

New England Biolabs Product Specification

Product Name:	<i>Tth1111</i>
Catalog #:	<i>R0185S</i>
Concentration:	<i>10,000 units/ml</i>
Unit Definition:	<i>One unit is defined as the amount of enzyme required to digest 1 µg of pBC4 DNA in rCutSmart Buffer in 1 hour at 65°C in a total reaction volume of 50 µl.</i>
Shelf Life:	<i>24 months</i>
Storage Temp:	<i>-20°C</i>
Storage Conditions:	<i>500 mM NaCl, 10 mM Tris-HCl, 1 mM DTT, 0.1 mM EDTA, 50 % Glycerol, 200 µg/ml rAlbumin, (pH 7.4 @ 25°C)</i>
Specification Version:	<i>PS-R0185S v2.0</i>
Effective Date:	<i>15 Apr 2024</i>

Assay Name/Specification (minimum release criteria)

Exonuclease Activity (Radioactivity Release) - A 50 µl reaction in rCutSmart™ Buffer containing 1 µg of a mixture of single and double-stranded [³H] *E. coli* DNA and a minimum of 50 units of Tth1111 incubated for 4 hours at 65°C releases <0.1% of the total radioactivity.

Functional Testing (15 minute Digest) - A 50 µl reaction in rCutSmart™ Buffer containing 1 µg of pBC4 DNA and 1 µl of Tth1111 incubated for 15 minutes at 65°C results in complete digestion as determined by agarose gel electrophoresis.

Non-Specific DNase Activity (16 hour) - A 50 µl reaction in rCutSmart™ Buffer containing 1 µg of pBC4 DNA and a minimum of 10 units of Tth1111 incubated for 16 hours at 65°C results in a DNA pattern free of detectable nuclease degradation as determined by agarose gel electrophoresis. NOTE: although no nuclease degradation is detected under these conditions, extended incubations and/or high concentrations of this enzyme may result in star activity. See the product FAQ for recommended reaction conditions for this enzyme.

qPCR DNA Contamination (*E. coli* Genomic) - A minimum of 10 units of Tth1111 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.

Ligation and Recutting (Terminal Integrity) - After a 5-fold over-digestion of pBC4 DNA with Tth1111, ~25% of the DNA fragments can be ligated with T4 DNA ligase in 16 hours at 16°C. Of these ligated fragments, >95% can be recut with Tth1111.

Protein Purity Assay (SDS-PAGE) - Tth1111 is >95% pure as determined by SDS PAGE analysis using Coomassie Blue detection.





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Date 15 Apr 2024

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Quality Approver

