

DNA Marker/Ladder



MD101/102/103/104/105

Version 8.1

Vazyme biotech co., Ltd.

Introduction

This product consists of double-stranded DNA fragments of specific molecular weight and is mixed with loading buffer containing a blue dye, which is suitable for DNA molecular weight standards in agarose gel electrophoresis. All fragments in the DNA Marker/Ladder are obtained by enzymatic digestion and purification. The bands of Marker/Ladder are more clear and dense during electrophoresis; the mass ratio between the bands is more accurate and true. The 750 bp fragment in the DL2000 Plus DNA Marker, the 1,000 bp fragment in the DL5000 DNA Marker, the 500 bp fragment in the 100 bp DNA Ladder, and the 5 kb fragment in the 1 kb DNA Ladder have a DNA concentration of 100 ng/5 μ l, which are bright bands; all remaining bands have a DNA concentration of 50 ng/5 μ l.

Components

Components	250 μ l (50 rxn, 5 μ l/rxn)	500 μ l (100 rxn, 5 μ l/rxn)
DL2000 Plus DNA Marker	MD101-01	MD101-02
DL5000 DNA Marker	MD102-01	MD102-02
DL15000 DNA Marker	MD103-01	MD103-02
100 bp DNA Ladder	MD104-01	MD104-02
1 kb DNA Ladder	MD105-01	MD105-02

*Stored in 10 mM Tris-HCl (pH 8.0); 10 mM EDTA; 6% Glycerol; 0.01% Bromophenol Blue; 0.005% Xylene Cyanol.

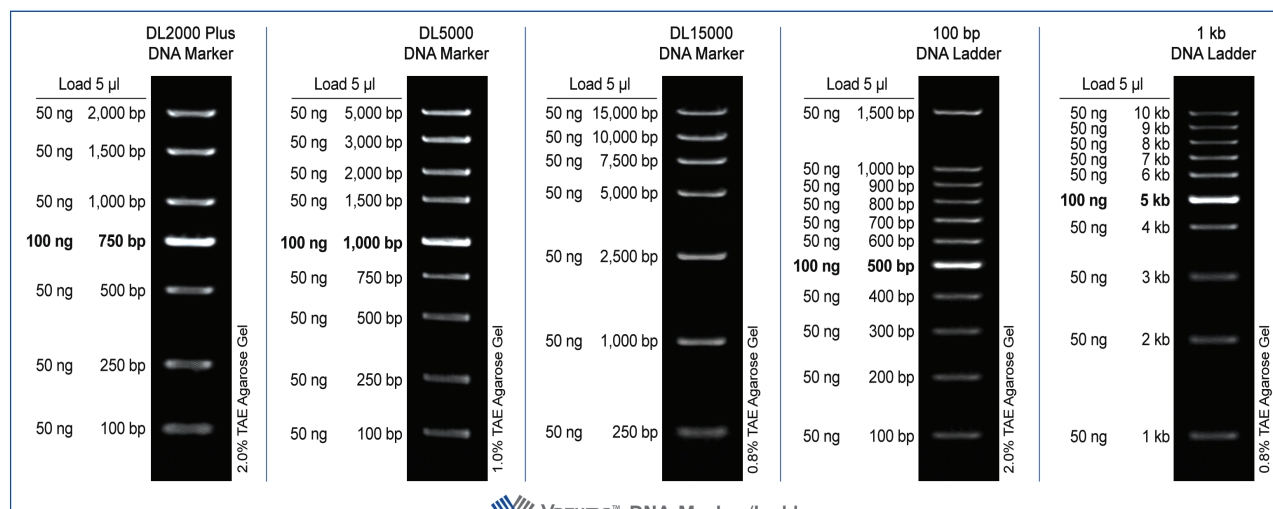
Storage

Store at -20°C; store at 4°C after thawing to avoid repeated freezing and thawing.

Experiment Process

1. This product is a ready-to-use product. You can directly take 5 μ l of the product into the wells of the agarose gel for electrophoresis (if the spotting holes are wide, you can increase the loading amount).
2. It is recommended that the electrophoresis conditions be 1 \times TAE buffer, 0.8 - 2.0% agarose gel, and the voltage between positive and negative electrodes is 4 - 10 V/cm.
3. Two electrophoretic indicators of Xylene Cyanol and Bromophenol Blue have been added to this product. If a 1% agarose gel is used, the xylene cyanide band is located at approximately 2 kb and the bromophenol blue band is located at approximately 400 bp.
4. The electrophoresis bands are observed under UV light by staining with nucleic acid dyes.

The figure below shows the results of Vazyme DNA Marker/Ladder electrophoresis.



Vazyme™ DNA Marker/Ladder



Vazyme Biotech Co., Ltd.
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Order: global@vazyme.com

Support: support@vazyme.com

For research use only, not for use in diagnostic procedures.

Notes

1. Please thaw and mix thoroughly before use.
2. The quality of the electrophoresis image is related to the agarose gel and the electrophoresis buffer. Please use high quality agarose, freshly prepared agarose gel, and replace the running buffer in time to avoid affecting the electrophoresis results.
3. The concentration of the agarose gel is critical for the separation of DNA fragments. Higher concentrations of agarose gel have better separation performance for short fragments of DNA, while lower concentrations of agarose gel facilitate separation of long fragment DNA. According to the actual situation, a suitable concentration of agarose gel can be selected for electrophoresis.
4. After the DNA fragments of equal mass are stained by electrophoresis or nucleic acid dye, the fragments with smaller molecular weight are lightly colored and the bands are thick; the fragments with larger molecular weight are darker and the bands are fine. It is a normal phenomenon.
5. If EB is used as the nucleic acid dye, it should be noted that the electrical properties of EB are opposite to those of DNA, and EB and DNA migrate in the opposite direction during electrophoresis. If EB is added in the preparation of agarose gel, when the electrophoresis is carried out for a long time, the smaller molecular weight fragments in the DNA Marker/Ladder may be lightly colored and the bands are blurred, which is a normal phenomenon.



ISO 9001: 2015