



Product name:

AlloSeq Tx 17

Product code:

ASTX17.1(96)-A-RUO

Lot number:

23-1390

Expiry date:

2024-10-09

Number of tests:

96

Storage:

Box 1, 3 and 4:

-15°C to -25°C

Box 2:

Room temperature

Box 5:

2°C to 8°C

Kit contents:

Warnings and Precautions

This product contains DNA sequencing tests designed to detect variants of the tissue typing genes for research use only.

Product Insert: Room temperature

It is strongly recommended that these kits are validated by the user prior to implementation in the laboratory using

samples whose genotype has been determined by other molecular based procedures.

It is strongly recommended that the user follows all instructions provided via product labelling. Deviations from the described procedure are not recommended, may not be supported and could lead to typing errors. It is recommended that a positive control (human DNA) and negative/no template control (using sterile water in place of DNA) be included on every library preparation run. The positive control must produce a quantifiable library and the resultant sequence must be concordant with the sample's expected genotype. There must be no quantifiable library (measured by QuBit) in the negative template control for each experiment. If a quantifiable library is produced for the no template control the

run must be repeated.

Changes compared to the previous IFU:

IFU083 updated: Sections 1.1, 1.4, 1.6, 1.7, 1.9 and 1.10 were edited for clarity. Section 1.3 were modified to reflect verification studies performed by CareDx. Section 6 was modified to include thermal cycler parameters requirements for compatibility with Tx assay.

The following documents, and their translations, may be downloaded from https://www.caredx.com/lab-products/ or upon request from Manufacturer.

- IFU083 AlloSeg Tx Instructions for Use (Version 7.0, issued Jan 2023)
- IFU083-1_AlloSeq Tx Library Preparation Workbook RUO (Version 6.0, issued Apr 2022)
- IFU083-2_AlloSeq Tx Hybrid Capture Workbook RUO (Version 6.0, issued Apr 2022)
- IFU083-3_AlloSeq Tx Sequencing Workbook RUO (Version 4.0, issued Apr 2022)
- IFU083-5_AlloSeq Tx Early Pooling Workbook RUO (Version 9.1, issued Sep 2023)
- TEC478 AlloSeg Tx Safety Data Sheets (Version 4.2, issued Apr 2022)

ADDRESSES:

Manufacturer:

CareDx Pty Ltd,

20 Collie Street, Fremantle, WA, Australia, 6160.

Tel: +61-8-9336-4212 Email: orders-aus@caredx.com

Website: http://www.caredx.com

Distributed by:

CareDx AB,

Franzéngatan 5, SE-112 51 Stockholm, Sweden.

Tel: +46-8-508 939 00 Fax: +46-8-717 88 18

E-mail: orders-se@caredx.com Website: http://www.caredx.com/ CareDx Lab Solutions Inc.,

901 S. Bolmar St., Suite R, West Chester, PA 19382

Tel: 1-877-OLERUP1 Fax: 610-344-7989

Email: orders-us@caredx.com Website: http://www.caredx.com

October 2023 Version No: 11.0 For Research Use Only

2023-047



CareDx Pty Ltd 20 Collie Street Fremantle WA 6160 **AUSTRALIA**

CERTIFICATE OF ANALYSIS

CERTIFICATE NUMBER: 23-0047

MANUFACTURER AND TESTING LABORATORY

CareDx Pty Ltd 20 Collie Street Fremantle WA 6160 **AUSTRALIA**

PRODUCT INFORMATION

Product Name:

AlloSeq Tx 17

ACCEPTANCE CRITERIA

Kit size:

96

Batch number:

TEST

23-1390

Expiry date:

2024/10/09

RESULT

TESTING PROCEDURE

A panel of DNA samples (whose genotype was known) was used to assess the library preparation, enrichment and sequencing performance on a single kit from this batch against defined acceptance criteria

TEST RESULTS

| Note: Milot Sill Ellin | HESOLI |
|---|---|
| | |
| ≥ 41.36 ng/uL | PASS |
| ≥703 bp | PASS |
| | |
| ≥ 3 ng/uL | PASS |
| ≥ 675 bp | PASS |
| | |
| Correct genotypes obtained for all | PASS |
| ≥ 95% of loci tested with ≥ 87% reads with quality score of | PASS |
| Q30 or higher | |
| ≥ 95% of loci tested with ≥ 38% average | PASS |
| ≥ 95% of loci tested with ≥ 150x average | PASS |
| | ≥703 bp ≥ 3 ng/uL ≥ 675 bp Correct genotypes obtained for all ≥ 95% of loci tested with ≥ 87% reads with quality score of Q30 or higher ≥ 95% of loci tested with ≥ 38% average |

CONCLUSION

Compliance with acceptance criteria? YES NO

CERTIFICATION

I hereby certify that the testing was performed according to the company's standard operating procedures. I also certify that the results obtained were reviewed and released by myself prior to the issue of this certificate.

Signed: Lay

Name: Loric Longley

Position: OA Associate

Date of Issue: 25 0 ct 23