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# **ProductInformation**

### 123 BP DNA LADDER

Product Number **D 5042** Storage Temperature 0 to -20 °C

**Product Summary** 

Storage buffer: 10 mM Tris-HCl, pH 7.5, 50 mM NaCl,

and 0.1 mM EDTA

Concentration: 0.8 to 1.2 μg/μl

See label for lot specific concentration.

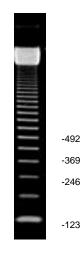
## **Suitability Assay and Results**

123 bp ladder was prepared for electrophoresis as follows:

0.5-2.0 μg 123 BP Ladder
2.5 μl Gel Loading Solution (Product No. G 2526)
(0.05% w/v bromophenol blue, 40% w/v sucrose, 0.1 M EDTA, pH 8.0)
Q.S. to 10 μl with 1X TE Buffer (10 mM TrisHCl, pH approx. 8.0, 1 mM EDTA)

The above solution was loaded on 1.5% agarose (Product No. A 9539) submarine type minigel. The samples were run in 1X TAE (40 mM Tris-acetate, 1 mM EDTA, pH approx. 8.3) with appropriate DNA fragment size standards at 80 volts for 2 hours. After staining 15-20 minutes in 10  $\mu$ g/ml ethidium bromide, the bands were clearly resolved and the pattern was consistent with the indicated fragment sizes.

## Fragment Sizes: base pairs (bp)



123 to 4182 bp in crements of 123 bp

### Notes:

- 1. Background can be reduced by destaining 30-45 minutes in 1X TAE.
- 2. The bromophenol blue tracking dye will migrate just ahead of the 492 bp fragment.
- 3. The 123 bp band will stain noticeably brighter than the other bands.

02/01