

102.101-01 – including *Taq* polymerase, IFU-01

102.101-01u – without *Taq* polymerase, IFU-02

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

Lot No.: 2D1

Lot-specific information

www.olerup-ssp.com

CERTIFICATE OF ANALYSIS

Olerup SSP® HLA Wipe Test – Negative Control

Product number: 102.101-01 – including *Taq* polymerase
Product number: 102.101-01u – without *Taq* polymerase
Lot number: 2D1
Expiry date: 2018-May-01
Number of tests: 96
Number of wells per test: 1-2

Well specification:

Well No.	Production No.
1	2015-545-01

The negative control primer solution has been tested in a dilution series of the corresponding PCR products, 1 to 10³ down to 1 to 10⁹.

The Positive Control DNA has been tested with the HLA Wipe Test kit and gives rise to PCR amplicons.

The negative control primer pairs can detect contamination with the corresponding PCR products diluted 1 to 10⁷.

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

Date of approval: 2015-12-07

Approved by:



Production Quality Control

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Declaration of Conformity

Product name: Olerup SSP® HLA Wipe Test – Negative Control
Product number: 102.101-01/01u
Lot number: 2D1

Intended use: Detection of contamination with HLA amplicons.

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 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.

Stockholm, Sweden
2015-Dec-17



Daniel Malica
Head of QA and Regulatory Affairs