

Certificates

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101.862-12 – including *Taq* **polymerase**, IFU-01 **101.862-12u** – without *Taq* polymerase, IFU-02

Visit www.olerup.com for "Instructions for Use" (IFU)

Lot No.: 0H8

Lot-specific information

CERTIFICATE OF ANALYSIS

Olerup SSP® HLA-C*04 Add-on

Product number:

101.862-12 - including Taq polymerase

101.862-12u - without Tag polymerase

Lot number:

0H8

Expiry date:

2021-04-01

Number of tests:

12

Number of wells per test:

4+1

Well specifications:

Well No.	Production No.
1	2015-477-01
2	2015-477-02
3	2015-477-03
4	2015-477-04

The negative control primer pairs, Production No. 2018-947-01, can detect contamination with PCR products diluted 10⁻⁷.

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

obtained.

Date of approval: 2018-11-09

Approved by: Rebecka Salme

Production Quality Control



Certificates

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101.862-12 – including *Taq* polymerase, IFU-01 **101.862-12** u – without *Taq* polymerase, IFU-02

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Lot No.: 0H8

Lot-specific information

Declaration of Conformity

Product name:

Olerup SSP® HLA-C*04 Add-on

Product number:

101.862-12/12u

Lot number:

0H8

Intended use:

HLA-C*04 histocompatibility testing

Manufacturer:

Olerup SSP AB

Franzengatan 5

20181113

SE-112 51 Stockholm, Sweden

Phone: +46-8-717 88 27 **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) ISO 9001:2008 and EN ISO 13485:2012, 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 *Olerup* SSP AB, Franzengatan 5, SE-112 51 Stockholm, Sweden.

Stockholm, Sweden

Emil Jonsson

Head of QA