HLA-C low resolution

Certificates

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101.601-24/12 - including *Taq* polymerase 101.601-24u/12u - without Taq polymerase

Visit www.caredx.com for "Instructions for Use" (IFU)

Lot No.: 2V8

Lot-specific information

CERTIFICATE OF ANALYSIS

Olerup SSP® HLA-C low resolution

Product number:

101.601-24/12 – including *Taq* pol.

101.601-24u/12u - without Tag pol.

Lot number:

2V8

Expiry date:

2029-07-01

Number of tests:

24 tests - Product No. 101.601-24/24u 12 tests - Product No. 101.601-12/12u

31 + 1

Number of wells per test:

Well specifications:

won openioanene.			
Well No.	Production No.	Well No.	Production No.
_1	2025-637-01	17	2025-637-17
2	2025-637-02	18	2025-637-18
3	2025-637-03	19	2025-637-19
4	2025-637-04	20	2025-637-20
5	2025-637-05	21	2025-637-21
6	2025-637-06	22	2025-637-22
7	2025-637-07	23	2021-288-23
8	2025-637-08	24	2025-638-24
9	2025-637-09	25	2021-288-25
10	2025-637-10	26	2021-288-26
11	2025-637-11	27	2021-288-27
12	2025-637-12	28	2021-288-28
13	2025-637-13	29	2021-288-29
14	2025-637-14	30	2021-288-30
15	2021-288-15	31	2021-288-31
16	2025-637-16		

The negative control primer pairs, Production No. 2025-648-01, can detect contamination with PCR products diluted 10⁻⁷.

Results of Quality Control: No false positive or false negative amplifications

obtained.

Date of approval: 2025-08-20

Approved by:

Production Quality Control

For In Vitro Diagnostic Use MA099 v05 CoA_DoC General IVDs Date: August 2025, Rev. No: 00



HLA-C low resolution

Certificates

Page 2 of 2

101.601-24/12 – including *Taq* polymerase 101.601-24u/12u – without *Taq* polymerase

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

Lot No.: 2V8

Lot-specific information

Declaration of Conformity

Product name:

Olerup SSP® HLA-C low resolution

Product number:

101.601-24/24u, -12/12u

Lot number:

2V8

Intended use:

HLA-C low resolution histocompatibility testing

Manufacturer:

CareDx AB

Franzéngatan 5

SE-112 51 Stockholm, Sweden **Phone:** +46-8-508 939 00 **Fax:** +46-8-717 88 18

We, CareDx 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) EN ISO 13485:2016, 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 *CareDx* AB, Franzéngatan 5, SE-112 51 Stockholm, Sweden.

The Authorized Representative located within the Community is: CareDx AB.

Stockholm, Sweden Date:

2025-08-20 P.

Quality Assurance