DNA Polymerase γ (POLG), His-FLAG-Tag Recombinant

Catalog: 21001 Lot: 230309-1

Product Information

Description: Recombinant human DNA polymerase γ (gamma), full length encompassing amino

acids 2-1239 (end). This construct contains an N-terminal His-tag (6xHis) followed by a

FLAG-tag. The recombinant protein was affinity purified.

Background: Polymerases are the enzymes responsible for synthesizing nucleic acids. Polymerase γ,

also known as DNA polymerase subunit gamma, POLG or POLG1, belongs to the Family A of DNA polymerases and corresponds to the catalytic subunit of the mitochondrial DNA polymerase. It works in conjunction with POLG2 to replicate mitochondrial DNA. In addition to its DNA polymerase activity, it has a 3'-5' exonuclease and a 5'dRP lyase activity for proofreading and repair. Mutations in polymerase γ cause mitochondrial dysfunction, as observed in SANDO (sensory ataxia neuropathy dysarthia and ophthalmoparesis), AHS (Alpers-Huttenlocher Syndrome) and MNGIE (mitochondrial neurogastrointestinal encephalopathy syndrome). Alternatively, the inhibition of DNA synthesis can be used as a therapeutic approach for diseases in which cell division or energetic needs are uncontrolled, such as cancer. Further studies into DNA polymerase

inhibitors will bring new therapy options to cancer patients.

Species: Human

Construct: DNA Polymerase γ (His-FLAG-2-1239(end))

Concentration: 0.59 mg/ml

Expression System: Sf9
Purity: ≥90%

Format: Aqueous buffer solution.

Formulated In: 40 mM Tris-HCl, pH 8.0, 110 mM NaCl, 2.2 mM KCl, 0.04% Tween-20, 20% glycerol, and

100 μg/ml FLAG peptide

MW: 141 kDa Genbank Accession: NM 002693

Stability: At least 6 months at -80°C.

Storage: -80°C

Instructions for Use: Thaw on ice and gently mix prior to use. DO NOT VORTEX. Perform a quick spin before

opening. Aliquot into small volumes and flash freeze for long term storage. Avoid

multiple freeze/thaw cycles.

Assay Conditions: Assay was performed using Human DNA Polymerase Assay Kit (ProFoldin #DPB100K)

and with various amounts of DNA Polymerase γ enzyme. The assay was run in 384-well format with a reaction volume of 15 μ l supplemented with 15 μ l of detection reagent (total 30 μ l) prior plate reading. Plate was incubated for 1 hour at 37°C prior detection.

Applications: Useful for the study of enzyme kinetics, screening inhibitors, and selectivity profiling.

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Quality Control Data

