

DL1500 Ladder

产品组成

产品名称	产品规格	Cat. No.
DL1500 Ladder	250 μ l	MD1019
DL1500 Ladder	250 μ l $\times$ 5	MD1119

产品储存与有效期

产品可在常温（0-30℃）储存三年。如果长期不用，为防止水分蒸发请于 - 20℃ 储存。

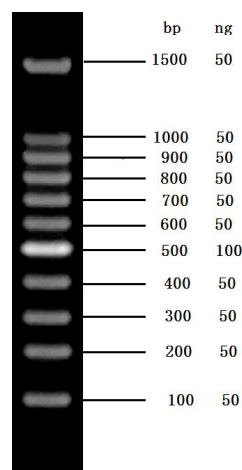
技术支持

杭州新景生物试剂开发有限公司研发部：e-mail: technical@simgen.cn, 电话：400-0099-857。

产品介绍

DL1500 Ladder 由 11 种长度在 100 bp 至 1,500 bp 的 DNA 片段组成，溶解于 1 $\times$ Loading Buffer 中，使用时可取 5-10 μ l 直接电泳，使用非常方便。

特别添加的红色和黄色两种电泳指示染料，不会削弱 DNA 在紫外线下的显色效果，较常用的电泳指示染料（溴酚蓝、二甲苯青等）具有更佳的使用效果。



注意事项

1. 电泳时的加样孔宽度小于 5 mm 时，每次取 5 μ l DNA Ladder 电泳便可得到清晰条带。如果加样孔增宽，须适当增加 DNA Ladder 的加样量。
2. 如果预计需要鉴定的 DNA 片段在 100~500 bp 之间，推荐使用 2.5%~3% 的 Agarose 凝胶电泳；如果预计需要鉴定的 DNA 片段在 500~1500 bp 之间，推荐使用 1.5%~2% 的 Agarose 凝胶电泳。
3. 对 DNA 电泳而言，Agarose 的纯度对 DNA 条带的清晰度影响很大。因此，电泳时应尽量选用质量好的 Agarose。
4. 进行 Agarose 电泳时，Agarose 浓度越大，对短片段 DNA 分离性能越好；反之，Agarose 浓度越小，越有利于长片段 DNA 的分离。
5. 使用 GelRed 作为电泳染料时，由于嵌入 DNA 中的 GelRed 分子较溴化乙锭更大，可能出现条带弥散或分离不理想的情况，请尝试减少 GelRed 用量或使用泡染法。

DL1500 Ladder

PRODUCT FORMATION

Components	Specification	Cat. No.
DL1500 Ladder	250 µl	MD1019
DL1500 Ladder	250 µl×5	MD1119

STORAGE

The product can be stored at normal temperature (0-30 °C) for up to three years.If the product is not used for a long period of time, please store at -20°C to prevent the evaporation of water.

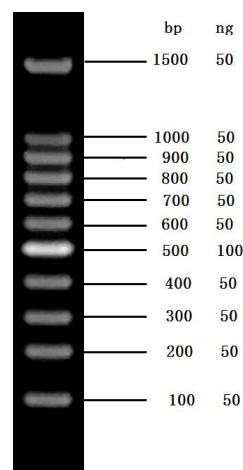
TECHNICAL SUPPORT

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INTRODUCTION

DL1500 Ladder is composed of 11 individual DNA fragments ranging from 100 bp to 1500 bp. DL1500 Ladder contains 1×Loading Buffer, users can apply 5 - 10 µl in agarose gel electrophoresis directly. The red and yellow tracking dye in DL1500 Ladder will not weakened the DNA bands under UV light, better than bromophenol blue and xylene cyanol FF.



PRECAUTION

1. Clear bands can be obtained by applying 5 µl DNA Ladder when the lane width is less than 5 mm. If the lane is wider, loading volume of DNA Ladder should be increased appropriately.
2. If the DNA fragment is expected to be identified between 100~500 bp, gel concentration of 2.5% - 3% is recommended. If the DNA fragment is expected to be identified between 500~1500 bp, gel concentration of 1.5% - 2% is recommended.
3. For DNA electrophoresis, agarose purity is of great significance to DNA band definition. Therefore, agarose with good quality should be used.
4. During agarose electrophoresis, the concentration of agarose is closely associated with the separation of DNA fragments. High agarose concentration is ideal for the separation of the short DNA fragments. While low agarose concentration is ideal to separate the long DNA fragments.
5. When using GelRed as an electrophoretic dye, try reducing the amount of GelRed or using Post-Staining Protocol if the bands are diffuse or do not separate well.