

For research use only ISO9001

Phi29 DNA Polymerase

Product	Quantity	Cat. No.	Remarks
Phi29 DNA Polymerase	500 unit	EBT-3004	10 unit/μℓ

Description

Phi29 DNA Polymerase is one of the enzymes showing strong strand displacement DNA synthetic activity. The polymerase has an inherent $3'\rightarrow 5'$ proof-reading exonuclease activity. The polymerase gene from the *Bacillus subtilis* phage phi29 (Φ 29) was overexpressed and purified from *E.coli* strain.

- Extreme strand displacement: best fit for RCA (rolling circle amplification) and WGA by MDA (whole genome amplification by multiple displacement amplification)
- High fidelity: 3'→5' proof-reading exonuclease activity
- Extreme processivity : up to 70 kbp

Concentration & Storage Condition

10 unit/μl. Store at -20℃.

Storage Buffer

10 mM Tris-HCl, pH 7.5, 100 mM KCl, 0.1 mM EDTA, 1 mM DTT, 0.1% Triton X-100, 50% glycerol.

10x Reaction Buffer

500 mM Tris-HCl, pH 7.5, 100 mM (NH₄)₂SO₄ 100 mM MgCl₂ 40 mM DTT.

Unit Definition

One unit is defined as the amount of enzyme required to incorporate 0.5 pmol of dNTP into acid insoluble material in 10 minutes at 30°C

QC Tests

Activity, exo and endonuclease activity test, DNA contamination test, SDS-PAGE purity, performance tests.



ISO9001

For research use only

Phi29 DNA Polymerase

Product	Quantity	Cat. No.	Remarks	
Phi29 DNA Polymerase	500 unit	EBT-3004	10 unit/μℓ	

Description

Phi29 DNA Polymerase is one of the enzymes showing strong strand displacement DNA synthetic activity. The polymerase has an inherent $3'\rightarrow 5'$ proof-reading exonuclease activity. The polymerase gene from the *Bacillus subtilis* phage phi29 (Φ 29) was overexpressed and purified from *E.coli* strain.

- Extreme strand displacement: best fit for RCA (rolling circle amplification) and WGA by MDA (whole genome amplification by multiple displacement amplification)
- High fidelity: 3´→5′ proof-reading exonuclease activity
- Extreme processivity : up to 70 kbp

Concentration & Storage Condition

10 unit/µl. Store at -20℃.

Storage Buffer

10 mM Tris-HCl, pH 7.5, 100 mM KCl, 0.1 mM EDTA, 1 mM DTT, 0.1% Triton X-100, 50% glycerol.

10x Reaction Buffer

500 mM Tris-HCl, pH 7.5, 100 mM (NH₄)₂SO₄ 100 mM MgCl₂ 40 mM DTT.

Unit Definition

One unit is defined as the amount of enzyme required to incorporate 0.5 pmol of dNTP into acid insoluble material in 10 minutes at 30°C

QC Tests

Activity, exo and endonuclease activity test, DNA contamination test, SDS-PAGE purity, performance tests.

