

101.612-12 – including Taq pol., IFU-01
 101.612-12u – without Taq pol., IFU-02

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

Lot No.: **1L9**

Lot-specific information

CERTIFICATE OF ANALYSIS

Olerup SSP® HLA-C*04

Product number: 101.612-12 – including Taq polymerase
 101.612-12u – without Taq polymerase
Lot number: 1L9
Expiry date: 2024-05-01
Number of tests: 12
Number of wells per test: 63+1

Well specifications:

Well No.	Production No.	Well No.	Production No.	Well No.	Production No.
1	2017-782-01	25	2017-782-25	49	2017-782-49
2	2017-782-02	26	2017-782-26	50	2017-782-50
3	2017-782-03	27	2017-782-27	51	2017-782-51
4	2017-782-04	28	2017-782-28	52	2017-782-52
5	2017-782-05	29	2017-782-29	53	2019-054-53
6	2017-782-06	30	2017-782-30	54	2019-054-54
7	2017-782-07	31	2017-782-31	55	2017-782-55
8	2017-782-08	32	2019-054-32	56	2017-782-56
9	2017-782-09	33	2017-782-33	57	2017-782-57
10	2017-782-10	34	2017-782-34	58	2017-782-58
11	2017-782-11	35	2019-054-35	59	2017-782-59
12	2017-782-12	36	2019-054-36	60	2017-782-60
13	2017-782-13	37	2017-782-37	61	2017-782-61
14	2017-782-14	38	2017-782-38	62	2020-200-62
15	2017-782-15	39	2017-782-39	63	2017-782-63
16	2019-054-16	40	2017-782-40		
17	2017-782-17	41	2017-782-41		
18	2017-782-18	42	2017-782-42		
19	2019-054-19	43	2017-782-43		
20	2019-054-20	44	2017-782-44		
21	2017-782-21	45	2017-782-45		
22	2017-782-22	46	2019-054-46		
23	2017-782-23	47	2017-782-47		
24	2017-782-24	48	2017-782-48		

The negative control primer pairs, **Production No. 2019-145-01**, can detect contamination with PCR products diluted 10⁻⁷.

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

Date of approval: 2020-07-10

Approved by: 

Production Quality Control



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Lot-specific information

Declaration of Conformity

Product name: Olerup SSP® HLA-C*04
Product number: 101.612-12/12u
Lot number: 1L9

Intended use: HLA-C*04 high resolution histocompatibility testing

Manufacturer: CareDx AB
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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:

2020-07-10

Quality Assurance


