103.202-100 Lot No.: **47M**

Olerup SSP® DNA Size Marker

Product number: 103.202-100

 $\begin{array}{lll} \text{Lot number:} & 47\text{M} \\ \text{Volume:} & 2 \times 500 \; \mu\text{l} \\ \text{Concentration:} & 20 \; \text{ng/} \; \mu\text{l} \\ \end{array}$

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

Expiry date: 2013-June-01

Storage: 2-8°C

DESCRIPTION 2 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** μ **I** 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

Product number: 103.202-100

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: 2011-August-19

Approved by:

Quality Control, Supervisor

103.202-100 Lot No.: **47M**

Declaration of Conformity

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

Lot number: 47M

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 13485:2003, 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.

The Authorized Representative located within the Community is: *Olerup* SSP AB.

Stockholm, Sweden 2011-August-19

Ann-Cathrin Jareman
Head of QA and Regulatory Affairs

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