

101.427-06 – including *Taq* pol., IFU-01  
101.427-06u – without *Taq* polymerase, IFU-02

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“Instructions for Use” (IFU)

Lot No.: **21V**

Lot-specific information  
**Olerup SSP® HLA-A\*66**

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Product number:	101.427-06 – including <i>Taq</i> pol. 101.427-06u – without <i>Taq</i> pol.
Lot number:	21V
Expiry date:	2016-July-01
Number of tests:	6
Number of wells per test:	15+1
Storage - pre-aliquoted primers:	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. 21V.

Complete product documentation consists of generic Instructions for Use (IFU), lot specific Product Insert, Worksheet and Certificate.

### CHANGES COMPARED TO THE PREVIOUS OLERUP SSP® HLA-A\*66 LOT (09R)

The HLA-A\*66 kit is updated for new alleles to enable separation of:

- Confirmed<sup>1</sup> alleles as listed in the IMGT/HLA database
- Polymorphisms in exons outside of the region encoding the peptide binding domain
- Null and Alternatively expressed alleles

A well containing Negative Control primer pairs has been added.

The format of the Product Insert and Worksheet have been changed.

One well has been added to the HLA-A\*66 kit, well **16**.

<sup>1</sup>As described in section Uniquely Identified Alleles.

The HLA-A\*66 specificity and interpretation tables have been updated for the HLA-A alleles described since the previous *Olerup SSP®* HLA-A\*66 lot was made (**Lot No. 09R**).

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As of lot series V, the Specificity Table is included in the lot-specific Product Insert, and the Interpretation Table is included in the Worksheet.

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

Well	5'-primer	3'-primer	rationale
12	-	Added	3'-primer added for the A*66:17 allele
13	Added	Added	Primer pair added for the A*66:19 allele.
15	-	Added	3'-primer added for the A*66:18 allele.
16	New	New	Negative Control

Change in revision R01 compared to R00:

1. Primer mix 6 does not amplify the A\*66:15 and the A\*11:98 and 31:03 alleles. This has been corrected in the Specificity and Interpretation tables.

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Well **16** contains Negative Control primer pairs, that will amplify more than 95% of the *Olerup SSP®* HLA Class I, DRB, DQB1 and DPB1 amplicons as well as amplicons generated by a control primer pair.

PCR product sizes range from 75 to 430 base pairs.  
The PCR product generated by the control primer pair is 430 base pairs.

Length of PCR product	105	200	105	80	75	80
<b>5'-primer<sup>1</sup></b>	<b>164</b>	<b>340</b>	<b>440</b>	<b>45</b>	<b>45</b>	<b>43</b>
	5'-CAC <sup>3'</sup>	5'-Agg <sup>3'</sup>	5'-TTA <sup>3'</sup>	5'-Tgg <sup>3'</sup>	5'-Tgg <sup>3'</sup>	5'-Tgg <sup>3'</sup>
<b>3'-primer<sup>2</sup></b>	<b>231</b>	<b>2<sup>nd</sup> I</b>	<b>507</b>	<b>59</b>	<b>58</b>	<b>57</b>
	5'-TgC <sup>3'</sup>	5'-AAA <sup>3'</sup>	5'-TTg <sup>3'</sup>	5'-CTC <sup>3'</sup>	5'-ggC <sup>3'</sup>	5'-CTC <sup>3'</sup>
<b>A*</b>	<b>+</b>	<b>+</b>	<b>+</b>			
<b>B*</b>	<b>+</b>	<b>+</b>	<b>+</b>			
<b>C*</b>	<b>+</b>	<b>+</b>	<b>+</b>			
<b>DRB1</b>				<b>+</b>	<b>+</b>	
<b>DRB3</b>				<b>+</b>	<b>+</b>	
<b>DRB5</b>				<b>+</b>		
<b>DQB1</b>					<b>+</b>	
<b>DPB1</b>						<b>+</b>

<sup>1</sup>The nucleotide position for HLA class I genes and the codon for HLA class II genes, in the 2<sup>nd</sup> or 3<sup>rd</sup> exon, matching the specificity-determining 3'-end of the primer is given. Nucleotide and codon numbering as on the [www.ebi.ac.uk/imgt/hla](http://www.ebi.ac.uk/imgt/hla) web site. The sequence of the 3 terminal nucleotides of the primer is given.

<sup>2</sup>The nucleotide position for HLA class I genes and the codon for HLA class II genes, in the 2<sup>nd</sup> or 3<sup>rd</sup> exon or the 2<sup>nd</sup> intron, matching the specificity-determining 3'-end of the primer is given in the anti-sense direction. Nucleotide and codon numbering as on the [www.ebi.ac.uk/imgt/hla](http://www.ebi.ac.uk/imgt/hla) web site. The sequence of the 3 terminal nucleotides of the primer is given.

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## PRODUCT DESCRIPTION

### HLA-A\*66 SSP subtyping

#### CONTENT

The primer set contains 5'- and 3'-primers for identifying the HLA-A\*66:01 to A\*66:19 alleles.

#### PLATE LAYOUT

Each test consists of 16 PCR reactions in a 16 well cut PCR plate.

<b>1</b>	<b>2</b>	<b>3</b>	<b>4</b>	<b>5</b>	<b>6</b>	<b>7</b>	<b>8</b>
<b>9</b>	<b>10</b>	<b>11</b>	<b>12</b>	<b>13</b>	<b>14</b>	<b>15</b>	<b>NC</b>

The 16 well cut PCR plate is marked with 'HLA-A\*66' in silver/gray ink.

Well No. 1 is marked with the Lot No. '21V'.

Wells 1 to 15 – HLA-A\*66 high resolution primers.

Well 16 – Negative Control (NC).

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 heat-sealed with a PCR-compatible foil.

**Please note:** When removing each 16 well PCR plate, make sure that the remaining plates stay sealed. Use a scalpel or a similar instrument to carefully cut the foil between the plates.

#### INTERPRETATION

Due to the sharing of sequence motifs between HLA-A alleles, non-HLA-A\*66 alleles will be amplified by primer mixes 1 to 3 and 5 to 15. In addition, one HLA-C allele will be amplified by primer mix 2.

For further details see Specificity Table.

#### UNIQUELY IDENTIFIED ALLELES

All the HLA-A\*66 alleles, i.e. **A\*66:01 to A\*66:19 alleles**, recognized by the HLA Nomenclature Committee in October 2013<sup>1,2,3</sup> will give rise to unique amplification patterns by the primers in the HLA-A\*66 subtyping kit.

The HLA-A\*66 kit enables separation of the confirmed HLA-A\*66 alleles as listed in the IMGT/HLA database. An HLA allele is listed as confirmed by IMGT/HLA if it has been sequenced by more than a single laboratory or from multiple sources. Current allele confirmation status for HLA-A\*66 alleles is listed below.

The HLA-A\*66 kit also enables identification of polymorphisms in exons outside of the region encoding the peptide binding domain and of null and alternatively expressed alleles

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The following HLA-A\*66 alleles can be distinguished by the different sizes of the HLA-specific PCR product:

Alleles	Primer mix
A*66:08, 66:17	12
A*66:13, 66:19	13

<sup>1</sup>HLA-A alleles listed on the IMGT/HLA web page 2013-October-11, release 3.14.0, [www.ebi.ac.uk/imgt/hla](http://www.ebi.ac.uk/imgt/hla).

<sup>2</sup>Alleles that have been deleted from or renamed in the official WHO HLA Nomenclature up to and including the last IMGT/HLA database release can be retrieved from web page <http://hla.alleles.org/alleles/deleted.html>.

<sup>3</sup>The A\*66:05 and A\*26:92 alleles give rise to identical amplification patterns with the HLA-A\*66 primer set. These two alleles can be distinguished by the HLA-A low resolution and/or HLA-A\*26 kits.

## ALLELE CONFIRMATION STATUS

Allele	Status <sup>1</sup>	Allele	Status <sup>1</sup>
<b>A*66:01:01</b>	<b>Confirmed</b>	<b>A*66:11</b>	<b>Confirmed</b>
<b>A*66:02</b>	<b>Confirmed</b>	<b>A*66:12</b>	<b>Confirmed</b>
<b>A*66:03</b>	<b>Confirmed</b>	<b>A*66:13</b>	<b>Confirmed</b>
A*66:04	Unconfirmed	A*66:14	Unconfirmed
A*66:05	Unconfirmed	A*66:15	Unconfirmed
A*66:06	Unconfirmed	A*66:16	Unconfirmed
A*66:07	Unconfirmed	<b>A*66:17</b>	<b>Confirmed</b>
A*66:08	Unconfirmed	A*66:18	Unconfirmed
<b>A*66:09</b>	<b>Confirmed</b>	A*66:19	Unconfirmed
A*66:10	Unconfirmed		

<sup>1</sup>Allele status “confirmed” or “unconfirmed” as listed on the IMGT/HLA web page 2013-October-11, release 3.14.0, [www.ebi.ac.uk/imgt/hla](http://www.ebi.ac.uk/imgt/hla).

## RESOLUTION IN HOMO- AND HETEROZYGOTES

Results file with resolution in HLA-A\*66 homo- and heterozygotes is available upon request.

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## SPECIFICITY TABLE

### HLA-A\*66 SSP subtyping

Specificities and sizes of the PCR products of the 15+1 primer mixes used for HLA-A\*66 SSP subtyping

Primer Mix	Size of spec. PCR product <sup>1</sup>	Size of control band <sup>2</sup>	Amplified HLA-A*66 alleles <sup>3</sup>	Other amplified HLA-A alleles <sup>4</sup>
1 <sup>6</sup>	175 bp	800 bp	*66:01:01, 66:04-66:15, 66:17-66:19	*01:01:56, 02:135, 25:01:01-25:05, 25:07-25:17, 25:19-25:23, 26:01:01-26:01:20, 26:01:22-26:03:02, 26:05-26:08, 26:10-26:33, 26:35-26:43:02, 26:45-26:72, 26:74-26:77, 26:79-26:90, 26:92-26:94, 43:01
2 <sup>5</sup>	100 bp	1070 bp	*66:01:01, 66:04, 66:06-66:11, 66:13-66:14, 66:17-66:19	*01:13, 01:17, 03:63, 03:88, 11:01:01-11:11, 11:13-11:16, 11:20-11:27, 11:29-11:39, 11:41-11:52Q, 11:54-11:95, 11:97, 11:99N-11:105, 11:107-11:120, 11:122-11:158, 25:02, 26:13, 26:19, 26:33, 34:01:01-34:06, 34:08, 34:10N-34:11, 69:02, <b>C*07:335</b>
3	430 bp	1070 bp	*66:01:01-66:02, 66:04, 66:06-66:19	*02:11:01-02:11:05, 02:34-02:35:03, 02:56:01-02:56:02, 02:62, 02:69, 02:78, 02:103, 02:128, 02:297-02:298, 02:308, 24:19, 26:13, 26:19, 34:01:01-34:11, 68:01:01:01-68:02:05, 68:04, 68:06-68:14, 68:16-68:19, 68:21:01-68:30, 68:32-68:35, 68:37-68:56, 68:58-68:89, 68:91-68:106, 69:01-69:02
4	175 bp	1070 bp	*66:02-66:03, 66:16	
5 <sup>5</sup>	70 bp	800 bp		*02:55, 26:03:01-26:03:02, 26:06, 26:21, 26:30, 26:78, 33:24, 68:05, 68:15, 68:20
	100 bp		*66:04, 66:07	
6 <sup>5</sup>	80 bp	800 bp	*66:01:01, 66:04-66:10, 66:13-66:14, 66:17-66:19	*01:13, 01:28, 02:346, 02:427, 03:63, 03:88, 11:01:01-11:01:50, 11:01:52-11:11, 11:13-11:16, 11:19-11:27, 11:29-11:39, 11:41-11:44, 11:46-11:52Q, 11:54-11:97, 11:99-11:110, 11:112-11:158, 24:19, 24:44, 26:03:01-26:03:02, 26:06, 26:21, 26:78, 26:92, 34:01:01-34:08, 34:10N-34:11, 69:02, 80:02
7	560 bp	1070 bp	*66:03	*02:16, 02:131
8 <sup>5</sup>	95bp	1070 bp	*66:05, 66:07, 66:15	*01:01:01:01-01:01:27, 01:01:29-01:01:56, 01:01:58-01:04N, 01:06, 01:08-01:12, 01:14-01:16N, 01:18N-01:33, 01:35-01:70, 01:72-01:99, 01:101-01:104, 01:106-01:142, 02:346, 02:427, 03:41, 11:17, 11:19, 11:40, 11:98, 11:121, 24:44 <sup>w</sup> , 24:109 <sup>w</sup> , 25:01:01-25:01:07, 25:03-25:12N, 25:14, 25:16, 25:18-25:23, 26:01:01-26:01:21, 26:01:23-26:01:27, 26:01:29-26:12, 26:14-26:18, 26:20-26:29, 26:31-26:32, 26:34-26:43:02, 26:45-26:63, 26:66-26:82, 26:84-26:94, 31:03, 33:13, 36:01-36:05, 43:01, 74:10, 80:01:01:01-80:03

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9	155 bp	1070 bp	*66:06	*02:309, 02:454, 03:01:19, 25:19, 26:43:01-26:43:02, 31:03-31:04, 34:02:01-34:04, 34:06-34:09	
	235 bp		*66:14		
10	205 bp	1070 bp	*66:09	*02:03:01-02:03:06, 02:25, 02:38, 02:117, 02:148, 02:171:01-02:171:02, 02:230, 02:253, 02:258, 02:264, 02:267, 02:280-02:281, 02:315, 02:345, 02:355, 02:370, 02:412, 02:427, 02:431, 02:447, 26:22	
	190 bp	1070 bp	*66:10	*01:01:56, 26:29, 26:49	
11	235 bp		*66:14		
	95 bp	1070 bp	*66:17	*02:453	
12 <sup>5</sup>	220 bp		*66:08	*02:294, 32:54, 34:05 <sup>2</sup>	
	155 bp	1070 bp	*66:19	*01:01:56	
13	305 bp		*66:13		
	440 bp		*66:11		
14	360 bp	1070 bp	*66:02-66:03, 66:12, 66:16	*02:135, 02:309, 02:454, 03:01:19, 25:13, 26:30, 26:65, 31:04, 34:09	
15	140 bp	1070 bp	*66:16, 66:18	*25:08, 26:47	
16 <sup>7</sup>	<b>Negative Control</b>				

<sup>1</sup>Alleles 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 HLA-A\*66 SSP typings.

When the primers in a primer mix can give rise to HLA-specific PCR products of more than one length this is indicated if the size difference is more than 20 base pairs. Size differences of 20 base pairs or less are not given. For high resolution SSP kits, the alleles listed are specified according to amplicon length.

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 inherent 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.

<sup>2</sup>The internal positive control primer pairs amplify segments of the human growth hormone gene. The internal positive control bands are 1070 or 800 base pairs respectively, well distribution as outlined in the table. Well number 1 contains the shorter, 800 bp, internal positive control band. The well distribution of the internal controls can help in orientation of the kit on gel photo, as well as allow for kit identification. In the presence of a specific amplification the intensity of the control band often decreases.

<sup>3</sup>For several HLA Class I alleles 1<sup>st</sup> and/or 4<sup>th</sup> exon(s) and beyond, as well as intron nucleotide sequences, are not 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. Assumption is made that unknown sequences in these regions are conserved within allelic groups.

<sup>4</sup>Due to the sharing of sequence motifs between HLA-A alleles, non-HLA-A\*66 alleles will be amplified by primer mixes 1 to 3 and 5 to 15. In addition, one HLA-C allele will be amplified by primer mix 2.

<sup>5</sup>HLA-specific PCR products shorter than 125 base pairs have a lower intensity and are less sharp than longer PCR products.

<sup>6</sup>Primer mix 1 may give rise to a PCR fragment approx. 500 bp in size. This band should be disregarded in the interpretation of HLA-A\*66 subtypings.

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<sup>7</sup>Primer mix 16 contains a negative control, which will amplify more than 95% of HLA amplicons as well as the amplicons generated by control primer pairs. PCR product sizes range from 75 to 200 base pairs. The PCR product generated by the control primer pair is 430 base pairs.

‘?’ , nucleotide sequence of the primer matching sequence is not known.  
‘w’ , might be weakly amplified.



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**PRIMER SPECIFICATION**

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Well No.	1	2	3	4	5	6	7	8	9	10	11	12
Length of spec. PCR product	175	100	430	175	70	80	560	95	155	205	190	95
					100				235		235	220
Length of int. pos. control <sup>1</sup>	<b>800</b>	1070	1070	1070	<b>800</b>	<b>800</b>	1070	1070	1070	1070	1070	1070
5'-primer(s) <sup>2</sup>	418	282	28	423	261	302	282	282	423	355	423	652
	5'-AgA 3'	5'-CAg 3'	5'-TCg 3'	5'-gCT 3'	5'-AAC 3'	5'-ggA 3'	5'-CAC 3'	5'-CAC 3'	5'-gCT 3'	5'-CCg 3'	5'-gCT 3'	5'-CTg 3'
	423				517			517				
	5'-gCT 3'				5'-AgA 3'			5'-AgA 3'				
3'-primer(s) <sup>3</sup>	559	341	282	559	292	341	559	341	539	517	570	704
	5'-CCg 3'	5'-CgT 3'	5'-gAC 3'	5'-CTC 3'	5'-gTg 3'	5'-CgT 3'	5'-CTC 3'	5'-CgT 3'	5'-TCA 3'	5'-CgT 3'	5'-CCg 3'	5'-CCA 3'
	559		290		566			566	616		616	829
	5'-CCg 3'		5'-CAA 3'		5'-CCg 3'			5'-CCg 3'	5'-CgC 3'		5'-CgC 3'	5'-CTC 3'
					583							
					5'-gTg 3'							
Well No.	1	2	3	4	5	6	7	8	9	10	11	12

Well No.	13	14	15
Length of spec. PCR product	155	360	140
	305		
	440		
Length of int. pos. control <sup>1</sup>	1070	1070	1070
5'-primer(s) <sup>2</sup>	28	341	423
	5'-TCg 3'	5'-ggC 3'	5'-gCT 3'
	423		
	5'-gCT 3'		
3'-primer(s) <sup>3</sup>	164	418	518
	5'-gCA 3'	5'-gTC 3'	5'-CCA 3'
	299		521
	5'-CCg 3'		5'-ggA 3'
	538		
	5'-CCg 3'		
Well No.	13	14	15

<sup>1</sup>The internal positive control primer pairs amplify segments of the human growth hormone gene. The internal positive control bands are 1070 or 800 base pairs respectively, well distribution as outlined in the table. Well number 1 contains the shorter, 800 bp, internal positive control band. The well distribution of the internal controls can help in orientation of the kit on gel photo, as well as allow for kit identification. In the presence of a specific amplification the intensity of the control band often decreases.

<sup>2</sup>The nucleotide position matching the specificity-determining 3'-end of the primer is given. Nucleotide numbering as on the [www.ebi.ac.uk/imgt/hla](http://www.ebi.ac.uk/imgt/hla) web site. The sequence of the 3 terminal nucleotides of the primer is given.

<sup>3</sup>The nucleotide position matching the specificity-determining 3'-end of the primer is given in the anti-sense direction. Nucleotide numbering as on the [www.ebi.ac.uk/imgt/hla](http://www.ebi.ac.uk/imgt/hla) web site. The sequence of the 3 terminal nucleotides of the primer is given.

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CELL LINE VALIDATION SHEET																			
HLA-A*66 SSP subtyping kit <sup>2</sup>																			
				Lot No.:	Well														
					1	2	3	4	5	6	7	8	9	10	11	12	13	14	15
					200964201	200853102	201182503	200964204	200853105	200853106	200853107	200853108	201182509	201328810	201182511	201328812	201328813	201182514	201328815
	IHWC cell line <sup>1</sup>	A*	A*																
1	9001 SA	*24:02			-	-	-	-	-	-	-	-	-	-	-	-	-	-	-
2	9280 LK707	*02:01			-	-	-	-	-	-	-	-	-	-	-	-	-	-	-
3	9011 E4181324	*01:01			-	-	-	-	-	-	-	+	-	-	-	-	-	-	-
4	9275 GU373	*30:01			-	-	-	-	-	-	-	-	-	-	-	-	-	-	-
5	9009 KAS011	*01:01			-	-	-	-	-	-	-	+	-	-	-	-	-	-	-
6	9353 SM	*02:01	*26:03		+	-	-	-	+	+	-	+	-	-	-	-	-	-	-
7	9020 QBL	*26:01			+	-	-	-	-	-	-	+	-	-	-	-	-	-	-
8	9025 DEU	*31:01			-	-	-	-	-	-	-	-	-	-	-	-	-	-	-
9	9026 YAR	*26:01			+	-	-	-	-	-	-	+	-	-	-	-	-	-	-
10	9107 LKT3	*24:02			-	-	-	-	-	-	-	-	-	-	-	-	-	-	-
11	9051 PITOUT	*29:02			-	-	-	-	-	-	-	-	-	-	-	-	-	-	-
12	9052 DBB	*02:01			-	-	-	-	-	-	-	-	-	-	-	-	-	-	-
13	9004 JESTHOM	*02:01			-	-	-	-	-	-	-	-	-	-	-	-	-	-	-
14	9071 OLGA	*31:01			-	-	-	-	-	-	-	-	-	-	-	-	-	-	-
15	9075 DKB	*24:02			-	-	-	-	-	-	-	-	-	-	-	-	-	-	-
16	9037 SWEIG007	*29:02			-	-	-	-	-	-	-	-	-	-	-	-	-	-	-
17	9282 CTM3953540	*03:01	*80:01		-	-	-	-	-	-	-	+	-	-	-	-	-	-	-
18	9257 32367	*33:03	*74:01		-	-	-	-	-	-	-	-	-	-	-	-	-	-	-
19	9038 BM16	*02:01			-	-	-	-	-	-	-	-	-	-	-	-	-	-	-
20	9059 SLE005	*02:01			-	-	-	-	-	-	-	-	-	-	-	-	-	-	-
21	9064 AMALA	*02:17			-	-	-	-	-	-	-	-	-	-	-	-	-	-	-
22	9056 KOSE	*02:01			-	-	-	-	-	-	-	-	-	-	-	-	-	-	-
23	9124 IHL	*02:01	*34:01		-	+	+	-	-	+	-	-	-	-	-	+	-	-	-
24	9035 JBUSH	*32:01			-	-	-	-	-	-	-	-	-	-	-	-	-	-	-
25	9049 IBW9	*33:01			-	-	-	-	-	-	-	-	-	-	-	-	-	-	-
26	9285 WT49	*02:05			-	-	-	-	-	-	-	-	-	-	-	-	-	-	-
27	9191 CH1007	*24:10	*29:01		-	-	-	-	-	-	-	-	-	-	-	-	-	-	-
28	9320 BEL5GB	*02:01	*29:02		-	-	-	-	-	-	-	-	-	-	-	-	-	-	-
29	9050 MOU	*29:02			-	-	-	-	-	-	-	-	-	-	-	-	-	-	-
30	9021 RSH	*30:01	*68:02		-	-	+	-	-	-	-	-	-	-	-	-	-	-	-
31	9019 DUCAF	*30:02			-	-	-	-	-	-	-	-	-	-	-	-	-	-	-
32	9297 HAG	*02:01			-	-	-	-	-	-	-	-	-	-	-	-	-	-	-
33	9098 MT14B	*31:01			-	-	-	-	-	-	-	-	-	-	-	-	-	-	-
34	9104 DHIF	*31:01			-	-	-	-	-	-	-	-	-	-	-	-	-	-	-
35	9302 SSTO	*32:01			-	-	-	-	-	-	-	-	-	-	-	-	-	-	-
36	9024 KT17	*02:06	*11:01		-	+	-	-	-	+	-	-	-	-	-	-	-	-	-
37	9065 HHKB	*03:01			-	-	-	-	-	-	-	-	-	-	-	-	-	-	-
38	9099 LZL	*02:17			-	-	-	-	-	-	-	-	-	-	-	-	-	-	-
39	9315 CML	*01:01	*03:01		-	-	-	-	-	-	-	+	-	-	-	-	-	-	-
40	9134 WHONP199	*02:07	*30:01		-	-	-	-	-	-	-	-	-	-	-	-	-	-	-
41	9055 H0301	*03:01			-	-	-	-	-	-	-	-	-	-	-	-	-	-	-
42	9066 TAB089	*02:07			-	-	-	-	-	-	-	-	-	-	-	-	-	-	-
43	9076 T7526	*02:06	*02:07		-	-	-	-	-	-	-	-	-	-	-	-	-	-	-
44	9057 TEM	*66:01			+	+	+	-	-	+	-	-	-	-	-	-	-	-	-
45	9239 SHJO	*23:01	*24:02		-	-	-	-	-	-	-	-	-	-	-	-	-	-	-
46	9013 SCHU	*03:01			-	-	-	-	-	-	-	-	-	-	-	-	-	-	-
47	9045 TUBO	*02:16	*03:01		-	-	-	-	-	-	+	-	-	-	-	-	-	-	-
48	9303 TER-ND	*02:01	*11:01		-	+	-	-	-	+	-	-	-	-	-	-	-	-	-

<sup>1</sup>The provided cell line HLA specificities are retrieved from the <http://www.ihwg.org/hla> web site. The specificity of an individual cell line may thus be subject to change.

<sup>2</sup>The specificity of each primer solution in the kit has been tested against 48 well characterized cell line DNAs

**101.427-06 – including *Taq* pol., IFU-01**  
**101.427-06u – without *Taq* polymerase, IFU-02**

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

101.427-06 – including *Taq* pol., IFU-01  
101.427-06u – without *Taq* polymerase, IFU-02

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Lot No.: **21V**

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

**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](mailto: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](mailto: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](mailto:info.us@olerup.com)

**Web page:** <http://www.olerup.com>

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