©LERUPSSP® DNA Size Marker 103.202-100/500 Lot No.: 8D5

## **Olerup SSP® DNA Size Marker**

Product number:	103.202-100/500
Lot number:	8D5
Volume:	Product No. 103.202-100 – 2 x 500 μl
	Product No. 103.202-500 – 10 x 500 μl
Concentration:	20 ng/µl
Format:	Ready-to-load, 10 µl per gel lane
Expiry date:	2018-03-01
Storage:	2-8°C

DESCRIPTION 103.202-100: 2 vials, 500 μl each. 103.202-500: 10 vials, 500 μl each. DNA ladder, 7 bands: 50, 100, 200, 300, 400, 500 and 1 000 bp, 20 ng/μl in 10 mM Tris-HCl (pH 9.0), 50 mM KCl, 5% (v/v) glycerol (99.5%), cresol red 100 ng/μl.

The DNA ladder contains 7 fragments of equal intensity:

<b>Fragment</b>	Number of base pairs
1	1000
2	500
3	400
4	300
5	200
6	100
7	50

**PRODUCT USE** The DNA Size marker can be resolved well in standard LE agarose gels of 1-2%, in NuSieve 3:1 or Metaphor agarose gels up to 4%. The DNA Size Marker is supplied in a *ready-to-load format*.

Typically load  $10 \ \mu l \ per \ gel \ lane$  of the ready-to-load DNA Size marker, equals 200 ng of DNA Size Marker.

**STORAGE** Store at 2-8°C.

**SHIPPING** The DNA Size Marker is shipped at ambient temperature.

**STABILITY** The DNA Size Marker is stable for 24 months when stored at 2-8°C.

## **CERTIFICATE OF ANALYSIS**

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10  $\mu I$  DNA Size Marker was size-separated on a 2% agarose gel stained by ethidium bromide.

*Results:* Seven fragments of 50, 100, 200, 300, 400, 500 and 1000 bp, were visible on a UV transilluminator.

Date of approval:

Approved by:

Production Quality Control

©LERUPSSP® DNA Size Marker 103.202-100/500 Lot No.: 8D5

**Declaration of Conformity** 

Product name: Product number: Lot number:	DNA Size Marker 103.202-100/500 8D5
Intended use:	DNA Size Marker
Manufacturer:	<i>Olerup</i> SSP AB Franzengatan 5 SE-112 51 Stockholm, Sweden <i>Phone:</i> +46-8-717 88 27 <i>Fax:</i> +46-8-717 88 18

We, *Olerup* SSP AB, hereby declare that this product, to which this Declaration of Conformity relates is in conformity with the following Standard(s) and other normative document(s) ISO 9001:2008 and EN ISO 13485:2012, following the provisions of the 98/79/EC Directive on *in vitro* diagnostic medical devices, Annex III, as transposed into the national laws of the Member States of the European Union.

The Technical Documentation File is maintained at *Olerup* SSP AB, Franzengatan 5, SE-112 51 Stockholm, Sweden.

Stockholm, Sweden 2016-Mar-21

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