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New England Biolabs Certificate of Analysis

Product Name:	<i>Tth1111</i>
Catalog #:	R0185S/L
Concentration:	5,000 units/ml
Unit Definition:	One unit is defined as the amount of enzyme required to digest 1 μ g of pBC4 DNA in 1 hour at 65 °C in a total reaction volume of 50 μ l.
<i>Lot</i> #:	0351712
Assay Date:	12/2017
Expiration Date:	12/2019
Storage Temp:	-20°C
Storage Conditions:	500 mM NaCl, 10 mM Tris-HCl (pH 7.4), 1 mM DTT, 0.1 mM EDTA, 50% Glycerol, 200 μg/ml BSA
Specification Version:	PS-R0185S/L v1.0
Effective Date:	30 Apr 2014

Assay Name/Specification (minimum release criteria)	Lot #0351712
Exonuclease Activity (Radioactivity Release) - A 50 μ l reaction in CutSmart TM Buffer containing 1 μ g of a mixture of single and double-stranded [³ H] <i>E. coli</i> DNA and a minimum of 50 units of Tth1111 incubated for 4 hours at 65°C releases <0.1% of the total radioactivity.	Pass
Ligation and Recutting (Terminal Integrity) - After a 5-fold over-digestion of pBC4 DNA with Tth111I, \sim 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 Tth111I.	Pass
Non-Specific DNase Activity (16 hour) - A 50 μ l reaction in CutSmart TM Buffer containing 1 μ g of pBC4 DNA and a minimum of 5 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.	Pass
Protein Purity Assay (SDS-PAGE) - Tth1111 is >95% pure as determined by SDS PAGE analysis using Coomassie Blue detection.	Pass

* The BSA in this product has been granted an EDQM "Certificate of Suitability" from the European Directorate for the Quality of Medicines (# R1-CEP-2003-204-Rev00) and has been granted a USDA Certificate for Export of Bovine Blood Plasma/Serum for Manufacture into Pharmaceutical Products.

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Authorized by Derek Robinson 30 Apr 2014



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Inspected by Anthony Francis 18 Dec 2017