For life science research only. Not for use in diagnostic procedures.



DNA Molecular Weight Marker IV

Version: 19
Content Version: July 2021

Fragment sizes: 0.07 to 19.3 kbp

λDNA and pSPTBM 20 DNA × Sty I and Sau I digested

Cat. No. 11 418 009 001 50 μg

200 µl

50 gel lanes

Store the product at -15 to -25°C.

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1. General Information

1.1. Contents

Vial / bottle	Label	Function / description	Content
1	DNA Molecular Weight Marker IV	 Ready-to-use solution in 10 mM Tris-HCl, 1 mM EDTA, pH 8.0, (250 µg/ml). 50 µg corresponds to 1 A₂₆₀ unit. 	1 Vial, 50 µg (200 µl)

1.2. Storage and Stability

Storage Conditions (Product)

When stored at -15 to -25°C, the product is stable through the expiry date printed on the label.

Vial / bottle	Label	Storage
1	DNA Molecular Weight Marker IV	Store at -15 to -25 °C.
		After thawing, store at +2 to +8°C.
		Avoid repeated freezing and thawing.

1.3. Application

Use DNA Molecular Weight Marker IV as a size standard for DNA in agarose gels.

2. How to Use this Product

2.1. Before you Begin

General Considerations

Size distribution

Fragment mixture prepared by cleavage of equimolar amounts of λDNA and pSPTBM 20 DNA with restriction endonucleases Sty I and Sau I. The mixture contains 14 DNA fragments with the following base pair lengths (1 base pair = 660 daltons).

Fragment lengths are derived from computer analysis of the λDNA and pSPTBM 20 sequences.

bp												
19,329 7,743	5,526	4,254	3,140	2,690	2,322	1,882	1,489	1,150	925	697	421	74

Improved visualization of the bands

The 19,329 and 4,254 bp fragments contain the cos-ends of lambda. These bands are visible after heating the marker at +65°C for 10 minutes, and quickly chilling on ice.

⚠ Fragments containing the 12 base cos-sites of lambda may anneal upon storage. This leads to a gel pattern where one band is of lower intensity than expected (or absent completely) and a larger fragment has an increased intensity. Denaturation of the cos-sites can be performed immediately before loading the gel, by heating at +65°C for 10 minutes and quick-chilling on ice.

3. Results

Typical analysis

The DNA fragment mixture shows the typical pattern of 13 bands in agarose gel electrophoresis, see Figure 1.

- After gel electrophoresis of 1 µg of the fragment mixture in a 1% Agarose MP* gel, 13 bands are visible.
- The smallest band is only visible in overloaded gels.

IV

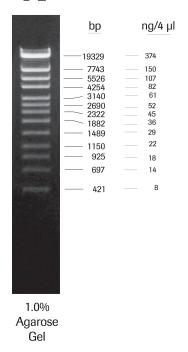


Fig. 1: Separation of 1 µg DNA Molecular Weight Marker IV on a 1% Agarose MP gel, stained with ethidium bromide.

4. Supplementary Information

4.1. Conventions

To make information consistent and easier to read, the following text conventions and symbols are used in this document to highlight important information:

Text convention and symbols					
1 Information Note: Additional information about the current topic or procedure.					
⚠ Important Note: Information critical to the success of the current procedure or use of the product.					
1 2 3 etc.	Stages in a process that usually occur in the order listed.				
1 2 3 etc. Steps in a procedure that must be performed in the order listed.					
* (Asterisk)	The Asterisk denotes a product available from Roche Diagnostics.				

4.2. Changes to previous version

Layout changes. Editorial changes.

4.3. Trademarks

All product names and trademarks are the property of their respective owners.

4.4. License Disclaimer

For patent license limitations for individual products please refer to: **List of biochemical reagent products**.

4.5. Regulatory Disclaimer

For life science research only. Not for use in diagnostic procedures.

4.6. Safety Data Sheet

Please follow the instructions in the Safety Data Sheet (SDS).

4.7. Contact and Support

To ask questions, solve problems, suggest enhancements or report new applications, please visit our **Online Technical Support Site**.

To call, write, fax, or email us, visit **sigma-aldrich.com**, and select your home country. Country-specific contact information will be displayed.

