



1-800-632-7799 info@neb.com www.neb.com

BioLabs.

1-800-632-7799

info@neb.com

www.neb.com

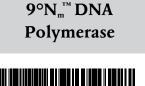


2.000 U/ml 200 units Lot: 0031209

RECOMBINANT Store at -20°C Exp: 9/14

Description: 9°N DNA Polymerase is a thermophilic DNA polymerase that has been genetically engineered to have a decreased (1% - 5%) of the wildtype) $3' \rightarrow 5'$ proofreading exonuclease activity. 9°N_DNA Polymerase features a half-life of 6.7 hours at 95°C.

Source: An E. coli strain that carries the 9°N (E143D) DNA Polymerase gene (1.2), a genetically engineered form of the native DNA polymerase from the extremely thermophilic marine archaea







200 units 2,000 U/ml Lot: 0031209 RECOMBINANT Store at -20°C Exp: 9/14

Description: 9°N DNA Polymerase is a thermophilic DNA polymerase that has been genetically engineered to have a decreased (1% - 5%) of the wildtype) $3' \rightarrow 5'$ proofreading exonuclease activity. 9°N __ DNA Polymerase features a half-life of 6.7 hours at 95°C.

Source: An E. coli strain that carries the 9°N (E143D) DNA Polymerase gene (1.2), a genetically engineered form of the native DNA polymerase from the extremely thermophilic marine archaea

Thermococcus species 9°N-7. The archaea was isolated from a submarine thermal vent. at a depth of 2.500 meters. 9° north of the equator at the East Pacific Rise (3).

Supplied in: 100 mM KCl, 0.1 mM EDTA, 10 mM Tris-HCI (pH 7.4), 1 mM DTT and 50% glycerol.

Applications:

- Primer extension
- SNP Analysis

Reagents Supplied with Enzyme:

10X ThermoPol[™] Reaction Buffer.

Reaction Conditions: 1X ThermoPol Reaction Buffer, 200 µM each dNTP, DNA template, primer and 1-2 units 9°N_ DNA Polymerase in a total reaction volume of 100 µl.

1X ThermoPol Reaction Buffer:

20 mM Tris-HCI 10 mM (NH₄)₂SO₄ 10 mM KCl 2 mM MgSO 0.1% Triton® X-100 pH 8.8 @ 25°C

Thermococcus species 9°N-7. The archaea was isolated from a submarine thermal vent. at a depth of 2,500 meters, 9° north of the equator at the East Pacific Rise (3).

Supplied in: 100 mM KCl. 0.1 mM EDTA. 10 mM Tris-HCI (pH 7.4), 1 mM DTT and 50% glycerol.

Applications:

- Primer extension
- SNP Analysis

Reagents Supplied with Enzyme:

10X ThermoPol[™] Reaction Buffer.

Reaction Conditions: 1X ThermoPol Reaction Buffer, 200 µM each dNTP, DNA template, primer and 1-2 units 9°N_ DNA Polymerase in a total reaction volume of 100 µl.

1X ThermoPol Reaction Buffer:

20 mM Tris-HCI 10 mM (NH₄)₂SO₄ 10 mM KCl 2 mM MgSO 0.1% Triton® X-100 pH 8.8 @ 25°C

Unit Definition: One unit is defined as the amount of enzyme that will incorporate 10 nmol of dNTP into acid-insoluble material in 30 minutes at 75°C.

Unit Assav Conditions: 1X ThermoPol Reaction Buffer, 200 µM dNTPs including [³H]-dTTP and 15 nM primed single-stranded M13mp18.

Heat Inactivation: No

Quality Control Assays

Endonuclease Activity: Incubation of a 50 µl reaction in ThemoPol Reaction Buffer supplemented with 400 uM each dNTP containing a minimum of 20 units of 9°N, DNA Polymerase with 1 μ g of supercoiled ϕ X174 DNA for 4 hours at either 37°C or 75°C results in < 10% conversion to the nicked form as determined by agarose gel electrophoresis.

Notes on use:

It is suggested that the number of units be optimized with each primer:template.

Unit Definition: One unit is defined as the amount

into acid-insoluble material in 30 minutes at 75°C.

of enzyme that will incorporate 10 nmol of dNTP

Unit Assay Conditions: 1X ThermoPol Reaction

Buffer. 200 µM dNTPs including [3H]-dTTP and

15 nM primed single-stranded M13mp18.

Endonuclease Activity: Incubation of a

supplemented with 400 µM each dNTP

50 ul reaction in ThemoPol Reaction Buffer

containing a minimum of 20 units of 9°N., DNA

DNA for 4 hours at either 37°C or 75°C results

It is suggested that the number of units be opti-

Polymerase with 1 μ g of supercoiled ϕ X174

in < 10% conversion to the nicked form as

determined by agarose gel electrophoresis.

mized with each primer:template.

Heat Inactivation: No

Notes on use:

Quality Control Assays

(see other side)

CERTIFICATE OF ANALYSIS

References:

References:

447-462

#B1003S

Diluent E

#B8005S

#B9004S

#B9005S

#B9013S

Pack

1. Southworth, M. W. et al. (1996) Proc. Natl.

2. Rodriguez, A. C. et al. (2000) J. Mol. Biol.

3. Thermococcus sp. (strain 9°N-7) isolated by

Dr. Holger Jannasch, Woods Hole Oceano-

6.0 ml

4.0 ml

6.0 ml

6.0 ml

6.0 ml

ThermoPol II (Mg-free) Reaction Buffer Pack

ThermoPol DF (Detergent-free) Reaction Buffer

Acad. Sci. USA, 5281-5285.

graphic Institute, 1991.

Companion Products Sold Separately:

Magnesium Sulfate (MgSO₄) Solution

ThermoPol Reaction Buffer Pack

- 1. Southworth, M. W. et al. (1996) Proc. Natl. Acad. Sci. USA, 5281-5285.
- 2. Rodriquez, A. C. et al. (2000) J. Mol. Biol. 447-462
- 3. Thermococcus sp. (strain 9°N-7) isolated by Dr. Holger Jannasch, Woods Hole Oceanographic Institute, 1991.

Companion Products Sold Separately:

Magnesium Sulfate (MgSO₄) Solution #B1003S 6.0 ml Diluent E #B8005S 4.0 ml ThermoPol Reaction Buffer Pack #B9004S 6.0 ml ThermoPol II (Mg-free) Reaction Buffer Pack #B9005S 6.0 ml ThermoPol DF (Detergent-free) Reaction Buffer Pack #B9013S 6.0 ml

(see other side)



DeoxynucleotideSolutionSet#N0446S25 μmol eachDeoxynucleotideSolution#N0447S8 μmol each#N0447L40 μmol each

9°N_m[™] and THERMOPOL[™] are trademarks of New England Biolabs, Inc.

TRITON® is a registered trademark of Union Carbide Corporation.



Page 2 (M0260)

Deoxynucleotide Solution Set #N0446S 25 µmol each Deoxynucleotide Solution Mix #N0447S 8 µmol each #N0447L 40 µmol each

9°N_m[™] and THERMOPOL[™] are trademarks of New England Biolabs, Inc.

TRITON® is a registered trademark of Union Carbide Corporation.

