**101.554-06** – including *Taq* pol., IFU-01 **101.554-06u** – without *Taq* pol., IFU-02

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

Lot No.: 18R

Lot-specific information

## **CERTIFICATE OF ANALYSIS**

Olerup SSP® HLA-B\*59 SSP

Product number: 101.554-06 – including *Taq* pol.

101.554-06u - without Tag pol.

Lot number: 18R

Expiry date: 2015-March-01

Number of tests: 6 Number of wells per test: 8

## Well specifications:

Well No.	Production No.
1	2012-060-01
2	2006-248-02
3	2006-248-03
4	2010-814-04
5	2009-644-06
6	2008-505-06
7	2009-644-07
8	2009-644-08

The specificity of each primer solution of the HLA-B\*59 primer set has been tested against 48 well characterized IHWC cell line DNAs.

No DNAs carrying the alleles to be amplified by primer solution 4 and 8 were available. The specificities of the primers in primer solution 4 were tested by separately adding one additional 5'-primer, respectively one additional 3'-primer. In primer solution 8 it was only possible to test the 5'-primer, the 3'-primer was not possible to test.

**Results:** No false positive or false negative amplifications were obtained.

Date of approval: 2012-September-21

Approved by:

Du Olaus

**Production Quality Control** 

0088

101.554-06 – including *Taq* pol., IFU-01 101.554-06u – without *Taq* pol., IFU-02 Visit <u>www.olerup-ssp.com</u> for "Instructions for Use" (IFU)

Lot No.: 18R

Lot-specific information

## **Declaration of Conformity**

Product name:

Olerup SSP® HLA-B\*59

Product number:

101.554-06/06u

Lot number:

18R

Intended use:

HLA-B\*59 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 ISO 13485:2003, 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.

Notified Body: Lloyd's Register Quality Assurance Limited, Hiramford, Middlemarch Office Village, Siskin Drive, Coventry CV3 4FJ, United Kingdom. (Notified Body number: 0088.)

Stockholm, Sweden 2012-September-21

Ann-Cathrin Jareman

Head of QA and Regulatory Affairs

Alar-lathouis Inecuais