

PRODUCT UNDER TEST

Organization Name: [Healthland](#)

Address of Vendor: [1600 Utica Avenue South, Suite 300 Minneapolis MN 55416](#)

Test Product Name:

[Centriq](#)

Test Product Version-with-Release: [14](#)

TEST EVENT RESULT

Criteria Tested and Passed:

[b.1 - Transition of Care](#)

[b.2 - Clinical Information Reconciliation and Incorporation](#)

[b.5 - Common Clinical Data Set summary record – receive](#)

[f.1 - Xmit to Immunization Registries](#)

[f.2 - Xmit to Public Health Agencies – syndromic surveillance](#)

[f.3 - Xmit to Public Health Agencies – reportable laboratory tests](#)

[g.2 - Automated Measure Calculation](#)

[g.3 - Safety-enhanced Design](#)

[g.4 - Quality Management System](#)

[g.5 - Accessibility-Centered Design](#)

[g.6 - Consolidated CDA Creation Performance](#)

Additional Software Used in this Test Event:

[Updox 2016](#)

Clinical Quality Measures Tested in this Test Event:

Ambulatory:

[None](#)

Inpatient:

[None](#)

CERTIFICATION INFORMATION

Certifying ONC-ACB: [Drummond Group](#)

Certified Modules Inherited / Self-declare Without Testing Pending Final ONC-ACB Certification Evaluation and Decision:

- a.1 - CPOE-Medication
- a.2 - CPOE-Laboratory
- a.3 - CPOE-Imaging
- a.4 - Drug-drug, Drug-allergy Interaction Checks
- a.5 - Demographics
- a.6 - Problem List
- a.7 - Medication List
- a.8 - Medication Allergy List
- a.9 - Clinical Decision Support
- a.10 - Drug-Formulary Checks
- a.11 - Smoking Status
- a.12 - Family Health History
- a.13 - Patient-specific Education Resources
- a.14 - Implantable Device List
- a.15 - Social, Psychological, and Behavioral Data
- b.4 - Common Clinical Data Set summary record – create
- b.6 - Data Export
- d.1 - Authentication, Access Control, Authorization
- d.2 - Auditable Events and Tamper Resistant
- d.3 - Audit Report(s)
- d.4 - Amendments
- d.5 - Automatic Access Time-Out
- d.6 - Emergency Access
- d.7 - End-user Device Encryption
- d.8 - Integrity
- d.9 - Trusted connection

d.10 - Auditing actions on health information

f.7 - Xmit to Public Health Agencies – health care surveys

All Modules (Tested, Inherited, Self-declare) to be Evaluated for Certification by ONC-ACB:

a.1 - CPOE-Medication

a.2 - CPOE-Laboratory

a.3 - CPOE-Imaging

a.4 - Drug-drug, Drug-allergy Interaction Checks

a.5 - Demographics

a.6 - Problem List

a.7 - Medication List

a.8 - Medication Allergy List

a.9 - Clinical Decision Support

a.10 - Drug-Formulary Checks

a.11 - Smoking Status

a.12 - Family Health History

a.13 - Patient-specific Education Resources

a.14 - Implantable Device List

a.15 - Social, Psychological, and Behavioral Data

b.1 - Transition of Care

b.2 - Clinical Information Reconciliation and Incorporation

b.4 - Common Clinical Data Set summary record – create

b.5 - Common Clinical Data Set summary record – receive

b.6 - Data Export

d.1 - Authentication, Access Control, Authorization

d.2 - Auditable Events and Tamper Resistant

d.3 - Audit Report(s)

d.4 - Amendments

d.5 - Automatic Access Time-Out

- d.6 - Emergency Access
- d.7 - End-user Device Encryption
- d.8 - Integrity
- d.9 - Trusted connection
- d.10 - Auditing actions on health information
- f.1 - Xmit to Immunization Registries
- f.2 - Xmit to Public Health Agencies – syndromic surveillance
- f.3 - Xmit to Public Health Agencies – reportable laboratory tests
- f.7 - Xmit to Public Health Agencies – health care surveys
- g.2 - Automated Measure Calculation
- g.3 - Safety-enhanced Design
- g.4 - Quality Management System
- g.5 - Accessibility-Centered Design
- g.6 - Consolidated CDA Creation Performance

Additional Software Used:

Updox 2016, Medispan, Health Language Clinical Vocabulary Engine, Health Language HLI Web Services, Medline Plus Connect, Up To Date, Intellichart Patient Portal, Health Language Clinical Vocabulary Engine, Health Language HLI Web Services, Meinberg NTP

Clinical Quality Measures:

Ambulatory:
None

Inpatient:
None

APPROVED TEST SIGNATORY

Test Lab: Drummond Group EHR Test Lab

Personnel of Organization Leading Testing: [Earl Evans](#)

Location Where the Test Proctor Conducted the Testing (Remote/Onsite):
Remote

Test Report Serial Number: [EE-171206-2306](#)

Test Lab Approved Signatory printed name:
[Earl Evans](#)

Test Lab Approved Signatory signature: *Earl Evans*

Test Lab Approved Signatory title or function: [Test Proctor](#)

Date: [12/04/17](#), [12/05/17](#), [12/06/17](#)

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Overview of Test Report Document

The test report contains two core sections. The first section is the test result details on the first page of this report. This identifies the product under test, the modules successfully tested along, other test event details and the NVLAP approved signatory signature. The “Certification Status” section of that page is a non-normative section of the test report to provide some background on modules reported by the participating organization to inherit certification or qualify for gap certification pending approval from an ONC-ACB. This section does not imply certification, but it is a reference to note certified modules explicitly not tested.

The second section is a reference to the NVLAP accreditation of the Drummond EHR Testing Lab, a DG disclaimer of testing services and a reference to the testing process described in the Drummond Group EHR Testing Guide on the www.drummondgroup.com website.

Accreditation and Disclaimers

NVLAP Statement of Accreditation

For the scope of accreditation under [NVLAP Lab Code 200979-0](#), the Drummond Group EHR Test Lab fully conducted this test event in accordance to the approved quality procedures and within the scope of its accreditation.

This report must not be used by the customer to claim product certification, approval, or endorsement by ONC, NVLAP, NIST, or any agency of the Federal Government.



NVLAP Lab Code 200979-0

Drummond Group Disclaimer

Drummond Group Inc. (DG) conducts interoperability and conformance testing in a neutral test environment for various companies and organizations ("Customer"). The fact that the name of the Customer appears in the final report is not an endorsement of the Customer or its products or services, and DG therefore makes no warranties, either express or implied, regarding any facet of the business conducted by the Customer.

No warranty of the test product is implied over and above the publishing of the results of the as completed by the Customer during the specified time period of testing as described in this test report.

The Test Report shall not be reproduced except in full, without written approval of the laboratory.

Testing Process

For details on the test process, please refer to the Testing Guide on the Drummond Group website, or for further questions, please email EHR@drummondgroup.com.

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Change History

Date and description of changes made to original version of report are recorded here. The serial number of the test report which was modified is also recorded.

Date	Serial #	Description of Change
DRAFT1		Automated TRS Report

END OF DOCUMENT