

101.625-06 – including *Taq* polymerase, IFU-01
101.625-06u – without *Taq* polymerase, IFU-02

Visit www.olerup-ssp.com for
“Instructions for Use” (IFU)

Lot No.: **0D2**

Lot-specific information

www.olerup-ssp.com

CERTIFICATE OF ANALYSIS

Olerup SSP® HLA-C*14 SSP

Product number: 101.625-06 – including *Taq* polymerase
101.625-06u – without *Taq* polymerase
Lot number: 0D2
Expiry date: 2018-April-01
Number of tests: 6
Number of wells per test: 30+1

Well specifications:

Well No.	Production No.	Well No.	Production No.
1	2013-162-01	17	2013-162-17
2	2013-162-02	18	2015-583-18
3	2013-162-03	19	2014-402-19
4	2013-162-04	20	2013-162-20
5	2013-162-05	21	2013-162-21
6	2013-162-06	22	2013-162-22
7	2013-162-07	23	2013-162-23
8	2013-162-08	24	2014-402-24
9	2013-162-09	25	2014-402-25
10	2013-162-10	26	2014-402-26
11	2013-162-11	27	2014-402-27
12	2013-162-12	28	2014-402-28
13	2013-162-13	29	2015-583-29
14	2013-162-14	30	2015-583-30
15	2013-162-15		
16	2013-162-16		

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: 2015-10-19

Approved by:



Production Quality Control

101.625-06 – including *Taq* polymerase, IFU-01
101.625-06u – without *Taq* polymerase, IFU-02

Visit www.olerup-ssp.com for
“Instructions for Use” (IFU)

Lot No.: **0D2**

Lot-specific information

www.olerup-ssp.com

Declaration of Conformity

Product name: Olerup SSP® HLA-C*14
Product number: 101.625-06/06u
Lot number: 0D2

Intended use: HLA-C*14 high 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-Oct-19

Daniel Malica
Head of QA and Regulatory Affairs