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New England Biolabs Product Specification

Product Name: AscI

Catalog #: R0558S/L
Concentration: 10,000 units/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.

Shelf Life: 24 months
Storage Temp: -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-R0558S/L v2.0
Effective Date: 08 Sep 2022

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Assay Name/Specification (minimum release criteria)

Blue-White Screening (Terminal Integrity) - A sample of pNEB193 vector linearized with a 10-fold excess of AscI, religated and transformed into an *E. coli* strain expressing the LacZ beta fragment gene results in <1% white colonies.

Endonuclease Activity (Nicking) - A 50 μ l reaction in rCutSmartTM Buffer containing 1 μ g of supercoiled PhiX174 DNA and a minimum of 30 units of AscI incubated for 4 hours at 37°C results in <10% conversion to the nicked form as determined by agarose gel electrophoresis.

Exonuclease Activity (Radioactivity Release) - A 50 μl reaction in rCutSmartTM Buffer containing 1 μg of a mixture of single and double-stranded [³H] *E. coli* DNA and a minimum of 100 units of AscI incubated for 4 hours at 37°C releases <0.1% of the total radioactivity.

Functional Testing (15 minute Digest) - A 50 μ l reaction in rCutSmartTM Buffer containing 1 μ g of Lambda DNA and 1 μ l of AscI incubated for 15 minutes at 37°C results in complete digestion as determined by agarose gel electrophoresis.

Ligation and Recutting (Terminal Integrity) - After a 10-fold over-digestion of Lambda DNA with AscI, >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 AscI.

Non-Specific DNase Activity (16 Hour) - A 50 μ l reaction in rCutSmartTM Buffer containing 1 μ g of Lambda DNA and a minimum of 30 units of AscI incubated for 16 hours at 37°C results in a DNA pattern free of detectable nuclease degradation as determined by agarose gel electrophoresis.

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









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Date 08 Sep 2022

Derek Robinson Quality Approver





