101.624-12– including *Taq* polymerase, IFU-01
101.624-12u – without *Taq* polymerase, IFU-02

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

Lot No.: 6N5 **Lot-specific Information**

CERTIFICATE OF ANALYSIS

Olerup SSP® HLA-C*12

Product number: **101.624-12 – including *Taq* polymerase**
101.624-12u – without *Taq* polymerase

Lot number: **6N5**
Expiry date: **2025-11-01**
Number of tests: **12**
Number of wells per test: **47+1**

Well specifications:

Well No.	Production No.	Well No.	Production No.	Well No.	Production No.
1	2020-146-01	17	2020-146-17	33	2020-146-33
2	2020-146-02	18	2020-146-18	34	2020-146-34
3	2020-146-03	19	2020-146-19	35	2020-146-35
4	2020-146-04	20	2020-146-20	36	2020-146-36
5	2020-146-05	21	2020-146-21	37	2020-146-37
6	2020-146-06	22	2020-146-22	38	2020-146-38
7	2020-146-07	23	2020-146-23	39	2020-146-39
8	2020-146-08	24	2020-146-24	40	2020-146-40
9	2020-146-09	25	2020-146-25	41	2020-146-41
10	2020-146-10	26	2020-146-26	42	2020-146-42
11	2020-146-11	27	2020-146-27	43	2020-146-43
12	2020-146-12	28	2020-146-28	44	2020-146-44
13	2020-146-13	29	2020-146-29	45	2020-146-45
14	2020-146-14	30	2021-365-30	46	2020-146-46
15	2020-146-15	31	2020-146-31	47	2020-146-47
16	2020-146-16	32	2020-146-32		

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: 2022-01-12

Approved by:

Production Quality Control

CE

101.624-12– including *Taq* polymerase, IFU-01
101.624-12u – without *Taq* polymerase, IFU-02

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

Lot No.: 6N5 **Lot-specific Information**

Declaration of Conformity

Product name: *Olerup* SSP® HLA-C*12
Product number: 101.624-12/12u
Lot number: 6N5

Intended use: HLA-C*12 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
Date: 2022-01-17

Quality Assurance

CE