

Lot No.: 5E6

Olerup SSP® DNA Size Marker for short gel runs

Product number: 103.203-100/500
Lot number: 5E6
Volume: Product No. 103.203-100 – 2 x 500 µl
 Product No. 103.203-500 – 10 x 500 µl
Concentration: 20 ng/µl
Format: Ready-to-load, 10 µl per gel lane
Expiry date: 2018-10-01
Storage: 2-8°C

DESCRIPTION 103.203-100: 2 vials, 500 µl each.
 103.203-500: 10 vials, 500 µl each.
 DNA ladder, 4 bands: 50, **200**, 500 and 1 000 bp.
 Double intensity of 200 bp band.
 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 4 fragments, double intensity of 200 bp band:

| <u>Fragment</u> | <u>Number of base pairs</u> |
|-----------------|-----------------------------|
| 1 | 1000 |
| 2 | 500 |
| 3 | 200 |
| 4 | 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 for short gel runs is supplied in a **ready-to-load format**. Typically load **10 µl per gel lane** of the ready-to-load DNA Size Marker for short gel runs, equals 200 ng of DNA Size Marker for short gel runs.

STORAGE Store at 2-8°C.

SHIPPING The DNA Size Marker for short gel runs is shipped at ambient temperature.

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

Change in revision R01 compared to R00:

1. The Certificate Of Analysis and Declaration Of Conformity date has been corrected.

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CERTIFICATE OF ANALYSIS

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10 µl DNA Size Marker was size-separated on a 2% agarose gel stained by ethidium bromide.

Results: Four fragments of 50, 200, 500 and 1000 bp were visible on a UV transilluminator. Double intensity of 200 bp fragment.

Date of approval: 2016-Oct-31

Approved by:



Production Quality Control

Declaration of Conformity

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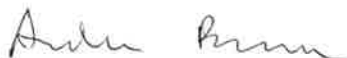
Intended use: DNA Size Marker for short gel runs

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
 2016-Oct-31



Andrea Boman
 Head of QA and Regulatory Affairs

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