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New England Biolabs Certificate of Analysis

Product Name: mRNA Decapping Enzyme

Catalog Number: M0608S
Concentration: 100,000 U/ml

Unit Definition: One unit is defined as the amount of mRNA Decapping Enzyme required

to convert 50% of a 500 nM m7G-capped substrate to a

5'-monophosphorylated form in a total reaction volume of 20 µl in 1

hour at 37°C.

Packaging Lot Number: 10140320
Expiration Date: 03/2024
Storage Temperature: -20°C

Storage Conditions: 10 mM Tris-HCI, 300 mM NaCI, 1 mM DTT, 0.1 mM EDTA, 50 %

Glycerol, (pH 7.4 @ 25°C)

Specification Version: PS-M0608S v2.0

mRNA Decapping Enzyme Component List			
NEB Part Number	Component Description	Lot Number	Individual QC Result
M0608SVIAL	mRNA Decapping Enzyme	10140319	Pass
B0608SVIAL	10X MDE Decapping Buffer	10122104	Pass

Assay Name/Specification	Lot # 10140320
Endonuclease Activity (Nicking) A 50 μl reaction in 1X mRNA Decapping Enzyme Reaction Buffer containing 1 μg of supercoiled PhiX174 DNA and a minimum of 100 units of mRNA Decapping Enzyme incubated for 4 hours at 37°C results in <10% conversion to the nicked form as determined by agarose gel electrophoresis.	Pass
Phosphatase Activity (pNPP) A 200 µl reaction in 1M Diethanolamine, pH 9.8, 0.5 mM MgCl2 containing 2.5 mM p-Nitrophenyl Phosphate (pNPP) and a minimum of 100 units of mRNA Decapping Enzyme incubated for 4 hours at 37°C yields <0.00001 unit of alkaline phosphatase activity as determined by spectrophotometric analysis.	Pass
Exonuclease Activity (Radioactivity Release) A 50 μl reaction in 1X mRNA Decapping Enzyme Reaction Buffer containing 1 μg of a mixture of single and double-stranded [³H] E. coli DNA and a minimum of 100 units of mRNA Decapping Enzyme incubated for 4 hours at 37°C releases <0.1% of the total radioactivity.	Pass



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Assay Name/Specification	Lot # 10140320
RNase Activity (Extended Digestion) A 10 μ I reaction in NEBuffer 4 containing 40 ng of a 300 base single-stranded RNA and a minimum of 100 units of mRNA Decapping Enzyme 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.	Pass
Protein Purity Assay (SDS-PAGE) mRNA Decapping Enzyme is ≥ 95% pure as determined by SDS-PAGE analysis using Coomassie Blue detection.	Pass

This product has been tested and shown to be in compliance with all specifications.

One or more products referenced in this document may be covered by a 3rd-party trademark. Please visit www.neb.com/trademarks for additional information.

Bhairavi Jani Production Scientist

09 Mar 2022

Michael Tonello

Packaging Quality Control Inspector

09 Mar 2022



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