Certificates

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101.903-24 - including Taq polymerase, IFU-01Visit www.olerup-ssp.com for 101.903-24u – without *Taq* polymerase, IFU-02"Instructions for Use" (IFU)

Lot No.: 86Y

Lot-specific information

CERTIFICATE OF ANALYSIS

Olerup SSP® DQA1*02,05;DQB1*02,03:02 SSP

Product number:

101.903-24 – including *Taq* polymerase

101.903-24u – without *Tag* polymerase

Lot number:

Expiry date:

2018-February-01

Number of tests: Number of wells per test: 24 17 + 1

Well specifications:

| Well No. | Production No. | Well No. | Production No. | Well No. | Production No. |
|----------|-----------------------|----------|-----------------------|----------|-----------------------|
| 1 | 2015-490-01 | 9 | 2015-490-10 | 17 | 2015-490-17 |
| 2 | 2015-490-02 | 10 | 2015-490-10B | | |
| 3 | 2015-490-03 | 11 | 2015-490-11 | | |
| 4 | 2015-490-04 | 12 | 2015-490-12 | | |
| 5 | 2015-490-06 | 13 | 2015-490-13 | | |
| 6 | 2015-490-07 | 14 | 2015-490-14 | | |
| 7 | 2015-490-08 | 15 | 2015-490-15 | | |
| 8 | 2015-490-09 | 16 | 2015-490-16 | | |

The negative control primer pairs, Production No. 2015-545-01, can detect contamination with PCR products diluted 10⁻⁷.

Results of Quality Control: No false positive or false negative amplifications

obtained.

Date of approval: 10150820

Approved by:

Production Quality Control

CE

Certificates

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101.903-24 – including *Taq* polymerase, IFU-01Visit <u>www.olerup-ssp.com</u> for 101.903-24u – without *Taq* polymerase, IFU-02"Instructions for Use" (IFU)

Lot No.: 86Y Lot-specific information

Declaration of Conformity

Product name: Olerup SSP® DQA1*02,05;DQB1*02,03:02

Product number: 101.903-24/24u

Lot number: 86Y

Intended use: DQA1*02, DQA1*05, DQB1*02 and DQB1*03 medium

resolution histocompatibility testing

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 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.

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

Stockholm, Sweden

2015-AUG-20

ennor Mala

Daniel Malica

Head of QA and Regulatory Affairs