

101.101-48/12 – including *Taq* pol., IFU-01
 101.101-48u/12u – without *Taq* pol., IFU-02

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

Lot No.: **09Y**

Lot-specific information

CERTIFICATE OF ANALYSIS

Olerup SSP® DR low resolution

Product number: 101.101-48/12 – including *Taq* pol.
 101.101-48u/12u – without *Taq* pol.
Lot number: 09Y
Expiry date: 2017-August-01
Number of tests: 48 tests – Product No. 101.101-48/48u
 12 tests – Product No. 101.101-12/12u
Number of wells per test: 31 + 1

Well specifications:

Well No.	Production No.	Well No.	Production No.
1	2014-401-01	17	2015-492-17
2	2014-401-02	18	2014-401-18
3	2015-492-03	19	2015-492-19
4	2014-401-04	20	2015-492-20
5	2014-394-05	21	2015-492-21
6	2014-401-06	22	2015-492-22
7	2014-401-07	23	2015-492-23
8	2014-401-08	24	2015-492-24
9	2015-492-09	25	2015-492-25
10	2014-394-10	26	2015-492-26
11	2014-401-11	27	2015-492-27
12	2014-401-12	28	2015-492-28
13	2014-401-13	29	2014-401-29
14	2014-401-14	30	2014-401-30
15	2014-401-15	31	2014-401-31
16	2015-492-16		

The negative control primer pairs, **Production No. 2015-499-01**, 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-March-03

Approved by:

Karin Mattsson

Production Quality Control

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Lot-specific information

Declaration of Conformity

Product name: *Olerup* SSP® DR low resolution
Product number: 101.101-48/48u, -12/12u
Lot number: 09Y

Intended use: DRB1 low resolution histocompatibility testing

Manufacturer: *Olerup* SSP AB
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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
 2015-March-03



Daniel Malica
 Head of QA and Regulatory Affairs