Tel: 609-228-6898

Email: tech@MedChemExpress.com



500 bp DNA Marker

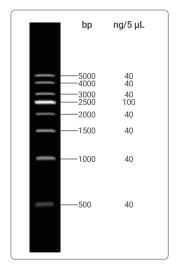
1 Contents

Components	HY-K0804-250 μL	HY-K0804-500 μL	HY-K0804-1 mL
500 bp DNA Marker	250 μL	250 μL × 2	250 μL × 4

2 Introduction

The 500 bp DNA Marker is provided in a solution of $1 \times DNA$ Loading Buffer, which can be directly used for nucleic acid electrophoresis analysis. The Marker contains 14 double-stranded DNA fragments ranging from 500 bp to 5000 bp. 5 μ L of this product contains about 100 ng for the 2500 bp band, and about 40 ng for the other bands.

3 Electrophoresis illustration



1.4% Agarose 1x TAE Buffer $5 \mu L/lane 7 V/cm, 45 min$

4 Protocol

- 1. Add 5 μ L of 500 bp DNA Marker to sample well of the agarose gel and perform electrophoresis.
- 2. After electrophoresis, stain with Nucleic Acid Gel Stain and detect the electrophoresis results.

Note: a) 1.0-1.5% agarose gel at 5-10 V/cm is recommended.

- b) Adjust the loading volume of DNA Marker for different loading well format.
- c) Pre-dyeing or post-dyeing is suitable when using the Nucleic Acid Gel Stain.

Storage

-20°C, 2 years.

Avoid repetitive freeze-thaw cycles.

Precautions

- 1. For short-term use, DNA Marker may be stored at 2-8°C.
- 2. Replace the electrophoresis buffer in time and use fresh agarose gels to achieve better electrophoretic results.
- 3. When the concentration of agarose gel is or too high or too low, the bands will not be easy to separate.
- 4. This product is for R&D use only, not for drug, household, or other uses.
- 5. Please consult the Material Safety Data Sheet for information regarding hazards and safe handling practices

Recommended products used for Nucleic Acid Gel Electrophoresis

Cat. No	Name	Application	
HY-K1031	Agarose		
HY-K1029	Agarose With TAE Powder (1%)		
HY-K1016	TBE Powder (1 L of 1×)		
HY-K1015	TAE Powder (1 L of 1×)	- Electrophoresis buffer	
HY-K1017	Rapid Running Buffer Powder (1 L of 1x)		
HY-K1004	SYBR Green I Nucleic Acid Gel Stain	– Nucleic Acid Gel Stain	
HY-K1007	Red Nucleic Acid Gel Stain (10,000×)		

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