CERTIFICATE OF ANALYSIS

Olerup SSP® HLA-C*04:09N SSP

Product number:	**101.861-12 – including *Taq* polymerase** **101.861-12u – without *Taq* polymerase**
Lot number:	**51X**
Expiry date:	**2017-May-01**
Number of tests:	**12**
Number of wells per test:	**2+1**

Well specifications:

Well No.	Production No.
1	2014-435-01
2	2014-435-02

The negative control primer pairs, **Production No. 2014-382-01**, 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-January-09

Approved by:

Kurin Claesson

Production Quality Control

Lot-specific information

Declaration of Conformity

Product name:	*Olerup* SSP® HLA-C*04:09N
Product number:	101.861-12/12u
Lot number:	51X
Intended use:	HLA-C*04:09N 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 13485:2012, 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-January-09

Daniel Malica
Head of QA and Regulatory Affairs