Tel: (34) 91 710 00 74

C/ Valle de Tobalina, 52 - Nave 39 E-28021 Madrid (Spain) Fax: (34) 91 505 31 18 e-mail: info@biotools.eu www.biotools.eu

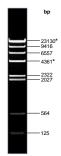
Lambda DNA/HindIII Marker

(Ref. 31.011)

Concentration: 0.5 mg/ml (50 µg) Store at -20°C

Description

The Lambda DNA is completely digested with HindIII to yield bands ranging from 0.125 kb to 23 kb, suitable for use as molecular weight standards for agarose gels. The marker is composed to 8 purified individual DNA fragments (in base pairs): 23130*, 9416, 6557, 4361*, 2322, 2027, 564 and 125.



Lambda DNA/HindIII Marker 0.5ug/lane 1% agarose gel stained with ethidium bromide

Note: *The cohesive ends of the 12nt cos site of bacteriophage lambda from fragments of 23130 bp and 4361 bp (indicated*) may anneal and form an additional band at 27491 bp. These fragments may be separated by heating at 65 °C for 5 min and then cooling on ice for 3 min.

Storage buffer (TE buffer) 10 mM Tris-HCI (pH 7.6), 1 mM EDTA

Storage temperature

Store at -20°C. For frequent use divide in aliquots to avoid multiple freeze/thaw cycles. or store at 4°C in the presence of loading buffer.

Protocol

1- Prepare loading mixture (for the 5 mm gel lane⁺):

Lambda DNA/HindIII Marker 1 ul (0.5 ug)

5X Loading Dye
Distilled water
3 μl

2- Mix gently

3- Heat for 5 min at 65 °C and then cool on ice for 3 min*

- 4- Load onto the gel
- 5- Visualise DNA by staining with ethidium bromide or with SYBR® Green I

The Lambda DNA/HindIII Marker was not designed for precise quantification of DNA mass, but can be used for semi-quantification (see Table 1). For quantification, adjust the concentration of the sample to equalize it approximately with the amount of DNA in the nearest band of the ladder.

Table 1. Percentage and mass of individual fragments (for 0.5 μg Lambda DNA/HindIII Marker)

Fragment	Size	%	mass (ng/0.5µg)
1	23130	47.7	238.4
2	9416	19.4	97.1
3	6557	13.5	67.6
4	4361	9.0	45.0
5	2322	4.8	23.9
6	2027	4.2	20.9
7	564	1.2	5.8
8	125	0.3	1.3

Notice to users

Research and in vitro use only.

^{*}The mixture should be scaled up or down, depending on the width of the gel. Use 0.1 µg of DNA marker/mm of lane.