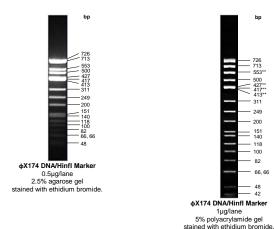
# ΦX174 DNA/Hinfl Marker

(Ref. 31.001)

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

## Description

The ΦX174 DNA is completely digested with Hinfl to yield bands ranging from 24 bp to 726 bp, suitable for use as molecular weight standards for agarose and polyacrylamide gels. The marker is composed to 21 purified individual DNA fragments (in base pairs): 726, 713, 553, 500, 427, 417, 413, 311, 249, 200, 151, 140, 118, 100, 82, 66, 66, 48, 42, 40\* and 24\*.



\*The shortest fragments (40 and 24 bp) are not visible in standard electrophoresis

### 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.

<sup>\*\*</sup>The 553, 427, 417 and 413 bp fragments migrate anomalously on polyacrilamyde gels

### Protocol

1- Prepare loading mixture (for the 5 mm gel lane<sup>+</sup>):

	Agarose Gels	Polyacrylamide Gels
<ul> <li>DNA Marker (0.5-1 μg)</li> </ul>	1 µl	2 µl
<ul> <li>5X Loading Dye</li> </ul>	1 µl	0.5 µl
<ul> <li>Distilled water</li> </ul>	3 µl	-
Final volume	5 µl	2.5 µl

#### 2- Mix gently

#### 3- Do not heat

4- Load onto the gel

5- Visualise DNA by staining with ethidium bromide or with SYBR® Green I

The \$\phiX174 DNA/Hinfl 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 φX174 DNA/Hinfl Marker)

Fragment	Size	%	mass (ng/0.5µg)
1	726	13.5	67.4
2	713	13.2	66.2
3	553	10.3	51.3
4	500	9.3	46.4
5	427	7.9	39.6
6	417	7.7	38.7
7	413	7.7	38.3
8	311	5.8	28.9
9	249	4.6	23.1
10	200	3.7	18.6
11	151	2.8	14.0
12	140	2.6	13.0
13	118	2.2	11.0
14	100	1.9	9.3
15	82	1.5	7.6
16	66,66	1.2, 1.2	6.1, 6.1
17	48	0.9	4.5

#### Notice to users

Research and in vitro use only.

<sup>&</sup>lt;sup>⋆</sup>The mixture should be scaled up or down, depending on the width of the gel. Use 0.1-0.2 µg of DNA marker/mm of lane.