

pBR322 DNA– MspI Digest



1-800-632-7799
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N3032S

50 gel lanes (50 µg) Lot: 1531204 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 MspI digest of pBR322 DNA yields 26 fragments.

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

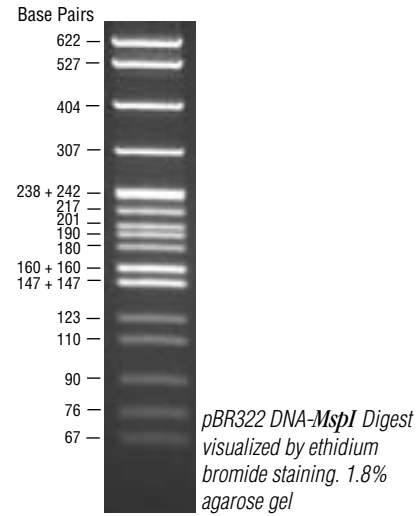
Reagents supplied:
6X Gel Loading Dye, Blue

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 MspI, 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- MspI Digest is as follows (assuming a 1.0 µg loading):



Fragment	Base Pairs	DNA Mass
1	622	143 ng
2	527	121 ng
3	404	93 ng
4	307	70 ng
5	242	55 ng
6	238	55 ng
7	217	50 ng
8	201	46 ng
9	190	44 ng
10	180	41 ng
11, 12	160	74 ng
13, 14	147	68 ng
15	123	28 ng
16	110	25 ng
17	90	21 ng
18	76	17 ng
19	67	15 ng
20, 21	34	16 ng
22, 23	26	12 ng
24	15	3 ng
25, 26	9	4 ng

(see other side)

CERTIFICATE OF ANALYSIS

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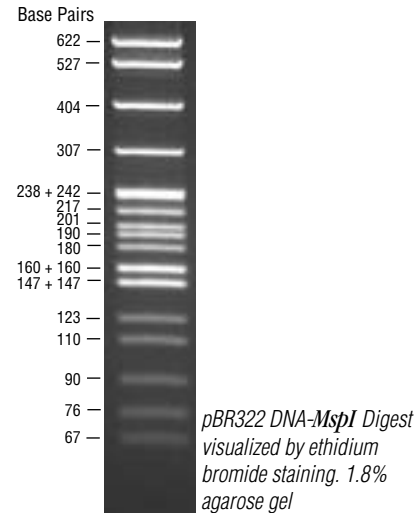
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CERTIFICATE OF ANALYSIS

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	<u>6 μl</u>

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.

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