101.615-24/04 – including *Taq* pol., IFU-01 101.615-24u/04u - without Taq pol., IFU-02

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

Lot No.: 42S Lot-specific information

CERTIFICATE OF ANALYSIS

Olerup SSP® HLA-C*07 SSP

Product number: 101.615-24/04 – including *Taq* pol.

101.615-24u/04u – without *Taq* pol.

Lot number: **42S**

Expiry date: 2015-December-01

Number of tests: 24 tests - Product No. 101.615-24/24u

4 tests - Product No. 101.615-04/04u

Number of wells per test: 74

Wall enecifications:

Well No.	Production No.	Well No.	Production No.	Well No.	Production No
1	2010-698-01	33	2011-895-33	65	2013-193-65
2	2011-895-02	34	2013-193-34	66	2013-193-66
3	2011-895-03	35	2013-193-35	67	2013-193-67
4	2010-698-04	36	2012-08136	68	2013-193-68
5	2012-081-05	37	2012-081-37	69	2013-193-69
6	2010-698-06	38	2013-193-38	70	2013-193-70
7	2010-698-07	39	2010-698-39	71	2013-193-71
8	2010-698-08	40	2012-081-40	72	2013-193-72
9	2013-193-09	41	2010-698-41	73	2013-193-73
10	2010-698-10	42	2012-081-42	74	2013-193-74
11	2012-005-11	43	2012-081-43		
12	2010-698-12	44	2012-081-44	1	
13	2010-698-13	45	2012-081-45	1	
14	2012-081-14	46	2012-005-46	1	
15	2012-081-15	47	2012-005-47	1	
16	2013-193-16	48	2012-081-48		
17	2010-698-17	49	2012-005-49		
18	2012-081-18	50	2012-005-50	1	
19	2013-193-19	51	2012-081-51	1	
20	2012-081-20	52	2012-081-52	1	
21	2010-698-21	53	2012-081-53	1	
22	2010-698-22	54	2012-081-54		
23	2012-081-23	55	2012-081-55	1	
24	2013-193-24	56	2012-081-56]	
O.F.	2040 000 05		2042 004 57	1	
25	2010-698-25	57	2012-081-57	-	
26	2010-698-26	58	2012-081-58	-	
27	2012-081-27	59	2012-081-59	-	
28	2011-895-28	60	2012-081-60	-	
29	2011-895-29	61	2012-081-61	-	
30	2012-081-30	62	2012-081-62	-	
31	2012-005-31	63	2012-081-63	-	
32	2010-698-32	64	2012-081-64	J	

101.615-24/04 – including *Taq* pol., IFU-01 **101.615-24u/04u** – without *Taq* pol., IFU-02

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Lot No.: 42S

Lot-specific information

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

No DNAs carrying the alleles to be amplified by primer solutions 5, 10, 11, 13 to 15, 18, 19, 23 to 31, 34 to 36, 38, 39, 41 to 44 and 46 to 74 were available.

The specificities of the primers in primer solutions 5, 10, 11, 13 to 15, 18, 19, 24 to 27, 31, 36, 42 to 44, 46 to 50, 53, 54, 56 to 60, 63, 64, 71, 72 and 74 were tested by separately adding one 5'-primer, respectively one 3'-primer.

In primer solutions 35, 51, 61, 65 to 70 and 73 it was only possible to test the 5'-primer, the 3'-primer were not possible to test.

In primer solutions 23, 28 to 30, 34, 38, 39, 41, 52, 55 and 62 it was only possible to test the 3'-primers, the 5'-primers were not possible to test.

In primer solutions 8, 11, 20, 22, 26, 33, 42, 44, 46 to 49, 53, 56, 57, 59 and 63 one or two 5'-primers were not possible to test, and in primer solutions 5, 9, 13, 15, 18, 21, 25, 32, 36, 37, 40, 43, 46 to 48, 50, 54, 58, 60, 63, 64 and 74 one or two 3'-primers were not possible to test.

Additional primers in primer solutions 2, 8, 20, 21, 33, 37, 40 and 45 were tested by separately adding either one 5'-primer or one 3'-primer.

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

Date of approval: 2013-July-16

Approved by:

Production Quality Control

Lacin Mattre

101.615-24/04 – including *Taq* pol., IFU-01 **101.615-24u/04u** – without *Taq* pol., IFU-02

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

Lot No.: 42S

Lot-specific information

Declaration of Conformity

Product name: Product number:

Olerup SSP® HLA-C*07 101.615-24/24u, -04/04u

Lot number:

42S

Intended use:

HLA-C*07 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 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 2013-July-16

Ann-Cathrin Jareman

Head of QA and Regulatory Affairs

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