

101.215-12 – including *Taq* pol., IFU-01
 101.215-12u – without *Taq* pol., IFU-02

Visit <https://labproducts.caredx.com> for
 “Instructions for Use” (IFU)

Lot No.: **2L9**

Lot-specific Information

CERTIFICATE OF ANALYSIS

Olerup SSP® DQB1*04

Product number: 101.215-12 – including *Taq* polymerase
 101.215-12u – without *Taq* polymerase
Lot number: 2L9
Expiry date: 2024-08-01
Number of tests: 12
Number of wells per test: 21+1

Well specifications:

Well No.	Production No.	Well No.	Production No.	Well No.	Production No.
1	2020-211-01	9	2020-211-09	17	2020-211-17
2	2020-211-02	10	2020-211-10	18	2020-211-18
3	2020-211-03	11	2020-211-11	19	2020-211-19
4	2020-211-04	12	2020-211-12	20	2020-211-20
5	2020-211-05	13	2020-211-13	21	2020-211-21
6	2020-211-06	14	2020-211-14		
7	2020-211-07	15	2020-211-15		
8	2020-211-08	16	2020-211-16		

The negative control primer pairs, **Production No. 2020-205-01**, can detect contamination with PCR products diluted 10⁻⁷.

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

Date of approval: 2020-09-16

Approved by:



Production Quality Control



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Lot No.: **2L9**

Lot-specific Information

Declaration of Conformity

Product name: *Olerup* SSP® DQB1*04
Product number: 101.215-12/12u
Lot number: 2L9

Intended use: DQB1*04 high 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, Franzégatan 5, SE-112 51 Stockholm, Sweden.

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

Stockholm, Sweden

Date: 2020-09-18



Quality Assurance

