

101.704-48/12 – including *Taq* polymerase
 101.704-48u/12u – without *Taq* polymerase

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

Lot No.: **7R3**

Lot-specific information

CERTIFICATE OF ANALYSIS

Olerup SSP® DQ-DR SSP Combi Tray

Product number: 101.704-48/12 – including *Taq* pol.
 101.704-48u/12u – without *Taq* pol.
Lot number: 7R3
Expiry date: 2027-02-01
Number of tests: 48 tests – Product No. 101.704-48/48u
 12 tests – Product No. 101.704-12/12u
Number of wells per test: 46 + 1

Well specifications:

Well No.	Production No.	Well No.	Production No.
1	2023-491-01	9	2023-491-09
2	2023-491-02	10	2023-491-10
3	2023-491-03	11	2023-491-11
4	2023-491-04	12	2023-491-12
5	2023-491-05	13	2023-491-13
6	2023-491-06	14	2023-491-14
7	2023-491-07	15	2023-491-15
8	2023-491-08		

Well No.	Production No.	Well No.	Production No.
16	2022-438-01	32	2021-371-17
17	2022-438-02	33	2022-438-18
18	2022-438-03	34	2022-438-19
19	2022-438-04	35	2022-438-20
20	2022-438-05	36	2021-371-21
21	2022-438-06	37	2021-371-22
22	2022-438-07	38	2021-371-23
23	2022-438-08	39	2021-371-24
24	2021-371-09	40	2021-371-25
25	2022-438-10	41	2021-371-26
26	2022-438-11	42	2021-371-27
27	2022-438-12	43	2021-371-28
28	2022-438-13	44	2023-481-29
29	2022-438-14	45	2022-438-30
30	2022-438-15	46	2022-438-31
31	2022-438-16		

The negative control primer pairs, **Production No. 2022-404-01**, can detect contamination with PCR products diluted 10⁻⁷.



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For *In Vitro* Diagnostic Use
 MA100 v06 CoA_DoC IVD Annex II List B
 Date: March 2023, Rev. No: 00

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Lot-specific information

Results of Quality Control:

No false positive or false negative amplifications
obtained.

Date of approval: 2023-03-13

Approved by:



Production Quality Control



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Lot-specific information

Declaration of Conformity

Product name: Olerup SSP® DQ-DR SSP Combi Tray
Product number: 101.704-48/48u, -12/12u
Lot number: 7R3

Intended use: DQB1 and DRB1 low resolution histocompatibility testing

Manufacturer: CareDx AB
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 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 II List B, conformity assessed using Annex IV, as transposed into the national laws of the Member States of the European Union.

The Technical Documentation File is maintained at CareDx AB, Franzengatan 5, SE-112 51 Stockholm, Sweden.

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

Notified Body: TÜV Rheinland LGA products, Tillystrasse 2, D-90431 Nürnberg, Germany. (Notified Body number: 0197.)

Stockholm, Sweden

Date:

2023-04-27

Quality Assurance




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