th>Vol (μL)</th>
<th>Calibrator (μM)</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>100 μL + 0 μL</td>
<td>100</td>
<td>300</td>
</tr>
<tr>
<td>2</td>
<td>60 μL + 40 μL</td>
<td>100</td>
<td>180</td>
</tr>
<tr>
<td>3</td>
<td>30 μL + 70 μL</td>
<td>100</td>
<td>90</td>
</tr>
<tr>
<td>4</td>
<td>0 μL + 100 μL</td>
<td>100</td>
<td>0</td>
</tr>
</tbody>
</table>

Transfer 10 μL diluted standards into separate wells of a clear flat bottom 96-well plate. Samples: transfer 10 μL of each sample into separate wells of the plate.

3. Color reaction. Prepare enough Working Reagent by mixing, for each well, 85 μL Assay Buffer, 1 μL Enzyme Mix, 1 μL Dye Reagent and 12 μL Substrate. Add 90 μL Working Reagent to each well. Tap plate to mix. Incubate at desired temperature and protect from light. At 10 and 30 min, read optical density 570 nm (550-585 nm).

✓ Fluorimetric assay

The fluorimetric assay procedure is similar to the colorimetric procedure except that (1) 0, 9, 18 and 30 μM calibrator and (2) a black 96-well plate are used. Read fluorescence intensity at λ<sub>ex</sub> = 530 nm and λ<sub>em</sub> = 585 nm.
Data Analysis

Calculation of Results

Subtract blank value (#4) from the standard values and plot the ΔOD or ΔF against standard concentrations. Determine the slope and calculate the PLD activity of Sample,

\[
\text{[Phospholipase D]} = \frac{R_{30} - R_{10}}{\text{Slope \times 20}} \times n \text{ (U/L)}
\]

\( R_{30} \) and \( R_{10} \) are optical density or fluorescence intensity readings of the Sample at 30 min and 10 min, respectively. 20 is the enzyme reaction time (30 min - 10 min). \( n \) is the sample dilution factor. Note: if the calculated PLD activity of a sample is higher than 10 U/L in the Colorimetric Assay or 1 U/L in the Fluorimetric Assay, dilute sample in assay buffer and repeat the assay. Multiply result by the dilution factor.

Unit definition: 1 unit of PLD catalyzes formation of 1 μmole of choline per minute under the assay conditions (pH 7.4).
Resources

References

