

Catalog No: DPG100K

Catalog No: DPG100KE

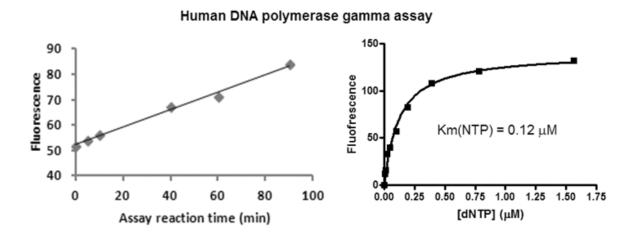
INSTRUCTIONS

ProFoldin Human DNA Polymerase Gamma Assay Kits

Human DNA Polymerase Gamma Assay Kit Human DNA Polymerase Gamma Assay Kit Plus

Introduction

Human DNA polymerase gamma is the only DNA polymerase in human mitochondria and is responsible for the DNA replication, recombination and repairing in human mitochondria. It plays critical roles in mitochondrial diseases and aging. Some antiviral nucleoside analogs were reported to inhibit DNA polymerase gamma after intracellular phosphorylation and cause severe chronic toxicity. The human DNA polymerase gamma assay is based on measurement of the DNA molecules synthesized by the DNA polymerase. The assay is performed in a 384-well or 96-well plate format. The assay can be used for detection of DNA polymerase gamma activity and high throughput screen of human DNA polymerase gamma inhibitors.



The **Human DNA Polymerase Gamma Assay Kit (Catalog No. DPG100K)** includes all the assay kit components except the enzyme for 100 assays in a 384-well plate assay format: 400 μ l of 10 x Buffer, 33 μ l of 100 x DNA template (3 μ M), 33 μ l of 100 x dNTP mix (1 mM), 1550 μ l of 2 x Dye, 1550 μ l of 50 mM EDTA.

The Human DNA Polymerase Gamma Assay Kit Plus (Catalog No. DPG100KE) includes all the assay kit components for 100 assays in 384-well plate assay format: 400 μ l of 10 x Buffer, 33 μ l of 100 x DNA template (3 μ M), 33 μ l of 100 x dNTP mix (1 mM), 33 μ l of 100 x human DNA polymerase gamma, 1550 μ l of 2 x Dye, 1550 μ l of 50 mM EDTA.

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Assay protocol

The following assay protocol is based on the 384-well plate assay format (plate type: Matrix 4318 or alike). The reaction volume is 30 μ l and the final assay volume is 60 μ l. For 96-well plate assays (plate type: Costar 3915 or alike), the reaction volume is 60 μ l and the final assay volume is 120 μ l.

1. Reagent preparation:

- (1) 10 x DNA: Dilute the 100 x DNA 10-fold with water. Each assay uses 3 µl of 10 x DNA.
- (2) 10 x enzyme: Dilute the 100 x Human DNA polymerase gamma 10-fold with the 1 x assay buffer. Each assay uses 3 µl of 10 x enzyme.
- (3) 10 x dNTP mix: Dilute the 100 x dNTP mix 10-fold with water. Each assay uses 3 μl of 10 x dNTP mix.
- (4) 1 x dye: Dilute the 2 x fluorescence dye 2-fold with 50 mM EDTA. Each assay uses 30 µl of 1 x dye.

2. **Reaction**:

The total volume of each reaction mixture is 30 μ l including 18 μ l of H₂O, 3 μ l of 10 x Buffer, 3 μ l of 10 x DNA template, 3 μ l of 10 x enzyme, 3 μ l of 10 x dNTP mix. Incubate the reaction mixture at 37°C for 60 min.

3. **Detection**:

Mix 30 μ l of the 1 x fluorescence dye with 30 μ l of the reaction mixture. Measure the fluorescence intensity at 535 nm using the excitation wavelength at 485 nm.

Assay Protocol for enzyme inhibition

The assay can be optimized in terms of assay window, assay linearity and sensitivity to competitive inhibitors. ProFoldin offers HTS assay development service. For more information, please visit our website at http://www.profoldin.com/services.html.

Related Products

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