

101.903-24 – including *Taq* polymerase
 101.903-24u – without *Taq* polymerase

Visit www.caredx.com for
 “Instructions for Use” (IFU)

Lot No.: **7N7**

Lot-specific information

CERTIFICATE OF ANALYSIS

Olerup SSP® DQA1*02,05;DQB1*02,03:02

Product number: 101.903-24 – including *Taq* polymerase
 101.903-24u – without *Taq* polymerase
Lot number: 7N7
Expiry date: 2026-07-01
Number of tests: 24
Number of wells per test: 22+1

Well specifications:

Well No.	Production No.	Well No.	Production No.	Well No.	Production No.
1	2022-378-01	9	2022-378-09	17	2022-378-17
2	2022-378-02	10	2022-378-10	18	2022-378-18
3	2022-378-03	11	2022-378-11	19	2022-378-19
4	2022-378-04	12	2022-378-12	20	2022-378-20
5	2022-378-05	13	2022-378-13	21	2022-378-21
6	2022-378-06	14	2022-378-14	22	2022-378-22
7	2022-378-07	15	2022-378-15		
8	2022-378-08	16	2022-378-16		

The negative control primer pairs, **Production No. 2021-336-01**, can detect contamination with PCR products diluted 10⁻⁷.

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

Date of approval: 20220809

Approved by:



Production Quality Control



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Declaration of Conformity

Product name: Olerup SSP® DQA1*02,05;DQB1*02,03:02
Product number: 101.903-24/24u
Lot number: 7N7

Intended use: DQA1*02, DQA1*05, DQB1*02 and DQB1*03 medium resolution histocompatibility testing

Manufacturer: CareDx AB
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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: 20220812



Quality Assurance

