

Lot No.: 4H2

Lot-specific Information

CERTIFICATE OF ANALYSIS**Olerup SSP[®] HLA-B*82**

Product number:

101.552-06 – including *Taq* polymerase101.552-06u – without *Taq* polymerase

Lot number:

4H2

Expiry date:

2021-08-01

Number of tests:

6

Number of wells per test:

5+1

Well specifications:

Well No.	Production No.
1	2018-916-01
2	2015-556-02
3	2016-709-03
4	2016-709-04
5	2019-025-05

The negative control primer pairs, **Production No. 2018-947-01**, can detect contamination with PCR products diluted 10^{-7} .

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

Approved by:
2019-02-22**Production Quality Control**

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For *In Vitro* Diagnostic Use

MA100 v01 CoA_DoC IVD Annex II List B

Date: February 2019, Rev. No: 00

Lot No.: **4H2**

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Declaration of Conformity**Product name:** *Olerup* SSP® HLA-B*82**Product number:** 101.552-06/06u**Lot number:** 4H2**Intended use:** HLA-B*82 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

Date: 20190228



Emil Jonsson
Head of Quality Assurance

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