

101.211.24 – including *Taq* pol., IFU-01 Rev. No. 03  
 101.211.24u – without *Taq* pol., IFU-02 Rev. No. 03

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

Lot No.: **83N**

Lot-specific information

[www.olerup-ssp.com](http://www.olerup-ssp.com)

## CERTIFICATE OF ANALYSIS

### Olerup SSP® DQB1\*05 SSP

**Product number:** 101.211-24 – including *Taq* polymerase  
 101.211-24u - without *Taq* polymerase

**Lot number:** 83N

**Expiry date:** 2014-December-01

**Number of tests:** 24

**Number of wells per test:** 12

#### Well specifications:

Well No.	Production No.	Well No.	Production No.
1	2009-696-01	9	2012-024-09
2	2011-846-02	10	2012-024-10
3	2009-696-03	11	2012-024-11
4	2009-696-04	12	2012-024-12
5	2011-846-05		
6	2012-024-06		
7	2011-846-07		
8	2011-846-08		

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 5 to 12 were available. The specificities of the primers in primer solutions 5, 6 and 8 were tested by separately adding one 5'-primer, respectively one 3'-primer. In primer solutions 7 and 12 it was only possible to test the 3'-primer, the 5'-primers were not possible to test. In primer solutions 9 to 11 it was only possible to test the 5'-primer, the 3'-primers were not possible to test. In primer solution 5 one 3'-primer was not possible to test,

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

**Date of approval:** 2012-June-29

**Approved by:**



**Production Quality Control**

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## Declaration of Conformity

**Product name:** *Olerup* SSP® DQB1\*05  
**Product number:** 101.211-24/24u  
**Lot number:** 83N


**Intended use:** DQB1\*05 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 III, 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.

Stockholm, Sweden  
2012-June-29



Ann-Cathrin Jareman  
Head of QA and Regulatory Affairs