

*be* INSPIRED *drive* DISCOVERY *stay* GENUINE

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:	KpnI-HF®
Catalog Number:	R3142M
Concentration:	100,000 U/ml
Unit Definition:	One unit is defined as the amount of enzyme required to digest 1 µg of pXba DNA in rCutSmart™ Buffer in 1 hour at 37°C in a total reaction volume of 50 µl.
Packaging Lot Number:	10134135
Expiration Date:	01/2024
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-R3142M v2.0

KpnI-HF® Component List				
NEB Part Number	Component Description	Lot Number	Individual QC Result	
R3142MVIAL	KpnI-HF®	10137590	Pass	
B7024AVIAL	Gel Loading Dye, Purple (6X)	10130600	Pass	
B6004SVIAL	rCutSmart™ Buffer	10131975	Pass	

Assay Name/Specification	Lot # 10134135
<b>Blue-White Screening (Terminal Integrity)</b> A sample of Litmus28i vector linearized with a 10-fold excess of KpnI-HF®, religated and transformed into an E. coli strain expressing the LacZ beta fragment gene results in <1% white colonies.	Pass
Endonuclease Activity (Nicking) A 50 µl reaction in rCutSmart <sup>™</sup> Buffer containing 1 µg of supercoiled PhiX174 DNA and a minimum of 60 units of KpnI-HF® incubated for 4 hours at 37°C results in <20% conversion to the nicked form as determined by agarose gel electrophoresis.	Pass
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 200 units of KpnI-HF® 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 pXba DNA and 1 μl of	Pass





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Assay Name/Specification	Lot # 10134135
KpnI-HF® incubated for 15 minutes at 37°C results in complete digestion as determined by agarose gel electrophoresis.	
<b>Ligation and Recutting (Terminal Integrity)</b> After a 50-fold over-digestion of pXba DNA with KpnI-HF®, >95% 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 KpnI-HF®.	Pass
<b>qPCR DNA Contamination (E. coli Genomic)</b> A minimum of 20 units of KpnI-HF® 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
<b>Protein Purity Assay (SDS-PAGE)</b> KpnI-HF® is >95% pure as determined by SDS PAGE analysis using Coomassie Blue detection.	Pass
Non-Specific DNase Activity (16 Hour) A 50 µl reaction in rCutSmart <sup>™</sup> Buffer containing 1 µg of pXba DNA and a minimum of 100 units of KpnI-HF® incubated for 16 hours at 37°C results in a DNA pattern free of detectable nuclease degradation as determined by agarose gel electrophoresis.	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.

Penghua Zhang Production Scientist 04 Feb 2022

Michae

Michael Tonello Packaging Quality Control Inspector 04 Feb 2022

