INSTRUCTIONS

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S. aureus DNA Helicase ATPase Assay Kits

S. aureus DNA Helicase ATPase assay Kit Plus-100

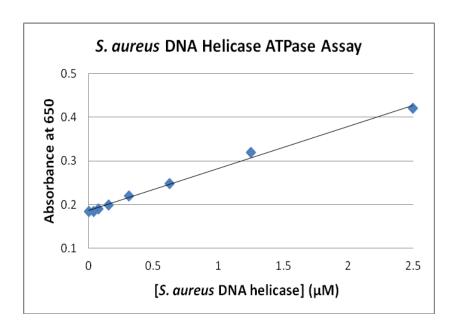
Catalog No. DNAB100KS

S. aureus DNA Helicase ATPase assay Kit Plus-500

Catalog No. DNAB500KS

Introduction

DNA helicase (DnaB) hydrolyzes ATP as the source of molecular energy to carry out DNA unwinding required by the DNA replication process. Inhibition of the ATPase activity of DNA helicase blocks its DNA unwinding function. The DNA helicase ATPase assay can be used for high-throughput screen of DNA helicase inhibitors in drug discovery. The **DNA Helicase ATPase Assay Kit** is based on detection of the phosphate produced by the ATP hydrolysis reaction in the presence of DNA. The assay is in a 384-well plate format and the phosphate is detected using light absorbance at 650 nm.



The *S. aureus* **DNA Helicase ATPase assay Kit Plus-100** (Catalog No. DNAB100KS) includes 500 μ l of 10 x assay buffer, 35 μ l of 100 x DNA, 150 μ l of 20 x *S. aureus* DNA helicase, 35 μ l of 100 x ATP and 5 ml of dye for 100 assays in a 384-well assay format.

The *S. aureus* **DNA Helicase ATPase assay Kit Plus-500** (Catalog No. DNAB500KS) includes 2 ml of 10 x assay buffer, 170 μl of 100 x DNA, 750 μl of 20 x *S. aureus* DNA helicase, 170 μl of 100 x ATP and 25 ml of dye for 500 assays in a 384-well assay format.

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Assay Protocol

1. Reagent preparation:

For each 10 assay reactions, prepare

- (1) 297 μl of premix composed of 247.5 μl of H₂O, 33 μl of 10 x Buffer and 16.5 μl of 20 x *S. aureus* DNA helicase;
- (2) 33 µl of 10 x Enzyme substrate by mixing 3.3 µl of 100 x ATP and 3.3 µl of 100 x DNA and 26.4 µl of water.

2. Reaction:

Mix 27 μ l of the premix with 3 μ l of the 10 x Enzyme substrate in each well. Incubate the reaction mixture at 37°C for 90 min.

3. Detection:

Add 45 µl of the Dye MPA3000 into the 30 µl of the reaction mixture. Incubate for 5 min. Measure the light absorbance at 650 nm.

Note: The final concentrations for the ATPase assays of the helicases are 20 mM HEPES, pH 7.5, 20 mM potassium glutamate, 1 mM DTT, 0.005% Triton X-100, 10 mM MgCl₂, 20 μ g/ml DNA, 0.25 mM ATP and 2 μ M DNA helicase. A negative control reaction can be the reaction mixture without addition of ATP or enzyme.

Assay optimization 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

| E. coli DNA Helicase ATPase assay Kit Plus-100 | Catalog No. DNAB100KE |
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| H. influenzae DNA Helicase ATPase assay Kit Plus-100 | Catalog No. DNAB100KH |
| S. pneumoniae DNA Helicase ATPase assay Kit Plus-100 | Catalog No. DNAB100KN |
| P. aeruginosa DNA Helicase ATPase assay Kit Plus-100 | Catalog No. DNAB100KP |
| E. coli DNA Primase Assay Kit Plus | Catalog No. EGA100KE |
| H. influenzae DNA Primase Assay Kit Plus | Catalog No. HGA100KE |
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