

Lot No.: **0H3**

Olerup SSP[®] DNA Size Marker

Product number:	103.202-100/500
Lot number:	0H3
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:	2020-10-01
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:

<u>Fragment</u>	<u>Number of base pairs</u>
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 µl 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.

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CERTIFICATE OF ANALYSIS

Olerup SSP[®] DNA Size Marker

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Lot number: 0H3
Volume: Product No. 103.202-100 – 2 x 500 µl
Product No. 103.202-500 – 10 x 500 µl
Concentration: 20 ng/µl
Expiry date: 2020-10-01

10 µl 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

Lot No.: OH3

Declaration of Conformity

Product name: DNA Size Marker
Product number: 103.202-100/500
Lot number: OH3

Intended use: DNA Size Marker

Manufacturer: *Olerup* SSP AB
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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

Emil Jonsson
Head of QA

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