**DQB1\*02** 

Certificates

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101.213-24 – including *Taq* polymerase 101.213-24u – without *Taq* polymerase

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

Lot No.: 8R0

Lot-specific information

## **CERTIFICATE OF ANALYSIS**

Olerup SSP® DQB1\*02

Product number: 101.213-24 – including *Taq* polymerase

101.213-24u - without Tag polymerase

Lot number: 8R0

Expiry date: 2027-10-01

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

## Well specifications:

Well No.	Production No.	Well No.	Production No.
1	2020-231-01	17	2023-498-17
2	2020-231-02	18	2020-231-18
3	2021-358-03	19	2020-231-19
4	2020-231-04	20	2020-231-20
5	2020-231-05	21	2020-231-21
6	2020-231-06	22	2020-231-22
7	2020-231-07	23	2020-231-23
8	2020-231-08	24	2020-231-24
9	2020-231-09	25	2020-231-25
10	2020-231-10	26	2020-231-26
11	2020-231-11	27	2023-498-27
12	2020-231-12	28	2020-231-28
13	2020-231-13	29	2020-231-29
14	2020-231-14	30	2020-231-30
15	2020-231-15	31	2020-231-31
16	2020-231-16		

The negative control primer pairs, **Production No. 2022-404-01**, 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: 2023 - 10 - 19

Approved by:

**Production Quality Control** 

omero

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For In Vitro Diagnostic Use MA099 v04 CoA\_DoC General IVDs Date: October 2023, Rev. No: 00 **ELERUP SSP** 

DQB1\*02

Certificates

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101.213-24 – including *Taq* polymerase 101.213-24u – without *Taq* polymerase

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

Lot No.: 8R0

Lot-specific information

**Declaration of Conformity** 

Product name:

Olerup SSP® DQB1\*02

**Product number:** 

101.213-24/24u

Lot number:

8R0

Intended use:

DQB1\*02 high 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

20231024

Date:

Quality Assurance