Olerup SSP® DNA Size Marker

Product number: 103.202-100/500

Lot number: 61N

Volume: Product No. 103.202-100 – 2 x 500 μl

Product No. 103.202-500 – 10 x 500 μ l

Concentration: 20 ng/ μl

Format: Ready-to-load, 10 µl per gel lane

Expiry date: 2014-March-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>	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 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.

103.202-100/500 Lot No.: **61 N**

CERTIFICATE OF ANALYSIS

Olerup SSP® DNA Size Marker

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Lot number: 61N

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: 2014-March-01

10 μ l 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: 2012-March-19

Approved by:

Production Quality Control

Lot No.: 61N

Declaration of Conformity

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

61N Lot number:

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 ISO 13485:2003, 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 2012-March-19

Ann-Cathrin Jareman Head of QA and Regulatory Affairs **DNA Size Marker Product Insert** Page 4 of 4

103.202-100/500 Lot No.: **61N**

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