DQA1*02,05;DQB1*02,03:02

Page 1 of 2

101.903-24 – including *Taq* polymerase, IFU-01 **101.903-24** u – without *Taq* polymerase, IFU-02

Visit <u>www.olerup.com</u> for "Instructions for Use" (IFU)

Lot No.: 6F0

Expiry date:

Lot-specific information

CERTIFICATE OF ANALYSIS

Olerup SSP® DQA1*02,05;DQB1*02,03:02

Product number: 101.903-24 – including *Taq* polymerase

101.903-24u - without Taq polymerase

Lot number:

2020-05-01

Number of tests: 24
Number of wells per test: 21+ 1

Well specifications:

Tren specifications.						
Well No.	Production No.	Well No.	Production No.	Well	No.	Production No.
1	2017-841-01	9	2017-841-09	17	2017	-841-16
2	2015-490-02	10	2017-841-21	18	2017	-841-17
3	2017-841-03	11	2017-841-10	19	2017	-841-18
4	2017-841-04	12	2017-841-11	20	2017	-841-19
5	2017-841-05	13	2017-841-12	21	2017	-841-20
6	2017-841-06	14	2017-841-13			
7	2017-841-07	15	2017-841-14			
8	2017-841-08	16	2017-841-15			

The negative control primer pairs, **Production No. 2017-845-01**, can detect contamination with PCR products diluted 10⁻⁷.

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

obtained.

Date of approval: 171130

Approved by:

Production Quality Control

CE

Certificates

Page 2 of 2

101.903-24 – including *Taq* polymerase, IFU-01 **101.903-24** u – without *Taq* polymerase, IFU-02

Visit <u>www.olerup.com</u> for "Instructions for Use" (IFU)

Lot No.: 6F0

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:

6F0

Intended use:

DQA1*02, DQA1*05, DQB1*02 and DQB1*03 medium resolution

histocompatibility testing

Manufacturer:

Olerup SSP AB

2017 1130

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

Emil Jonsson

Head of QA

CE