

Lonza Rockland, Inc. www.lonza.com biotechserv@lonza.com Tech Service: 800-521-0390 Customer Service: 800-638-8174 Document # 18142-0807-04 Rockland, ME 04841 USA

# **DNA Marker 50-2,500 bp**

#### Introduction

DNA Markers range in size from 50bp to 2,500 bp for rapid size estimation of PCR $^{\dagger}$  products and restriction fragments. Loading 5  $\mu$ l per lane yields approximately 50 ng of DNA per band. DNA can be visualized by ethidium bromide staining or kinased with radiolabeled  $^{32}$ P for detection by autoradiography.

#### Contents DNA Marker

250  $\mu$ l-50 applications Store at 4°C

## Loading Buffer 6X (250 µl)

Contains: bromophenol blue Store at 4°C or 25°C

## Triple Dye Loading Buffer 6X (1.1 ml)

Contains: Orange G, bromophenol blue, xylene cyanol Store at  $4^{\circ}\text{C}$ 

#### **Standard Procedure**

- 1. Mix 5 μl of DNA marker and 1 μl of 6X loading buffer.
- 2. Mix 5 parts of your sample to 1 part of 6X loading buffer.
- 3. Load DNA markers and samples onto an agarose gel.
- 4. Electrophorese, stain, and photograph following your standard protocol.
- Estimate the size of the sample DNA by reading its relative position to the closest marker.

**NOTE:** There is a higher concentration of dye material in the Triple Dye Loading Buffer. Therefore, we recommend it for use in large (40 cm) gels, run for extended times, (18 hours or more).

### Procedure for 5' End Radiolabeling

**NOTE:** The marker can be labeled directly or for more efficient labeling, ethanol precipitate first.

#### **Ethanol Precipitation**

- 1. Remove a 100 μl aliquot of the DNA marker.
- 2. Add 10 μl of 3 M potassium acetate, pH 7.4.
- 3. Add 300 μl of <u>absolute</u> ethanol.
- 4. Incubate at -70°C for 30 minutes.
- 5. Microcentrifuge at 4°C for 10 minutes.
- 6. Redissolve the pallet in 100 μl of distilled water.
- Quantitate by reading the absorption at 260 nm and label with [<sup>Υ32</sup>P]-ATP using T4 polynucleotide kinase and a standard protocol. See Sambrook, et al., 5.68 (1989) or Ausubel, et al. (1987).

#### **Product Safety:**

For details regarding product safety, see Material Safety Data Sheet (MSDS); call (800) 638-8174 for extra copies of the MSDS. Emergency after hours, call collect (303) 595-9048.

#### Warranty:

Because of the numerous factors affecting results, Lonza DNA markers are sold with the understanding that purchasers will make their own tests to determine the suitability of these markers for their particular purposes. The use suggested by Lonza is presented only to assist our customers in exploring possible applications for this product. All information and data presented are believed to be accurate and reliable but are presented without the assumption of any liability by Lonza.

#### References

- Ausubel, F.M., Brent, R., Kingston, R.E., Moore, D.D., Seidman, J.G., Smith, J.A., and Struhl, K.
   Current Protocols in Molecular Biology, John Wiley & Sons, New York 1987.
- Sambrook, J., Fritsch, E.F., and Maniatis, T.
   Molecular Cloning, A Laboratory Manual,
   Second Edition, Cold Spring Harbor:
   Cold Spring Harbor Laboratories 1989

# **Ordering Information**

Catalog No. Description Size

50631 DNA Marker 50-2,500 bp 50 Applications



GelStar® Nucleic Acid Gel Stain Latitude® HT Precast Agarose Gels Latitude® Precast Agarose Midigels Reliant® Gel System SYBR® Green Gel Stains SeaKem® LE Agarose SeaPlaque® GTG® Agarose SeaPlaque® Agarose MDE® Gel Solution

For more information contact Technical Service at (800) 521-0390 or visit our website at www.Lonza.com.



Manufactured for Lonza Rockland, Inc.

For Research Use Only.

Not for Use in Diagnostic Procedures.

<sup>†</sup>The PCR process may be covered by one or more third-party patents.

GelStar and SeaKem are trademarks of FMC Corporation. All other trademarks herein are marks of the Lonza Group or its affiliates. GelStar is covered by U.S. patents 5,436,134. Other U.S. and foreign patent pending. SYBR is a trademark of Molecular Probes.

© 2007 Lonza Rockland, Inc. All rights reserved.