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Date

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New England Biolabs Product Specification

Product Name: Exonuclease VIII, truncated

Catalog #: M0545S/L
Concentration: 10,000 units/ml

Unit Definition:

One unit is defined as the amount of enzyme required to produce 1 nmol of acid-soluble deoxyribonucleotide from double-

stranded DNA in a total reaction volume of 50 μ l in 30 minutes at 37°C in 1X NEBuffer 4 with 0.15 mM sonicated

 $duplex [^3H]-DNA.$

Shelf Life: 24 months
Storage Temp: -20°C

Storage Conditions: 50 mM Tris-HCl, 100 mM NaCl, 1 mM DTT, 0.1 mM EDTA, 50 % Glycerol, 0.1 % Triton X-100, (pH 7.5 @ 25°

C)

Specification Version: PS-M0545S/L v1.0
Effective Date: 22 May 2018

Assay Name/Specification (minimum release criteria)

Endonuclease Activity (Circular Single Stranded DNA) - A 50 μ l reaction in NEBuffer 4 containing 1 μ g of M13mp18 Single-stranded DNA and a minimum of 30 units of Exonuclease VIII, truncated incubated for 4 hours at 37°C results in <10% conversion to linear DNA as determined by agarose gel electrophoresis.

Endonuclease Activity (Nicked Circular DNA) - A 50 μ l reaction in NEBuffer 4 containing 1 μ g of PhiX174 RF II DNA and a minimum of 30 units of Exonuclease VIII, truncated incubated for 4 hours at 37°C results in <10% conversion to linear DNA as determined by agarose gel electrophoresis.

Endonuclease Activity (Nicking) - A 50 μ l reaction in NEBuffer 4 containing 1 μ g of supercoiled PhiX174 DNA and a minimum of 50 units of Exonuclease VIII, truncated incubated for 4 hours at 37°C results in <10% conversion to the nicked form as determined by agarose gel electrophoresis.

Protein Purity Assay (SDS-PAGE) - Exonuclease VIII, truncated is ≥ 95% pure as determined by SDS-PAGE analysis using Coomassie Blue detection.

RNase Activity (Extended Digestion) - A 10 μ l reaction in NEBuffer 4 containing 40 ng of a 300 base single-stranded RNA and a minimum of 10 units of Exonuclease VIII, truncated is incubated at 37°C. After incubation for 4 hours, >90% of the substrate RNA remains intact as determined by gel electrophoresis using fluorescent detection.

Derek Robinson

Director of Quality Control







22 May 2018