pBR322 DNA–BstNI Digest

1-800-632-7799
info@neb.com
www.neb.com

N3031S 060120414041

N3031S

50 gel lanes (50 µg) **Lot: 0601204** **Exp: 4/14**

1,000 µg/ml **Store at –20°C**

1.5 ml Gel Loading Dye, Blue (6X) **Store at 25°C**

Description: The BstNI Digest of pBR322 DNA yields 6 fragments suitable for use as molecular weight standards for agarose gel electrophoresis (1,2).

Supplied in: 10 mM Tris-HCl (pH 8.0), 1 mM EDTA.

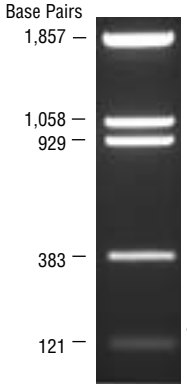
Reagents supplied:
6X Gel Loading Dye, Blue

1X Gel Loading Dye, Blue:
2.5% Ficoll-400
11 mM EDTA
3.3 mM Tris-HCl (pH 8.0@25°C)
0.017% SDS
0.015% bromophenol blue

Preparation: Prepared from *E. coli* ER2420 (dam$^+$ dcm$^+$ EcoKM$^-$) by a standard plasmid purification procedure, the double-stranded DNA is digested to completion with BstNI, phenol extracted and equilibrated to 10 mM Tris-HCl (pH 8.0) and 1 mM EDTA.

Usage Recommendation: The approximate mass of DNA in each of the bands in our pBR322 DNA-BstNI Digest is as follows (assuming a 1.0 µg loading):

Fragment	Base Pairs	DNA Mass
1	1,857	426 ng
2	1,058	243 ng
3	929	213 ng
4	383	88 ng
5	121	28 ng
6	13	3 ng



pBR322 DNA-BstNI Digest visualized by ethidium bromide staining. 1.4% agarose gel.

Note: For long term storage, store at –20°C. If samples need to be diluted, use TE or other buffer of minimal ionic strength. DNA may denature if diluted in dH_2O.

Suggested protocol for loading a sample:

The following protocol is recommended for a 5 mm wide lane.

1. Prepare loading mixture:

Distilled water	4 µl
6X Blue Loading Dye	1 µl
DNA Ladder	1 µl
Total volume	6 µl

2. Mix gently
3. Load onto the agarose gel

Note: The components of the mixture should be scaled up or down, depending on the width of the agarose gel.

References:

1. Sutcliffe, J. G. (1978) Cold Spring Harbor *Symp. Quant. Bio.* 43:77–90.
2. Peden, K. W. C. (1983) *Gene* 22, 277–280.
3. Forster, A.C. et al. (1985) *Nucl. Acids Res.* 13, 745-761

CERTIFICATE OF ANALYSIS

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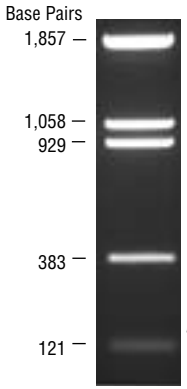

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