

PRB5 Certificates Page 1 of 2

101.123-24/06 – including *Taq* **polymerase**, IFU-01 **101.123-24u/06u – without** *Taq* **polymerase**, IFU-02

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

Lot No.: **25V**

Lot-specific information

CERTIFICATE OF ANALYSIS

Olerup SSP® DRB5 SSP

Product number: 101.123-24/06 – including *Tag* pol.

101.123-24u/06u - without Taq pol.

Lot number: 25V

Expiry date: 2016-July-01

Number of tests: 24 test – Product No. 101.123-24/24u

6 tests - Product No. 101.123-06/06u

Number of wells per test: 15+1

Well specifications:

Well No.	Production No.	Well No.	Production No.
1	2012-974-01	9	2012-974-09
2	2012-974-02	10	2012-974-10
3	2012-974-03	11	2012-974-11
4	2012-096-04	12	2012-974-12
5	2012-974-05	13	2012-974-13
6	2012-974-06	14	2012-974-14
7	2013-292-07	15	2012-974-15
8	2012-974-08		

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

Results of Quality Control:

No false positive or false negative

amplifications obtained.

Date of approval: 2014-February-06

Approved by:

Production Quality Control





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101.123-24/06 – including *Taq* **polymerase,** IFU-01 **101.123-24u/06u – without** *Taq* **polymerase,** IFU-02

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Lot No.: 25V Lot-specific information

Declaration of Conformity

Product name: Olerup SSP® DRB5 **Product number:** 101.123-24/06, -24u/06u

Lot number: 25V

Intended use: DRB5 high 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-February-06

Ann-Cathrin Jareman

Head of QA and Regulatory Affairs

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