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

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Lot No.: 66N 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: 66N

Expiry date: 2014-October-01

Number of tests: 24 tests – Product No. 101.615-24 4 tests – Product No. 101.615-04

Number of wells per test: 64

Well specifications:

Well No.	Production No.	Well No.	Production No.	Well No.	Production No.
1	2010-698-01	25	2010-698-25	49	2012-005-49
2	2011-895-02	26	2010-698-26	50	2012-005-50
3	2011-895-03	27	2012-005-27	51	2012-005-51
4	2010-698-04	28	2011-895-28	52	2012-005-52
5	2012-005-05	29	2011-895-29	53	2012-005-53
6	2010-698-06	30	2011-895-30	54	2012-005-54
7	2010-698-07	31	2012-005-31	55	2012-005-55
8	2010-698-08	32	2010-698-32	56	2012-005-56
9	2010-698-09	33	2011-895-33	57	2012-005-57
10	2010-698-10	34	2012-005-34	58	2012-005-58
11	2012-005-11	35	2012-005-35	59	2012-005-59
12	2010-698-12	36	2010-698-36	60	2012-005-60
13	2010-698-13	37	2010-698-37	61	2012-005-61
14	2010-698-14	38	2010-698-38	62	2012-005-62
15	2012-005-15	39	2010-698-39	63	2012-005-63
16	2012-005-16	40	2010-698-40	64	2012-005-64
17	2010-698-17	41	2010-698-41		
18	2010-698-18	42	2012-005-42	1	
19	2010-698-19	43	2012-005-43	1	
20	2012-005-20	44	2012-005-44	1	
21	2010-698-21	45	2010-698-45	1	
22	2010-698-22	46	2012-005-46	1	
23	2011-841-23	47	2012-005-47	1	
24	2011-895-24	48	2010-698-48	1	

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 39, 41 to 44 and 46 to 64 were available.

The specificities of the primers in primer solutions 5, 10, 11, 13 to 15, 18, 19, 24 to 27, 31, 36 to 38, 42 to 44, 46 to 50, 53, 54, 56 to 60, 63 and 64 were tested by separately adding one 5'-primer, respectively one 3'-primer. In primer solutions 35, 51 and 61 it was only possible to test the 5'-primer, the 3'-primer

101.615-24/04 – including *Taq* **pol., IFU-01 101. 615-24u/04u – without** *Taq* **pol., IFU-02**

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Lot No.: 66N Lot-specific information

were not possible to test. In primer solutions 23, 28 to 30, 34, 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, 37, 38, 42, 44, 46 to 49, 53, 56, 59 and 63 one or two 5'-primers were not possible to test, and in primer solutions 5, 9, 13, 15, 18, 19, 21, 25, 32, 38, 40, 43, 46 to 48, 50, 54, 58, 60, 63 and 64 one or two 3'-primers were not possible to test.

Additional primers in primer solutions 2, 8, 9, 20, 21 and 33 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: 2012-May-11

Approved by:

Production Quality Control

Karin Chutton

HLA-C*07 Product Insert Page 37 of 40

101.615-24/04 – including *Taq* **pol.,** IFU-01 **101. 615-24u/04u – without** *Taq* **pol.,** IFU-02

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Lot No.: 66N Lot-specific information

Declaration of Conformity

Product name: Olerup SSP® HLA-C*07 **Product number:** 101.615-24/24u, -04/04u

Lot number: 66N

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 13485:2003, 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 2012-May-11

Ann-Cathrin Jareman

Head of QA and Regulatory Affairs