101.123-24/06 – including *Taq* polymerase, IFU-01
101.123-24u/06u – without *Taq* polymerase, IFU-02

Visit https://labproducts.caredx.com for **"Instructions for Use" (IFU)**

Lot No.: 5F2 **Lot-specific information**

CERTIFICATE OF ANALYSIS

Olerup SSP® DRB5

Product number:	**101.123-24/06 – including *Taq* pol.** **101.123-24u/06u – without *Taq* pol.**
Lot number:	**5F2**
Expiry date:	**2021-10-01**
Number of tests:	**24 test – Product No. 101.123-24/24u** **6 tests – Product No. 101.123-06/06u**
Number of wells per test:	**18+1**

Well specifications:

Well No.	Production No.	Well No.	Production No.	Well No.	Production No.
1	2015-564-01	9	2017-833-09	17	2017-833-17
2	2015-564-02	10	2015-564-10	18	2017-833-18
3	2016-698-03	11	2016-698-11		
4	2015-564-04	12	2015-564-12		
5	2015-564-05	13	2016-698-13		
6	2015-564-06	14	2016-698-14		
7	2016-698-07	15	2015-564-15		
8	2016-698-08	16	2017-833-16		

The negative control primer pairs, **Production No. 2016-746-01**, can detect contamination with PCR products diluted 10^{-7}.

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

Date of approval: 2019-08-23

Approved by: Rebecka Ölund

Production Quality Control

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Date: August 2019, Rev. No: 01

101.123-24/06 – including *Taq* polymerase, IFU-01
101.123-24u/06u – without *Taq* polymerase, IFU-02

Visit https://labproducts.caredx.com for **"Instructions for Use" (IFU)**

Lot No.: 5F2

Lot-specific information

Declaration of Conformity

Product name:	*Olerup* SSP® DRB5
Product number:	101.123-24/06, -24u/06u
Lot number:	5F2
Intended use:	DRB5 high resolution histocompatibility testing
Manufacturer:	*Olerup* SSP AB Franzengatan 5 SE-112 51 Stockholm, Sweden ***Phone:*** +46-8-508 939 00 ***Fax:*** +46-8-717 88 18

We, *Olerup* SSP 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 *Olerup* SSP AB, Franzengatan 5, SE-112 51 Stockholm, Sweden.

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

Notified Body: Lloyd's Register Quality Assurance Limited, 1 Trinity Park, Bickenhill Lane, Birmingham B37 7ES, United Kingdom.
(Notified Body number: 0088.)

Stockholm, Sweden
Date: 2019-08-29

Quality Assurance

Changes in revision R01 compared to R00:

1. The expiry date has been altered due to extension of shelf-life.

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