**OLERUP SSP** 

HLA-B\*42 Certificates

101.543-06 - including Taq pol., IFU-01 **101.543-06u – without** *Taq* **pol.**, IFU-02

Page 1 of 2 Visit www.labproducts.caredx.com for "Instructions for Use" (IFU)

Lot No.: 7H8

Lot-specific information

## **CERTIFICATE OF ANALYSIS**

Olerup SSP® HLA-B\*42

**Product number:** 101.543-06 - including *Taq* pol.

101.543-06u – without *Taq* pol.

Lot number:

**7H8** 

**Expiry date:** 

2021-10-01

Number of tests:

Number of wells per test:

15+1

## Well specifications:

Well No.	<b>Production No.</b>	Well No.	<b>Production No.</b>
1	2015-488-01	9	2015-488-09
2	2015-488-02	10	2015-488-10
3	2015-488-03	11	2015-488-11
4	2015-488-04	12	2015-488-12
5	2015-488-05	13	2015-488-13
6	2015-488-06	14	2015-488-14
7	2015-488-07	15	2018-993-15
8	2015-488-08		

The negative control primer pairs, Production No. 2018-947-01, can detect contamination with PCR products diluted 10<sup>-7</sup>.

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

obtained.

Date of approval: 2019-05-16

Approved by:

RebukeSelne

**Production Quality Control** 

For In Vitro Diagnostic Use MA100 v02 CoA\_DoC IVD Annex II List B Date: April 2019, Rev. No: 00 OLERUP SSP

HLA-B\*42 101.543-06 - including Tag pol., IFU-01 **101.543-06u** – without *Taq* pol., IFU-02

Certificates

Page 2 of 2 Visit www.labproducts.caredx.com for "Instructions for Use" (IFU)

Lot No.: 7H8

Lot-specific information

## **Declaration of Conformity**

Product name:

Olerup SSP® HLA-B\*42

Product number:

101.543-06/06u

Lot number:

**7H8** 

Intended use:

HLA-B\*42 high resolution histocompatibility testing

Manufacturer:

Olerup SSP AB Franzengatan 5

SE-112 51 Stockholm, Sweden **Phone:** +46-8-508 939 00 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) EN ISO 13485:2016, following the provisions of the 98/79/EC Directive on in vitro diagnostic medical devices, Annex II List B, conformity assessed using Annex IV, 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.

Notified Body: Lloyd's Register Quality Assurance Limited, 1 Trinity Park, Bickenhill

Lane, Birmingham B37 7ES, United Kingdom.

(Notified Body number: 0088.)

Stockholm, Sweden 2019056

Date:

Emil Jonsson

Head of Quality Assurance