101.331-24u/06u - without Tag polymerase

General "Instructions for Use"

IFU-02 Rev. No. 01 can be downloaded from

Lot No.: 53K

Lot-specific information

www.olerup-ssp.com

CERTIFICATE OF ANALYSIS

Olerup SSP® DPA1 SSP

Product number:

101.331-24u/06u - without Tag pol.

Lot number:

53K

Expiry date:

2012-September-01

Number of tests:

24 test - Product No. 101.331-24u 6 tests - Product No. 101.331-06u

16

Number of wells per test:

Well specifications:

Well No.	Production No.	Well No.	Production No.
1	2009-682-01	9	2009-682-09
2	2009-682-02	10	2009-682-10
3	2010-774-03	11	2009-682-11
4	2009-682-04	12	2009-682-12
5	2009-682-05	13	2009-682-13
6	2009-682-06	14	2009-682-14
7	2009-682-07	15	2009-682-15
8	2009-682-08	16	2009-682-16

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

No DNAs carrying the allele to be amplified by primer solutions 6, 13, 14 and 16 were available. The specificities of the primers in primer solutions 6 and 14 were tested by separately adding one additional 5'-primer and one additional 3'primer, respectively. In primer solutions 13 and 16 it was only possible to test the 3'-primers, the 5'-primers were not possible to test. In primer solution 6, one 5'-primer and one 3'-primer was not possible to test.

Results:

No false positive or false negative amplifications were obtained.

Date of approval: 2010-November-24

Approved by:

Approved by:

Quality Control, Supervisor

DPA1 Product Insert Page 11 of 12 101.331-24u/06u – without *Taq* polymerase General "Instructions for Use"

IFU-02 Rev. No. 01 can be downloaded from

Lot No.: 53K

Lot-specific information

www.olerup-ssp.com

Declaration of Conformity

Product name: Product number: Olerup SSP® DPA1 101.331-24u/06u

Lot number:

53K

Intended use:

HLA-DPA1 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 III, as transposed into the national laws of the Member States of the European Union.

The Technical Construction 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.

Saltsjöbaden, Sweden 2010-November-24

Olle Olerup

Managing Director