DQB1*05 101.211-24 – including <i>Taq</i> polymera		Page 1 of 12 eneral "Instructions for Use" b. 02 can be downloaded from
Lot No.: 04M Lot	-specific information	www.olerup-ssp.com
Oleru	pSSP [®] DQB1*05	5
Product number:	101.211-24 – incl	uding <i>Taq</i> polymerase
Lot number:	04M	
Expiry date:	2013-October-01	
Number of tests:	24	
Number of wells per test:	8	
Storage - pre-aliquoted primers	s: dark at -20°C	
- PCR Master Mix:	-20°C	
- Adhesive PCR seals	RT	
- Product Insert	RT	

This Product Description is only valid for Lot No. 04M

CHANGES COMPARED TO THE PREVIOUS OLERUP SSP® DQB1*05 LOT

The DQB1*05 specificity and interpretation tables have been updated for the HLA-DQB1 alleles described since the previous *Olerup* SSP[®] DQB1*05 lot was made **(Lot No. 83G)**.

Two wells has been added to the DQB1*05 kit,
wells 7 and 8 .

The primers of the wells detailed below have been exchanged, added or modified compared to the previous lot.

Well	5'-primer	3'-primer	rationale
5	Added	Added	Primer pair added for the DQB1*05:10 allele.
6	Added, modified	Added	Primer pair added for the DQB1*05:08 allele, increased yield of HLA-specific PCR product.
7	New	New	New primer pairs for the DQB1*05:05 and DQB1*05:06 alleles.
8	New	New	New primer pair for the DQB1*05:09 allele.

Lot-specific information

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PRODUCT DESCRIPTION

DQB1*05 SSP subtyping

CONTENT

The primer set contains 5'- and 3'-primers for identifying the DQB1*05:01 to DQB1*05:11 alleles.

Please note that DQB1 amplifications usually are somewhat less pronounced than e.g. DRB and DQA1 amplifications even when using the same DNA preparation and exactly the same experimental procedures.

PLATE LAYOUT

Each test consists of 8 PCR reactions in an 8 well PCR plate.

1	2	3	4	5	6	7	8
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The 8 well cut PCR plate is marked with 'DQ5' in silver gray ink.

Well No. 1 is marked with the Lot No. '04M'.

A faint row of numbers is seen between wells 1 and 2 or wells 7 and 8 of the PCR trays. These stem from the manufacture of the trays, and should be disregarded.

The PCR plates are covered with a PCR-compatible foil.

Please note: When removing each 8 well PCR plate, make sure that the remaining plates stay covered. Use a scalpel or a similar instrument to carefully cut the foil between the plates.

INTERPRETATION

Only DQB1*05 alleles will be amplified by the DQB1*05 subtyping kit, except that primer mix 6 will amplify the DQB1*03:01:04 allele. Thus, the interpretation of DQB1*05 SSP subtypings is only influenced by this allele and not by other groups of DQB1 alleles or the DQB2 and DQB3 genes.

UNIQUELY IDENTIFIED ALLELES

All the DQB1*05 alleles, i.e. **DQB1*05:01 to DQB1*05:11**, recognized by the HLA Nomenclature Committee in January 2011¹ will give rise to unique amplification patterns by the primers in the DQB1*05 subtyping kit.

The DQB1*05 subtyping kit cannot distinguish the DQB1*05:01:01-05:01:03, the DQB1*05:02:01 and 05:02:03 alleles or the DQB1*05:03:01-05:03:04 alleles.

¹DQB1 alleles listed on the IMGT/HLA web page 2011-January-14, release 3.3.0, <u>www.ebi.ac.uk/imgt/hla</u>.

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RESOLUTION IN HOMO- AND HETEROZYGOTES

A total of 21 alleles generate 10 amplification patterns that can be combined in 55 homozygous and heterozygous combinations. 27 of these genotypes do not give rise to unique amplification patterns.

+++--+--*05:02:01, *05:11 = *05:05, *05:11 ++-+-++-*05:06, *05:11 = *05:07, *05:08 +-+++--- *05:02:01, *05:10 = *05:02:02, *05:04 = *05:02:02, *05:10 +-++-+-- *05:02:01, *05:08 = *05:02:02, *05:05 = *05:02:02, *05:08 = *05:05, *05:08 +-++--+-*05:02:01, *05:06 = *05:02:02, *05:06 +-++---+ *****05:02:01, *****05:09 = *****05:02:02, *****05:09 +-++----*05:02:01, *05:02:02 = *05:02:02, *05:02:02 +-+--+--*05:02:01, *05:05 = *05:05, *05:05 +--+++-- *05:04, *05:08 = *05:08, *05:10 +--++-+- *05:04, *05:06 = *05:06, *05:10 +--++--+ *05:04, *05:09 = *05:09, *05:10 +--++--- *05:04, *05:10 = *05:10, *05:10

*05:01:01 = *05:01:01-05:01:03

- *05:02:01 = *05:02:01 and 05:02:03
- *05:03:01 = *05:03:01-05:03:04

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SPECIFICITY TABLE

DQB1*05 SSP subtyping

Specificities and sizes of the PCR products of the 8 primer mixes used for DQB1*05 SSP subtyping

Primer Mix	Size of spec. PCR product ¹	Size of control band ²	Amplified DQB1*05 alleles ³	Other amplified DQB1 alleles ⁴
1	225 bp	515 bp	*05:01:01-05:11	
2	135 bp	430 bp	*05:01:01-05:01:03, 05:07, 05:11	
3 ⁵	120 bp	430 bp	*05:02:01-05:02:03, 05:05	
4 ⁵	95 bp	515 bp	*05:02:02, 05:03:01-05:03:04, 05:06, 05:08-05:10	
5 ^{5,7}	120 bp, 185 bp	430 bp	*05:04, 05:10	
6 ^{6,8}	135 bp, 185 bp	430 bp	*05:05, 05:08, 05:11	*03:01:04
7 ⁶	180 bp	430 bp	*05:06-05:07	
8	190 bp	430 bp	*05:09	

¹Alleles are assigned by the presence of specific PCR product(s). However, the sizes of the specific PCR products may be helpful in the interpretation of DQB1*05 SSP typings.

When the primers in a primer mix can give rise to specific PCR products of more than one length this is indicated if the size difference is 20 base pairs or more. Size differences shorter than 20 base pairs are not given. For high resolution SSP kits the respective lengths of the specific PCR product(s) of the alleles amplified by these primer mixes are given.

Nonspecific amplifications, i.e. a ladder or a smear of bands, may sometimes be seen. GC-rich primers have a higher tendency of giving rise to nonspecific amplifications than other primers.

PCR fragments longer than the control bands may sometimes be observed. Such bands should be disregarded and do not influence the interpretation of the SSP typings.

PCR fragments migrating faster than the control bands, but slower than a 400 bp fragment may be seen in some gel read-outs. Such bands can be disregarded and do not influence the interpretation of the SSP typings.

Some primers may give rise to primer oligomer artifacts. Sometimes this phenomenon is an inherit feature of the primer pair(s) of a primer mix. More often it is due to other factors such as too low amount of DNA in the PCR reactions, taking too long time in setting up the PCR reactions, working at elevated room temperature or using thermal cyclers that are not pre-heated.

²The internal positive control primer pairs amplify segments of the human growth hormone gene. The two different control primer pairs give rise to either an internal positive control band of 430 base pairs, for most wells, or a band of 515 base pairs, for some wells.

Well number 1 contains the primer pair giving rise to the longer, 515 bp, internal positive control band in order to help in the correct orientation of the DQB1*05 subtyping.

In addition, well number 4 contains the primer pair giving rise to the longer, 515 bp, internal positive control band in order to allow kit identification.

In the presence of a specific amplification the intensity of the control band often decreases.

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³For several DQB alleles only partial second exon nucleotide sequences are available. In these instances it is not known whether some of the primers of the SSP sets are completely matched with the target sequences or not. We assume that unknown sequences of codons 87 to 92 are conserved within allelic groups.

⁴Due to the sharing of sequence motifs, the DQB1*03:01:04 allele is amplified by primer mix 6. ⁵Specific PCR fragments shorter than 125 base pairs have a lower intensity than longer PCR bands.

⁶Primer mixes 6 and 7 may give rise to nonspecific amplifications.

⁷Primer mix 5: Specific PCR fragment of 120 bp in the DQB1*05:04 allele. Specific PCR fragment of 185 bp in the DQB1*05:10 allele.

⁸Primer mix 6: Specific PCR fragment of 135 bp in the DQB1*05:08 and the DQB1*03:01:04 alleles. Specific PCR fragment of 185 bp in the DQB1*05:05 and 05:11 alleles.

DQB1*05	Product Insert
101.211-24 - including	<i>Taq</i> polymerase

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		INTEF	RPRET	ATION ⁻	TABLE			
		DQE	31*05 SS	SP subt	yping			
Am	plification					1*05:11 al	leles	
	Well ⁴							
	1	2	3	4	5	6	7	8
Length of spec.	225	135	120	95	120	135	180	190
PCR product(s)					185	185		
Length of int.	515	430	430	515	430	430	430	430
pos. control ¹								
5'-primer ²	26(173)	26(173)	30(184)	30(184)	30(184)	38(209)	39(212)	37(205)
-	^{5'} -ggg ^{3'}	^{5'} -ggg ^{3'}	^{5'} -gAC ^{3'}	^{5'} -gAC ^{3'}	^{5'} -gAT ^{3'}	^{5'} -CgC ^{3'}	^{5'} -gCA ^{3'}	^{5'} -Agg ^{3'}
					135(500)	135(500)	40(216)	
					^{5'} -TgA ^{3'}	^{5'} -TgA ^{3'}	^{5'} -TTg ^{3'}	
3'-primer ³	87(356)	57(266)	57(265)	47(237)	57(265)	86(353)	87(356)	86(353)
	^{5'} -ggT ^{3'}	^{5'} -CAA ^{3'}	^{5'} -gCT ^{3'}	^{5'} -CgA ^{3'}	^{5′} -gCT ^{3′}	^{5'} -ACg ^{3'}	^{5'} -ggT ^{3'}	^{5'} -ACg ^{3'}
					182(642)	167(596)		
					^{5'} -ggT ^{3'}	^{5'} -CAT ^{3'}		
Well No.	1	2	3	4	5	6	7	8
DQB1 allele								
*05:01:01-	1	2						
05:01:03	•	-						
*05:02:01,	1		3					
05:02:03	-		•					
*05:02:02	1		3	4				
*05:03:01-	1			4				
05:03:04	-							
*05:04	1				5			
*05:05	1		3	-		6		
*05:06	1	-		4			7	
*05:07	1	2					7	
*05:08	1			4		6		•
*05:09	1			4				8
*05:10	1	•		4	5	•		
*05:11	1	2				6		
*03:01:04						6		
DQB1 allele	4	•			_	•		
Well No.	1	2	3	4	5	6	7	8

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Lot-specific information

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¹The internal positive control primer pairs amplify segments of the human growth hormone gene. The two different control primer pairs give rise to either an internal positive control band of 430 base pairs, for most wells, or a band of 515 base pairs, for some wells.

Well number 1 contains the primer pair giving rise to the longer, 515 bp, internal positive control band in order to help in the correct orientation of the DQB1*05 subtyping.

In addition, well number 4 contains the primer pair giving rise to the longer, 515 bp, internal positive control band in order to allow kit identification.

²The codon, and in parenthesis the nucleotide, in the 2nd and 3rd exons, matching the specificitydetermining 3'-end of the primer is given. Codon and nucleotide numbering as on the <u>www.ebi.ac.uk/imgt/hla</u> web site. The sequence of the 3 terminal nucleotides of the primer is given.

³The codon, and in parenthesis the nucleotide, in the 2nd or 3rd exons, matching the specificitydetermining 3'-end of the primer is given in the anti-sense direction. Codon and nucleotide numbering as on the <u>www.ebi.ac.uk/imgt/hla</u> web site. The sequence of the 3 terminal nucleotides of the primer is given.

⁴Primer mix 5: Specific PCR fragment of 120 bp in the DQB1*05:04 allele. Specific PCR fragment of 185 bp in the DQB1*05:10 allele.

Primer mix 6: Specific PCR fragment of 135 bp in the DQB1*05:08 and the DQB1*03:01:04 alleles. Specific PCR fragment of 185 bp in the DQB1*05:05 and 05:11 alleles.

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7 9020 QBL *02:01 - <th< td=""><td>-</td><td></td><td>-</td></th<>	-		-
8 9025 DEU *03:01 - <th< td=""><td>1-</td><td>-</td><td>-</td></th<>	1-	-	-
9 9026 YAR *03:02 - <th< td=""><td></td><td>- 1</td><td>-</td></th<>		- 1	-
10 9107 LKT3 *04:01	- 1	-	-
	- 1	-	-
	-	-	-
12 9052 DBB *03:03	-	-	-
12 3032 DBD 00.03	-	+ -	-
13 9004 JESTHOM 05:01 + + +	<u> </u> _	-	-
15 9075 DKB *03:03	-	-	-
10 9073 DKB 03.03 - <	+-	-	-
		-	-
	-	-	-
		-	-
19 9038 BM16 *03:01	-	-	-
20 9059 SLE005 *06:04	-	-	-
21 9064 AMALA *03:01		-	-
22 9056 KOSE *05:03 *06:04 + + -	-	-	-
23 9124 IHL *05:03 *06:01 + + -	-	-	-
24 9035 JBUSH *03:01	-	-	-
25 9049 IBW9 *02:02	-	-	-
26 9285 WT49 *02:01	-	-	-
27 9191 CH1007 *04:01 *05:01 + +	-	-	-
28 9320 BEL5GB *02:02 *03:01	-	-	-
29 9050 MOU *02:02	-	-	-
30 9021 RSH *04:02	-	-	-
31 9019 DUCAF *02:01	-	-	-
32 9297 HAG *03:01	-	-	-
33 9098 MT14B *03:02	-	-	-
34 9104 DHIF *03:01	-	-	-
35 9302 SSTO *03:05	-	-	-
36 9024 KT17 *03:02	-	-	-
37 9065 HHKB *06:03	-	-	-
38 9099 LZL *03:01	-	-	-
39 9315 CML *02:01 *03:01	-	-	-
40 9134 WHONP199 *02:02 *03:03	-	-	-
41 9055 H0301 *06:09 -	-	-	-
42 9066 TAB089 *06:01	-	-	-
43 9076 T7526 *03:03	-	-	-
44 9057 TEM *05:03 + + -	-	-	-
45 9239 SHJO *02:02	-	-	-
46 9013 SCHU *06:02	-	-	-
47 9045 TUBO *03:01	-	-	-
48 9303 TER-ND *05:01 + +	-	-	-

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Lot No.: **04M**

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CERTIFICATE OF ANALYSIS

Olerup SSP[®] DQB1*05 SSP

Product number:101.211-24 – including Taq polymeraseLot number:04MExpiry date:2013-October-01Number of tests:24Number of wells per test:8

Well specifications:

Well No.	Production No.
1	2009-696-01
2	2011-846-02
3	2009-696-03
4	2009-696-04
5	2011-846-05
6	2011-846-06
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 8 were available. The specificities of the primers in primer solutions 5, 6 and 8 were tested by separately adding one additional 5'-primer, respectively one additional 3'-primer. In primer solution 7 it was only possible to test the 3'-primer, the 5'-primers were not possible to test. In primer solutions 5 and 6 one 3'-primer was not possible to test,

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

Date of approval: 2011-May-26

Approved by:

Quality Control, Supervisor

Lot-specific information

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Declaration of Conformity

Product name: Product number: Lot number:	<i>Olerup</i> SSP [®] DQB1*05 101.211-24 04M
Intended use:	DQB1*05 resolution histocompatibility testing
Manufacturer:	<i>Olerup</i> SSP AB Franzengatan 5 SE-112 51 Stockholm, Sweden <i>Phone:</i> +46-8-717 88 27 <i>Fax:</i> +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.

The Authorized Representative located within the Community is: *Olerup* SSP AB.

Stockholm, Sweden 2011-May-26

Olle Olerup Managing Director

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Lot No.: 04M	Lot-specific information	www.olerup-ssp.com

Lot-specific information

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ADDRESSES:

Manufacturer: Olerup SSP AB, Franzengatan 5, SE-112 51 Stockholm, Sweden. *Tel:* +46-8-717 88 27 *Fax:* +46-8-717 88 18 *E-mail:* info-ssp@olerup.com *Web page:* http://www.olerup-ssp.com

Distributed by: Olerup GmbH, Löwengasse 47 / 6, AT-1030 Vienna, Austria. *Tel:* +43-1-710 15 00 *Fax:* +43-1-710 15 00 10 *E-mail:* support-at@olerup.com *Web page:* http://www.olerup.com

Olerup Inc., 901 S. Bolmar St., Suite R, West Chester, PA 19382 *Tel:* 1-877-OLERUP1 *Fax:* 610-344-7989 *E-mail:* info.us@olerup.com *Web page:* http://www.olerup.com

For information on Olerup SSP distributors worldwide, contact Olerup GmbH.