CERTIFICATE OF ANALYSIS

Olerup SSP® HLA-B*56 SSP

Product number: 101.571-06u – without *Taq* polymerase
Lot number: 60M
Expiry date: 2014-March-01
Number of tests: 6
Number of wells per test: 24

Well specifications:

Well No.	Production No.	Well No.	Production No.	Well No.	Production No.
1	2010-750-01	9	2011-893-09	17	2011-893-17
2	2010-750-02	10	2010-750-10	18	2011-893-18
3	2010-750-03	11	2010-750-11	19	2010-750-19
4	2010-750-04	12	2010-750-12	20	2010-750-20
5	2010-750-05	13	2011-893-13	21	2010-750-21
6	2011-893-06	14	2010-750-14	22	2010-750-22
7	2010-750-07	15	2010-750-15	23	2010-750-23
8	2010-750-08	16	2010-750-16	24	2010-750-24

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, 19, 21 and 23 were available. The specificities of the primers in primer solutions 8 and 23 were tested by separately adding one additional 5'-primer, respectively one or two additional 3'-primers. In primer solutions 19 and 21, it was only possible to test the 5'-primers, the 3'-primers were not possible to test. In primer solutions 1, 2, 3 and 15 one or two 3'-primers were not possible to test, and in primer solutions 13 and 23, one 5'-primer was not possible to test. One additional 5'-primer and on additional 3'-primer in primer solution 6 and 5'-primer in primer solution 15 were tested by separately adding one 5'-primer or 3'-primer.

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

Date of approval: 2011-November-28

Approved by:

Ísa Olausson

Production Quality Control

Declaration of Conformity

Product name: *Olerup* SSP® HLA-B*56
Product number: 101.571-06u
Lot number: 60M

Intended use: HLA-B*56 high resolution histocompatibility testing

Manufacturer: *Olerup* SSP AB
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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.

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-November-28

Ann-Cathrin Jareman
Head of QA and Regulatory Affairs