

240 County Road Ipswich, MA 01938-2723 Tel 978-927-5054 Fax 978-921-1350 www.neb.com info@neb.com

## New England Biolabs Certificate of Analysis

Product Name:	Mlyl
Catalog Number:	R0610L
Concentration:	10,000 U/ml
Unit Definition:	One unit is defined as the amount of enzyme required to digest 1 μg of Lambda DNA in rCutSmart Buffer in 1 hour at 37°C in a total reaction volume of 50 μl.
Packaging Lot Number:	10247972
Expiration Date:	06/2026
Storage Temperature:	-20°C
Storage Conditions:	10 mM Tris-HCl, 50 mM KCl, 1 mM DTT, 0.1 mM EDTA, 50% Glycerol, 200 µg/ml rAlbumin (pH 7.4 @ 25°C)
Specification Version:	PS-R0610S/L v2.0

Mlyl Component List				
NEB Part Number	Component Description	Lot Number	Individual QC Result	
R0610LVIAL	Mlyl	10245037	Pass	
B6004SVIAL	rCutSmart™ Buffer	10241728	Pass	

Assay Name/Specification	Lot # 10247972
Exonuclease Activity (Radioactivity Release) A 50 µl reaction in rCutSmart <sup>™</sup> Buffer containing 1 µg of a mixture of single and double-stranded [ <sup>3</sup> H] E. coli DNA and a minimum of 30 units of MlyI incubated for 4 hours at 37°C releases <0.1% of the total radioactivity.	Pass
Functional Testing (15 minute Digest) A 50 µl reaction in rCutSmart™ Buffer containing 1 µg of Lambda DNA and 1 µl of Mlyl incubated for 15 minutes at 37°C results in complete digestion as determined by agarose gel electrophoresis.	Pass
<b>Ligation and Recutting (Terminal Integrity)</b> After a 10-fold over-digestion of Lambda DNA with MlyI, ~75% of the DNA fragments can be ligated with T4 DNA ligase in 16 hours at 16°C. Of these ligated fragments, ~75% can be recut with MlyI.	Pass
Non-Specific DNase Activity (16 Hour) A 50 μl reaction in rCutSmart™ Buffer containing 1 μg of Lambda DNA and a minimum of 10 units of MlyI incubated for 16 hours at 37⁰C results in a DNA pattern free of	Pass





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detectable nuclease degradation as determined by agarose gel electrophoresis.	
Protein Purity Assay (SDS-PAGE) Mlyl is >95% pure as determined by SDS PAGE analysis using Coomassie Blue detection.	Pass
<b>qPCR DNA Contamination (E. coli Genomic)</b> A minimum of 10 units of MlyI 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 $\leq$ 1 E. coli genome.	Pass

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

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Ana Egana Production Scientist 12 Jun 2024

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Michael Tonello Packaging Quality Control Inspector 12 Jun 2024

