Certificates

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101.211-24 - including Taq pol., IFU-01 101.211-24u - without Taq pol., IFU-02

Visit https://labproducts.caredx.com for "Instructions for Use" (IFU)

Lot No.: 0R1

Lot-specific information

## CERTIFICATE OF ANALYSIS

Olerup SSP® DQB1\*05

101.211-24 - including Taq polymerase Product number:

101.211-24u - without Tag polymerase

0R1 Lot number:

2026-03-01 **Expiry date:** 

Number of tests: 24 31+1 Number of wells per test:

## Well specifications:

Well No.	Production No.	Well No.	Production No.
1	2022-402-01	17	2021-278-17
2	2022-402-02	18	2021-278-18
3	2021-278-03	19	2021-278-19
4	2022-402-04	20	2021-278-20
5	2021-278-05	21	2021-278-21
6	2021-278-06	22	2021-278-22
7	2021-278-07	23	2021-278-23
8	2022-402-08	24	2022-402-24
9	2021-278-09	25	2021-278-25
10	2021-278-10	26	2021-278-26
11	2021-278-11	27	2021-278-27
12	2022-402-12	28	2021-278-28
13	2021-278-13	29	2021-278-29
14	2021-278-14	30	2021-278-30
15	2021-278-15	31	2022-402-31
16	2021-278-16		

The negative control primer pairs, Production No. 2021-336-01, can detect contamination with PCR products diluted 10<sup>-7</sup>.

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

obtained.

Date of approval: 2072-04-(1

Approved by:

**Production Quality Control** 

Certificates

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101.211-24 – including *Taq* pol., IFU-01 101.211-24u – without *Taq* pol., IFU-02 Visit <a href="https://labproducts.caredx.com">https://labproducts.caredx.com</a> for "Instructions for Use" (IFU)

Lot No.: 0R1

Lot-specific information

**Declaration of Conformity** 

Product name:

Olerup SSP® DQB1\*05

**Product number:** 

101.211-24/24u

Lot number:

0R1

Intended use:

DQB1\*05 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

2022-04-12

Date:

**Quality Assurance** 

CE