Product Insert Page 21 of 24 **HLA-B\*78** 

101.551-06u - without Tag polymerase

General "Instructions for Use"

IFU-02 Rev. No. 03 can be downloaded from Lot-specific information www.olerup-ssp.com

## CERTIFICATE OF ANALYSIS

Olerup SSP® HLA-B\*78 SSP

Product number: 101.551-06u – without *Taq* polymerase

48K Lot number:

2012-September-01 Expiry date:

Number of tests: Number of wells per test: 15

## Well specifications:

Lot No.: 48K

Well No.	Production No.	Well No.	Production No.
1	2009-573-01	9	2009-573-09
2	2009-573-02	10	2010-765-10
3	2009-573-03	11	2009-573-11
4	2009-573-04	12	2009-573-12
5	2009-573-05	13	2009-573-13
6	2009-573-06	14	2010-765-14
7	2009-573-07	15	2010-765-15
8	2009-573-08		

The specificity of each primer solution of the HLA-B\*78 primer set has been tested against 48 well characterized cell line DNAs.

No DNAs carrying the alleles to be amplified by primer solution 4 and 13 were available. The specificities of the primers in primer solution 4 were tested by separately adding one additional 3'-primer, respectively one additional 5'-primer. In primer solution 13, only the 5'-primer could be tested, the 3'-primer was not possible to test

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

Date of approval: 2011-December-06

Approved by:

December 2011 Rev. No.: 01u

**Production Quality Control** 

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101.551-06u - without Tag polymerase

General "Instructions for Use"

www.olerup-ssp.com

IFU-02 Rev. No. 03 can be downloaded from Lot No.: 48K Lot-specific information

**Declaration of Conformity** 

Product name:

Olerup SSP® HLA-B\*78

Product number:

101.551-06u

Lot number:

48K

Intended use:

HLA-B\*78 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 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-December-06

Ann-Cathrin Jareman

Head of QA and Regulatory Affairs