

101.843-12 – including *Taq* polymerase, IFU-01
101.843-12u – without *Taq* polymerase, IFU-02

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

Lot No.: **55N**

Lot-specific information

CERTIFICATE OF ANALYSIS

Olerup SSP® HLA-A*23 Add-on SSP

Product number: 101.843-12 – including *Taq* polymerase
101.843-12u – without *Taq* polymerase
Lot number: 55N
Expiry date: 2014-August-01
Number of tests: 12
Number of wells per test: 3

Well specifications:

Well No.	Production No.
1	2012-960-01
2	2012-960-02
3	2012-992-03

The specificity of each primer solution of the kit has been tested against 48 well characterized IHWC cell line DNAs.

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

Date of approval: 2012-March-09

Approved by:



Production Quality Control

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Lot-specific information

Declaration of Conformity

Product name: Olerup SSP® HLA-A*23 Add-on
Product number: 101.843-12/12u
Lot number: 55N

Intended use: HLA-A*23 histocompatibility testing

Manufacturer: Olerup SSP AB
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Phone: +46-8-717 88 27
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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 ISO 13485:2003, following the provisions of the 98/79/EC Directive on *in vitro* diagnostic medical devices, Annex II List B, 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-May-10


Ann-Cathrin Jareman
Head of QA and Regulatory Affairs