

101.427-06 – including *Taq* pol., IFU-01 Rev. No. 03  
 101.427-06u – without *Taq* polymerase, IFU-02 Rev. No. 03

Visit [www.olerup-ssp.com](http://www.olerup-ssp.com) for  
 "Instructions for Use" (IFU)

Lot No.: **09R**

Lot-specific information

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## CERTIFICATE OF ANALYSIS

### Olerup SSP® HLA-A\*66 SSP

**Product number:** 101.427-06 – including *Taq* pol.  
 101.427-06u – without *Taq* pol.  
**Lot number:** 09R  
**Expiry date:** 2015-February-01  
**Number of tests:** 6  
**Number of wells per test:** 15

#### Well specifications:

Well No.	Production No.	Well No.	Production No.
1	2009-642-01	9	2011-825-09
2	2008-531-02	10	2009-642-10
3	2011-825-03	11	2011-825-11
4	2009-642-04	12	2009-642-12
5	2008-531-05	13	2011-825-13
6	2008-531-06	14	2011-825-14
7	2008-531-07	15	2012-050-15
8	2008-531-08		

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 11 to 13 and 15 were available. The specificities of the primers in primer solution 11 were tested by separately adding one additional 5'-primer, respectively one additional 3'-primer. In primer solutions 12, 13 and 15 it was only possible to test the 5'-primer, the 3'-primers were not possible to test. One of the 3'-primers in primer solutions 5, 8, 9 and 11 was not possible to test. The specificities of additional primers in primer solution 5 and 8 were tested by separately adding one additional 5'-primer, and/or one additional 3'-primer

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

**Date of approval:** 2012-August-20

**Approved by:**



**Production Quality Control**

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## Declaration of Conformity

**Product name:** *Olerup* SSP® HLA-A\*66  
**Product number:** 101.427-06/06u  
**Lot number:** 09R

**Intended use:** HLA-A\*66 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  
2012-August-20

Ann-Cathrin Jareman  
Head of QA and Regulatory Affairs