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Date

30 Sep 2016

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New England Biolabs Product Specification

Product Name: T4 RNA Ligase 1 (ssRNA Ligase)

Catalog #: M0204S/L
Concentration: 10,000 units/ml

Unit Definition: One unit is defined as the amount of enzyme required to convert 1 nanomole of 5'-[32P] rA16 into a phosphatase-resistant

form in 30 minutes at 37°C.

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

Storage Conditions: 50 mM KCl, 10 mM Tris-HCl, 1 mM DTT, 0.1 mM EDTA, 50 % Glycerol, (pH 7.4 @ 25°C)

Specification Version: PS-M0204S/L v1.0

Effective Date: 30 Sep 2016

Assay Name/Specification (minimum release criteria)

Endonuclease Activity (Nicking) - A 50 μ L reaction in T4 RNA Ligase 1 Reaction Buffer containing 1 μ g of supercoiled PhiX174 DNA and a minimum of 30 units of T4 RNA Ligase 1 (ssRNA Ligase) incubated for 4 hours at 37°C results in <10% conversion to the nicked form as determined by agarose gel electrophoresis.

Exonuclease Activity (Radioactivity Release) - A 50 μ l reaction in T4 RNA Ligase 1 Reaction Buffer containing 1 μ g of a mixture of single and double-stranded [3 H] *E. coli* DNA and a minimum of 30 units of T4 RNA Ligase 1 (ssRNA Ligase) incubated for 4 hours at 37°C releases <0.1% of the total radioactivity.

Protein Purity Assay (SDS-PAGE) - T4 RNA Ligase 1 (ssRNA Ligase) is ≥ 95% pure as determined by SDS-PAGE analysis using Coomassie Blue detection.

qPCR DNA Contamination (*E. coli* Genomic) - A minimum of 10 units of T4 RNA Ligase 1 (ssRNA Ligase) is screened for the presence of *E. coli* genomic DNA using SYBR® Green qPCR with primers specific for the *E. coli* 16S rRNA locus. Results are quantified using a standard curve generated from purified *E. coli* genomic DNA. The measured level of *E. coli* genomic DNA contamination is ≤ 1 *E. coli* genome.

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 1 μ l of T4 RNA Ligase 1 (ssRNA Ligase) is incubated at 37°C. After incubation for 16 hours, >90% of the substrate RNA remains intact as determined by gel electrophoresis using fluorescent detection.

Derek Robinson

Director of Quality Control





