

CERTIFICATE OF ANALYSIS**Olerup SSP® DRB1*15 SSP**

Product number: 101.125-24u/06u – without Taq pol.
Lot number: 27M
Expiry date: 2013-November-01
Number of tests: 24 test – Product No. 101.125-24u
 6 tests – Product No. 101.125-06u
Number of wells per test: 32

Well specifications:

Well No.	Production No.	Well No.	Production No.	Well No.	Production No.
1	2011-868-01	13	2011-868-13	25	2011-868-25
2	2011-868-02	14	2011-868-14	26	2011-868-26
3	2011-868-03	15	2011-868-15	27	2011-868-27
4	2011-868-04	16	2011-868-16	28	2011-868-28
5	2011-868-05	17	2011-868-17	29	2011-868-29
6	2011-868-06	18	2011-868-18	30	2011-868-30
7	2011-868-07	19	2011-868-19	31	2011-868-31
8	2011-868-08	20	2011-868-20	32	2011-868-32
9	2011-868-09	21	2011-868-21		
10	2011-868-10	22	2011-868-22		
11	2011-868-11	23	2011-868-23		
12	2011-868-12	24	2011-868-24		

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

No DNAs carrying the alleles to be amplified by primer solutions 8, 9, 11 to 20 and 22 to 32 were available. The specificities of the primers in primer solutions 8, 11 to 13, 15, 19, 20, 22, 23, 26, 28 and 32 were tested by separately adding one additional 5'-primer, respectively one additional 3'-primer. In primer solutions 14, 16 to 18 and 29 to 31 it was only possible to test the 5'-primers, the 3'-primers were not possible to test. In primer solutions 9, 24, 25 and 27 it was only possible to test the 3'-primers, the 5'-primers were not possible to test. In primer solutions 6 and 22 one respective two of the 5'-primers were not possible to test, and in primer solutions 2, 3, 8, 13, 19, 20, 23 and 32 one or two of the 3'-primers were not possible to test.

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

Date of approval: 2011-May-26

Approved by:



Quality Control, Supervisor

Declaration of Conformity

Product name: Olerup SSP® DRB1*15

Product number: 101.125-24u/06u

Lot number: 27M

Intended use: DRB1*15 high resolution histocompatibility testing

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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 13485:2003, 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.

The Authorized Representative located within the Community is: Olerup SSP AB.

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
2011-May-26



Olle Olerup
Managing Director