

101.401-48/12 – including *Taq* polymerase  
101.401-48u/12u – without *Taq* polymeraseVisit [www.caredx.com](http://www.caredx.com) for  
“Instructions for Use” (IFU)Lot No.: **3R0**

Lot-specific information

**CERTIFICATE OF ANALYSIS****Olerup SSP<sup>®</sup> HLA-A low resolution**

**Product number:** 101.401-48/12 – including *Taq* polymerase  
101.401-48u/12u – without *Taq* polymerase

**Lot number:** 3R0

**Expiry date:** 2026-09-01

**Number of tests:** 48 tests – Product No. 101.401-48/48u  
12 tests – Product No. 101.401-12/12u

**Number of wells per test:** 31 + 1

**Well specifications:**

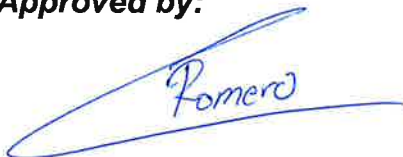
Well No.	Production No.	Well No.	Production No.
1	2021-369-01	17	2021-353-17
2	2021-353-02	18	2021-353-18
3	2021-369-03	19	2021-369-19
4	2021-353-04	20	2021-353-20
5	2021-353-05	21	2021-353-21
6	2022-442-06	22	2021-369-22
7	2021-353-07	23	2021-353-23
8	2021-353-08	24	2021-353-24
9	2021-353-09	25	2019-079-25
10	2022-442-10	26	2020-183-26
11	2021-353-11	27	2019-079-27
12	2021-353-12	28	2019-079-28
13	2021-353-13	29	2022-442-29
14	2021-353-14	30	2019-079-30
15	2021-353-15	31	2019-079-31
16	2022-442-16		

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

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

**Date of approval:** 2022-10-12

**Approved by:**



**Production Quality Control**

**0197**

For *In Vitro* Diagnostic Use  
MA100 v06 CoA\_DoC IVD Annex II List B  
Date: October 2022, Rev. No: 00

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Lot No.: **3R0**

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## Declaration of Conformity

**Product name:** Olerup SSP® HLA-A low resolution  
**Product number:** 101.401-48/48u, -12/12u  
**Lot number:** 3R0

**Intended use:** HLA-A low resolution histocompatibility testing

**Manufacturer:** CareDx AB  
Franzengatan 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 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: 20221013



Quality Assurance



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