Lot No.: 90K

Lot-specific information

www.olerup-ssp.com

CERTIFICATE OF ANALYSIS

Olerup SSP® HLA-B*14 SSP

Product number:

101.524-12u - without Taq polymerase

Lot number:

90K

Expiry date:

2013-July-01

Number of tests:

12

Number of wells per test:

16

Well specifications:

Well No.	Production No.	Well No.	Production No.
1	2010-813-01	9	2009-645-09
2	2010-813-02	10	2010-813-10
3	2010-813-03	11	2010-813-11
4	2010-813-04	12	2010-813-12
5	2007-323-05	13	2010-813-13
6	2010-813-06	14	2010-813-14
7	2007-323-07	15	2010-813-15
8	2009-645-08	16	2010-813-16

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 solution 4 to 6, 9, 15 and 16 were available. The specificities of the primers in primer solution 4 to 6 were tested by separately adding one additional 5'-primer, respectively one additional 3'-primer. In primer solution 9, 15 and 16 it was only possible to test the 5'primer, the 3'-primer was not possible to test. In primer solutions 5 and 6, one 5'primer was not possible to test, and in primer solutions 5 and 12 one 3'-primer was not possible to test.

Additional primers in primer solution 10 were tested by separately adding one additional 5'-primer, respective one additional 3'-primer.

Results:

No false positive or false negative amplifications were obtained.

Date of approval: 2011-February -04

Approved by:

Quality Control, Supervisor

Product Insert Page 22 of 24 HLA-B*14 General "Instructions for Use"

101.524-12u - without Tag polymerase

IFU -02 Rev. No. 02 can be downloaded from

Lot No.: 90K

Lot-specific information

www.olerup-ssp.com

Declaration of Conformity

Product name:

Olerup SSP® HLA-B*14

Product number:

101.524-12u

Lot number:

90K

Intended use:

HLA-B*14 high resolution histocompatibility testing

Manufacturer:

Olerup SSP AB Hasselstigen 1

SE-133 33 Saltsjöbaden, 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 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, Hasselstigen 1, SE-133 33 Saltsjöbaden, 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.)

Saltsjöbaden, Sweden 2011-February-04

Olle Olerup

Managing Director