Lot No.: 62S

Olerup SSP® DNA Size Marker

Product number: 103.202-100/500

Lot number: **62S**

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: 2015-June-01

2-8°C Storage:

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 na/ul

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:

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

The DNA Size marker can be resolved well in standard LE PRODUCT USE

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** μ**l per gel lane** of the ready-to-load DNA Size

marker, equals 200 ng of DNA Size Marker.

Store at 2-8°C. STORAGE

June 2013

Rev. No.: 00

The DNA Size Marker is shipped at ambient temperature. SHIPPING

The DNA Size Marker is stable for 24 months when stored at 2-STABILITY

8°C.

DNA Size Marker 103.202-100/500

Lot No.: 62S

CERTIFICATE OF ANALYSIS

Olerup SSP® DNA Size Marker

Product number: 103.202-100/-500

Lot number: **62S**

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

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

Concentration: **20 ng/μl Expiry date:** 2015-June-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: 2013-June-10

Approved by:

Production Quality Control

Nsa Olacessu

DNA Size Marker 103.202-100/500 Lot No.: **62S**

Declaration of Conformity

Product name: Product number:

DNA Size Marker 103.202-100/500

Lot number:

62S

Intended use:

DNA Size Marker

Manufacturer:

Olerup SSP AB Franzengatan 5

SE-112 51 Stockholm, Sweden

Phone: +46-8-717 88 27 **Fax:** +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 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 2013-June-10

Ann-Cathrin Jareman

Head of QA and Regulatory Affairs

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