102.101-01 – including *Taq* polymerase 102.101-01u – without *Taq* polymerase

Visit <u>www.caredx.com</u> for "Instructions for Use" (IFU)

Lot No.: 5N2 Lot-specific information Olerup SSP[®] HLA Wipe Test – Negative Control

| Product number: | 102.101-01 – including <i>Taq</i> polymerase |
|----------------------------------|--|
| Product number: | 102.101-01u – without Taq polymerase |
| Lot number: | 5N2 |
| Expiry date: | 2025-10-01 |
| Number of tests: | 96 |
| Number of wells per test: | 1-2 |
| Storage - pre-aliquoted primers: | dark at -20°C |
| - Positive Control DNA: | -20°C |
| - PCR Master Mix: | -20°C |
| - Adhesive PCR seals | RT |
| - Product Insert | RT |

This Product Description is only valid for Lot No. 5N2.

GENERAL DESCRIPTION

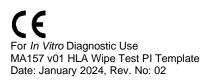
The *Olerup* SSP[®] HLA Wipe Test – Negative Control is intended to be used to monitor for contamination with amplicons generated by the *Olerup* SSP[®] and QTYPE product lines and can also be used as a negative control in *Olerup* SSP[®] typings.

The primer set contains Negative Control primer pairs that will amplify a majority of the *Olerup* SSP[®] HLA Class I, DRB, DQB1, DPB1 and DQA1 amplicons as well as amplicons generated by the control primer pairs matching the human growth hormone gene.

The Olerup SSP HLA Wipe Test – Negative Control has been established to be at least as sensitive as the Olerup SSP and QTYPE assays in detecting input DNA that could give rise to contamination issues when HLA typing with these products. The Olerup SSP[®] HLA Wipe Test – Negative Control has the sensitivity to detect approximately 25-50 copies of DNA template.

Changes in revision R01 compared to R00:

- 1. The size of the PCR product is corrected in the Protocol of the Product Insert.
- 2. The incubation temperature is corrected in step 6 of the Protocol of the Product Insert.



102.101-01 – including *Taq* polymerase 102.101-01u – without *Taq* polymerase

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Lot-specific information

PRODUCT DESCRIPTION

HLA Wipe Test – Negative Control

CONTENT

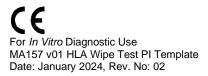
The primer set contains Negative Control primer pairs that will amplify the majority of the *Olerup* SSP[®] HLA Class I, DRB, DQB1, DPB1 and DQA1 amplicons as well as amplicons generated by the control primer pairs matching the human growth hormone gene for both the Olerup SSP and QTYPE product lines.

HLA-specific PCR product sizes range from 75 to 200 base pairs. The PCR product generated by the positive control primer pairs is 200 base pairs.

| Length of PCR | 105 | 200 | 105 | 80 | 75 | 80 | 85 |
|------------------------|----------------------------------|----------------------------------|----------------------------------|----------------------------------|----------------------------------|----------------------------------|----------------------------------|
| product | | | | | | | |
| 5'-primer ¹ | 164 | 340 | 440 | 45 | 45 | 43 | 36 |
| | ^{5'} -CAC ^{3'} | ^{5'} -Agg ^{3'} | ^{5'} -TTA3' | ^{5'} -Tgg ^{3'} | ^{5'} -Tgg ^{3'} | ^{5'} -Tgg ^{3'} | ^{5'} -TAC ^{3'} |
| | | | | | | | 36 |
| | | | | | | | ^{5'} -TAT ^{3'} |
| 3'-primer ² | 231 | 2 nd I | 507 | 59 | 58 | 57 | 47 |
| - | ^{5′} -TgC ^{3′} | ^{5'} -AAA ^{3'} | ^{5'} -TTg ^{3'} | ^{5'} -CTC ^{3'} | ^{5'} -ggC ^{3'} | ^{5′} -CTC ^{3′} | ^{5'} -ACA ^{3'} |
| | | | | | | | 48 |
| | | | | | | | ^{5'} -gCA ^{3'} |
| | | | | | | | 48 |
| | | | | | | | ^{5'} -gCC ^{3'} |
| | | | | | | | 52 |
| | | | | | | | ⁵'-TgT³' |
| A * | + | + | + | | | | |
| B* | + | + | + | | | | |
| C* | + | + | + | | | | |
| DRB1 | | | | + | + | | |
| DRB3 | | | | + | + | | |
| DRB5 | | | | + | | | |
| DQB1 | | | | | + | | |
| DPB1 | | | | | | + | |
| DQA1 | | | | | | | + |

¹The nucleotide position for HLA class I genes and the codon for HLA class II genes, in the 2nd or 3rd exon, matching the specificity-determining 3'-end of the primer is given. Nucleotide and codon 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.

²The nucleotide position for HLA class I genes and the codon for HLA class II genes, in the 2nd or 3rd exon or the 2nd intron, matching the specificity-determining 3'-end of the primer is given in the anti-sense direction. Nucleotide and codon 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.



102.101-01 – including *Taq* polymerase 102.101-01u – without *Taq* polymerase

Visit <u>www.caredx.com</u> for "Instructions for Use" (IFU)

Lot No.: 5N2

Lot-specific information

The primer solution is pre-aliquoted into 0.2 ml PCR wells. Each well contains the same dried primer solution.

PCR Master Mix complete with Taq, Taq polymerase, nucleotides, buffer, glycerol and cresol red, as well as adhesive PCR seals are included in the kit.

1-2 PCR reactions with a reaction volume of 10 μ l are performed per test.

Positive Control DNA is included in the kit, 20 ng/µl, 75 µl.

Sterile swabs, Dacron fiber tipped, plastic applicator are included in the kit. One per envelope, 100 per kit.

PCR Master Mix without Taq, nucleotides, buffer, glycerol and cresol red, as well as adhesive PCR seals are included in the kit.

1-2 PCR reactions with a reaction volume of 10 μ l are performed per test.

Positive Control DNA is included in the kit, 20 ng/µl, 75 µl.

Sterile swabs, Dacron fiber tipped, plastic applicator are included in the kit. One per envelope, 100 per kit.

PLATE LAYOUT

Each test consists of 1-2 PCR reactions. Each well of the 8 well PCR plates contains the same primer mix.

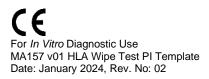
| | 1 | 1 | 1 | 1 | 1 | 1 | 1 | 1 |
|--|---|---|---|---|---|---|---|---|
|--|---|---|---|---|---|---|---|---|

The 8 well cut PCR plate is marked with 'WT' in silver/gray ink.

Well No. 1 is marked with the Lot No. '5N2'.

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 PCR well, make sure that the remaining plates/wells stay covered. Use a scalpel or a similar instrument to carefully cut the foil between the plates/wells.



102.101-01 – including *Taq* polymerase 102.101-01u – without *Taq* polymerase

Visit <u>www.caredx.com</u> for "Instructions for Use" (IFU)

Lot No.: 5N2

Lot-specific information **PROTOCOL**

It is recommended that 10 to 12 commonly used areas are tested for contamination; DNA preparation area, PCR setup area and post-amplification area. Test e.g. work benches, pipettes, centrifuges, refrigerator and freezer handles, door knobs and racks.

- 1. In a DNA free location label one 1.5 ml tube for each of the sample areas.
- 2. Add 500 µl sterile, distilled water to each tube.
- 3. Wet one of the provided sterile plastic applicator swab in each tube.
- 4. Wipe the area to be tested with the moistened applicator, and place it back in the original tube. Snap off the plastic stem of the applicator and close the cap of the tube.
- 5. Vortex briefly.
- 6. Incubate the samples at 50-60°C in a waterbath or heating block for 1 hour.
- 7. Centrifuge, 1 minute, 10 000 to 13 000 rpm in a microcentrifuge.
- 8. Remove and discard the applicator from the tubes with sterile forceps.

PCR AMPLIFICATION

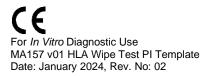
Positive control well.

Add 1 μ l of the provided Positive Control DNA and 1 μ l of dH₂O to well 1.

<u>Negative control well</u>. Add 2 μ l of dH₂O to well 2.

For each test area run two wells.

Add to the first well for each test area; 1 μ l of the test sample and 1 μ l of dH₂O. Add to the second well for each test area; 1 μ l of the test sample and 1 μ l of the Positive Control DNA. This well serves as an inhibition control.



■LERUP SSP

102.101-01 – including *Taq* polymerase 102.101-01u – without *Taq* polymerase

Visit <u>www.caredx.com</u> for "Instructions for Use" (IFU)

Lot No.: 5N2 Lot-specific information

102.101-01 – including Taq polymerase

In a 0.5 ml tube mix:

3 μ l of PCR Master Mix complete with *Taq* x the number of wells + 2 and

5 μ l of dH₂O x the number of wells + 2

e.g. for a Wipe Test consisting of (i) positive control well, (ii) negative control well and (iii) 10 tested areas (2 x 10 wells) mix: 24 x 3 = 72 μ l of PCR Master Mix complete with *Taq* and 24 x 5 = 120 μ l of dH₂O

Mix well, dispense 8 μ l of the PCR Master Mix complete with *Taq*-H₂O mixture into each of the wells of the HLA Wipe test. *Well No. 1 of the 8 well PCR plates is marked with the lot number.* Close the 8 well PCR plate(s) with the provided seals.

The HLA Wipe Test – Negative Control can also be used as a negative control when using the *Olerup* SSP[®] kits.

Cut off one well with a pair of scissors. Add to this well;

3 µl of PCR Master Mix complete with *Taq*

 $7 \mu l of dH_2O$

Include this well as a negative control when performing Olerup SSP® typings.

As of lot series V, the Negative Control is included in most Olerup SSP® typings.

102.101-01u – without Taq polymerase

In a 0.5 ml tube mix:

3 μ l of PCR Master Mix without *Taq* x the number of wells + 2, 0.1 μ l of Taq polymerase (5 units/ μ l) x the number of wells + 2 and 5 μ l of dH₂O x the number of wells + 2

e.g. for a Wipe Test consisting of (i) positive control well, (ii) negative control well and (iii) 10 tested areas (2 x 10 wells) mix: 24 x 3 = 72 μ l of PCR Master Mix without *Taq* and 24 x 0.1 = 2.4 μ l of *Taq* polymerase 24 x 5 - 2.4 = 117.6 μ l of dH₂O

Mix well, dispense 8 μ l of the PCR Master Mix without *Taq*-H₂O mixture into each of the wells of the HLA Wipe test. *Well No. 1 of the 8 well PCR plates is marked with the lot number.* Close the 8 well PCR plate(s) with the provided seals.

102.101-01 – including *Taq* polymerase 102.101-01u – without *Taq* polymerase

Visit <u>www.caredx.com</u> for "Instructions for Use" (IFU)

Lot No.: 5N2

5N2 Lot-specific information

The HLA Wipe Test – Negative Control can also be used as a negative control when using the *Olerup* $SSP^{\$}$ kits.

Cut off one well with a pair of scissors. Add to this well;

 $3 \ \mu l$ of PCR Master Mix without Taq

0.1 µl of Taq polymerase (5 units/µl)

6.9 μl of dH₂O

Include this well as a negative control when performing Olerup SSP® typings.

As of lot series V, the Negative Control is included in most Olerup SSP® typings.

Use a 96 well thermal cycler with a heated lid. The temperature gradient across the heating block should be $< 1^{\circ}$ C.

PCR cycling parameters:

| 1.1 cycle | 94°C | 2 min | denaturation |
|--------------|------|---------|-------------------------|
| 2. 10 cycles | 94ºC | 10 sec. | denaturation |
| | 65ºC | 60 sec. | annealing and extension |
| 3. 20 cycles | 94°C | 10 sec. | denaturation |
| | 61°C | 50 sec. | annealing |
| | 72°C | 30 sec. | extension |

The same PCR cycling parameters are used for all the Olerup SSP kits.

AGAROSE GEL ELECTROPHORESIS

Prepare a 2% (w/v) agarose gel in 0.5 x TBE buffer. Dissolve the agarose by boiling in a microwave oven. Let the gel solution cool to 60°C. Stain the gel prior to casting with ethidium bromide (10 mg/ml), 5 μ l per 100 ml gel solution. For maximal ease of handling use our ethidium bromide dropper bottles (Product No. 103.301-10), 1 drop of ethidium bromide solution per 50-75 ml of gel, or our GelRedTM dropper bottle (Product No. 103.302-05) 4 drops per 100-120 ml of gel solution. <u>Note:</u> Ethidium bromide is a powerful carcinogen.

Load the PCR products, preferably using an 8-channel pipette. Load a DNA size marker (100 base pair ladder, DNA Size Marker Product No. 103.202-100 or DNA Size Marker for short gel runs 103.203-100) in one well per row.

Run the gel in 0.5 x TBE buffer, without re-circulation of the buffer, for 15-20 minutes at 8-10 V/cm.

102.101-01 – including *Taq* polymerase 102.101-01u – without *Tag* polymerase

Visit <u>www.caredx.com</u> for "Instructions for Use" (IFU)

Lot No.: 5N2

Lot-specific information **DOCUMENTATION AND INTERPRETATION**

Put the gel on a UV transilluminator and document by photography. Record the presence and absence of PCR products.

HLA Wipe Test

In the positive control lane one or several PCR products should be seen. The sizes may range from 75 to 200 bp.

In the negative control lane no PCR product should be seen. Primer oligomer artefacts, approximately 40 to 50 bp in size, may be seen. This does not represent contamination.

The presence of PCR product(s) 75 to 200 bp in size in the sample lanes without Positive Control DNA indicates contamination.

The absence of PCR product in the sample lanes without Positive Control DNA shows that no detectable contamination is present.

In the sample lanes with the Positive Control DNA PCR product(s) of equal strength as in the positive control lane should be seen. If these PCR product(s) are weaker than the PCR product(s) in the positive control lane, then an inhibitor may be present in the sample. The test should then be repeated with the sample diluted 1:50 in sterile dH2O.

If contamination is detected, clean the area with a fresh 10% bleach solution and re-test the area.

Negative Control

In the negative control well no PCR product should be seen. The presence of PCR product(s) indicates contamination. Primer oligomer artefacts. approximately 40 to 50 bp in size, may be seen. This does not represent contamination.

If contamination is detected, wipe test and testing of all reagents should be performed in order to detect the source of contamination.

102.101-01 – including *Taq* polymerase 102.101-01u – without *Taq* polymerase

Visit <u>www.caredx.com</u> for "Instructions for Use" (IFU)

Lot No.: 5N2

Lot-specific information

ADDRESSES:

Manufacturer: CareDx AB, Franzengatan 5, SE-112 51 Stockholm, Sweden. Tel: +46-8-508 939 00 Fax: +46-8-717 88 18 E-mail: orders-se@caredx.com Web page: www.caredx.com/

CareDx Lab Solutions Inc., 901 S. Bolmar St., Suite R, West Chester, PA 19382 Tel: 1-877-653-78171 Fax: 610-344-7989 E-mail: orders-us@caredx.com Web page: www.caredx.com/

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