

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: 3E3

Expiry date: 2019-03-01

Number of tests: 12

Number of wells per test: 2+1

Well specifications:

Well No.	Production No.
1	2015-584-01
2	2014-435-02

The negative control primer pairs, **Production No. 2016-695-01**, can detect contamination with PCR products diluted 10⁻⁷.

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

Date of approval: 14 Sep 2016

Approved by:



Production Quality Control

HLA-C*04:09N
101.861-12 – including Taq polymerase, IFU-01
101.861-12u – without Taq polymerase, IFU-02
Lot No.: 3E3

Certificates

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

Lot-specific information
Declaration of Conformity

Product name: Olerup SSP® HLA-C*04:09N
Product number: 101.861-12/12u
Lot number: 3E3

Intended use: HLA-C*04:09N histocompatibility testing

Manufacturer: Olerup SSP AB
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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
 2016-Sep-14



Daniel Malica
 Head of QA and Regulatory Affairs