

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.: **93S**

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: 93S
Expiry date: 2016-April-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.	Well No.	Production No.
1	2013-219-01	13	2012-090-13	25	2012-055-25
2	2012-090-02	14	2013-254-14	26	2012-055-26
3	2012-090-03	15	2012-090-15	27	2013-254-27
4	2012-090-04	16	2012-090-16	28	2012-055-28
5	2012-090-05	17	2012-090-17	29	2012-090-29
6	2012-090-06	18	2012-090-18	30	2012-090-30
7	2012-090-07	19	2012-090-19	31	2012-090-31
8	2012-090-08	20	2012-219-20		
9	2012-254-09	21	2012-055-21		
10	2012-090-10	22	2012-055-22		
11	2012-090-11	23	2012-055-23		
12	2012-090-12	24	2013-254-24		

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

Additional 5'- and 3'-primers in primer solutions 11, 18 and 27 were tested by separately adding one 3'-primer, respectively one 5'-primer.

Additional 5'-primers in primer solutions 6, 12, 15 and 16 were tested by separately adding one 3'-primer. Additional 3'-primers in primer solutions 1, 3, 4, 9, 10, 20 and 22 were tested by separately adding one 5'-primer.

One, two or three of the 5'-primers in primer solution 1, 3, 4, 8 to 10, 13, 15, 16, 20, 22 and 27 were not possible to test. One or two of the 3'-primers in primer solution 1, 3, 4, 9, 13, 15, 21, 26 and 30 were not possible to test.

The negative control primer pairs, **Production No. 2013-271-01**, can detect contamination with PCR products diluted 10^{-7} .

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

Date of approval: 2013-October-24

Approved by:



Production Quality Control

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Declaration of Conformity

Product name: *Olerup* SSP® DR low resolution
Product number: 101.101-48/48u, -12/12u
Lot number: 93S

Intended use: DRB1 low resolution histocompatibility testing

Manufacturer: *Olerup* SSP AB
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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
2013-October-24



Ann-Cathrin Jareman
Head of QA and Regulatory Affairs