

100bp DNA Ladder

Code No. DNA-035
Lot No. *****
Storage Conditions -20°C
Shipping Conditions 4°C
Size 0.5ml
Contents 100bp DNA Ladder × 1
6 × Loading Dye × 1

Certificate of Analysis

Product : A mixture of ten double-stranded DNA fragments ranging from 100bp to 1,000bp in exactly 100bp increments, and 1,500bp of double-stranded DNA fragments. The 500bp band has increased intensity relative to the other bands. The 100bp DNA Ladder was dissolved in 10mM Tris-HCl (pH8.0), 5mM NaCl, and 10mM EDTA.

Concentration : 0.03 µg / µl

Volume : 500 µl / tube

| Fragment sizes | fragment No. | Number of Base Pairs(bps) | Concentration (ng/5µl) |
|----------------|--------------|---------------------------|------------------------|
| | 1 | 1,500 | 10 |
| | 2 | 1,000 | 10 |
| | 3 | 900 | 10 |
| | 4 | 800 | 10 |
| | 5 | 700 | 10 |
| | 6 | 600 | 10 |
| | 7 | 500 | 30 |
| | 8 | 400 | 10 |
| | 9 | 300 | 10 |
| | 10 | 200 | 15 |
| | 11 | 100 | 25 |

Contaminant assay (Nuclease assay) : After overnight incubation of 100bp DNA Ladder at 37°C, no visible degradation of banding pattern is observed on agarose gel electrophoresis analysis.

Recommended handling : 1. Centrifuge tubes before opening to improve recovery of content.
2. Add 1/5 volume of 6 × Loading Dye to 100bp DNA Ladder, and mix well.
3. Load 6 µl of the 100bp DNA Ladder per lane on agarose gels.

Note : Use 1.5~2 % agarose gels for good banding. Using 1% agarose gels may make lower bands faint.