

Catalogue	Pack Size
DL-100BP-200	200 μ L (40 lanes)
DL-100BP-500	500 μ L (100 lanes)

For research and educational use only.

Description

DNA Ladders, also known molecular-weight size markers, contain a set of predetermined DNA fragment sizes. These markers are a set of standards used to identify the approximate size of a DNA fragment run on gel electrophoresis.

The [100 bp DNA Ladder](#) is ideal for routine use during the electrophoresis of PCR-amplified DNA. It is supplied pre-mixed with loading dye: this allows easy pipetting into electrophoresis gel wells, and acts as a visual aid to track the extent of DNA migration during gel electrophoresis. Its formulation also allows room temperature usage and short-term storage.

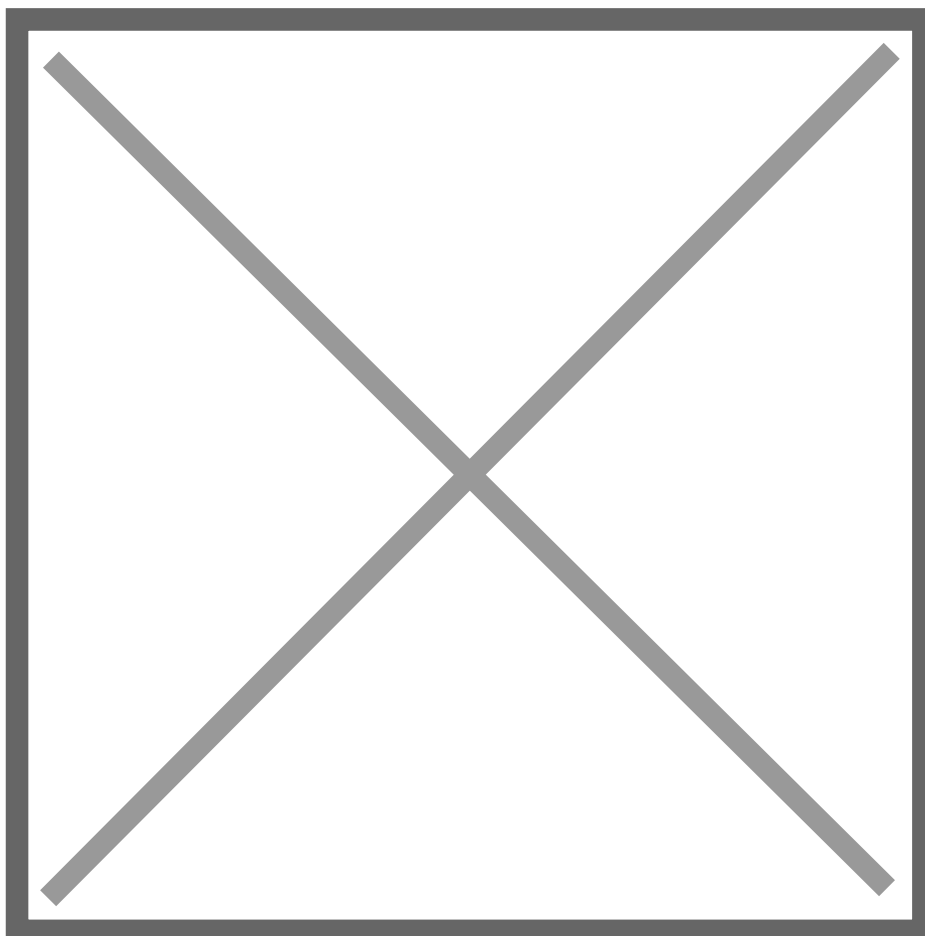


Figure 1: Size and concentration of DNA sizing fragments present in 100 bp DNA Ladder.

Application Recommendations

Gel electrophoresis

We recommend using 100 bp DNA Ladder to size DNA fragments of 100 – 3000 bp in length, such as DNA amplicons produced by many PCR applications.

Reagent Composition

100 bp Ladder (13 DNA fragments at 0.1 µg/µL (size range: 100 – 3000bp), 10 mM EDTA, 10% glycerol, 0.015% bromophenol blue, 0.17% SDS).

Storage & Stability

Store dry and at room temperature.

Long-term storage at -20°C.

Shipping conditions

Shipped at room temperature.

Safety warnings and precautions

This product and its components are not considered hazardous in their given concentrations. However, as with all scientific reagents this product should be handled and stored with care as standard practice. Wear gloves. Care should be taken to avoid contact with skin or eyes. In case of contact with skin or eyes, wash immediately with water.

Quick Start Protocol

1. Using a new pipette tip each time, transfer 5 µL of [100 bp DNA Ladder](#) in the electrophoresis gel well to the left of that assigned for the first sample, and the well to the right of the last sample. For example, the first and last wells if all wells are being used. No additional loading dye is needed because it is already included in the DNA ladder solution.
2. Load DNA samples as normal.
3. Run gel electrophoresis according to the appropriate protocol.