

HLA-B\*27 unit dose single well

Certificates

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**101.911-96** – including *Taq* pol., IFU-01 **101.911-96u** – without *Taq* pol., IFU-02

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

Lot No.: 49V

Lot-specific information

www.olerup-ssp.com

## CERTIFICATE OF ANALYSIS

Olerup SSP® HLA-B\*27 unit dose single well SSP

**Product number:** 

101.911-96 - including *Taq* polymerase

101.911-96u – without *Taq* polymerase

Lot number:

49V

**Expiry date:** 

2016-September-01

Number of tests:

96

Number of wells per test:

1

Well specifications:

Well No.

**Production No.** 

1

2014-323-01

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

obtained.

Date of approval:

Asa Olauss

2014-March-19

Approved by:

**Production Quality Control** 



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Lot No.: 49V Lot-specific information

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**Declaration of Conformity** 

Product name:

Olerup SSP® HLA-B\*27 unit dose single well

**Product number:** 

101.911-96/96u

Lot number:

101.911-96/960 49V

Intended use:

HLA-B\*27 low 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 13485:2012, 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 2014-March-19

Ann-Cathrin Jareman

Head of QA and Regulatory Affairs

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