Ankhos Oncology Software

Real World Testing Results Report CY 2023

General Information

Plan Report ID Number: 20221108ank Developer Name: Ankhos Oncology Software Product Name(s):Ankhos Version Number(s): 4 Certified Health IT Product List (CHPL) Product Number(s): 15.05.05.2959.ANKH.01.00.0.210111 Developer Real World Testing Plan Page URL: <u>https://www.ankhos.com/real-world-testing</u> Developer Real World Testing Results Report Page URL: <u>https://www.ankhos.com/real-world-testing</u>

Changes to Original Plan

No changes were made to the original test plan

Withdrawn Products

None

Summary of Testing Methods and Key Findings

Testing experience

There were multiple key findings from obtained while conducting Ankhos RWT. Results were obtained with a combination of queries, as well as visual validation of those queries. For some tests, manual document validation was required.

Challenges and lessons learned

Calendar year 2023 saw dramatic improvement in the number of valid CCDA files received. This is likely due to wider adoption as well as measures put in place to accommodate wider parsing of non-CCDA-compliant documents received.

There were few Direct messages that contained .rtf files or unsolicited HL7v2 content.

Even with these challenges in mind, we continue to demonstrate that users are actively using the CCDA import functionality as a consistent percentage of CCDA documents were being imported and integrated into the medical record.

One challenge remains: that CCDA documents are provided for patients who are not a part of the practice. In these cases the messages are archived as they are not able to be incorporated into the medical record.

Non-conformance

No non-conformities were identified during testing.

Standards Updates (SVAP and USCDI)

Standard (and version)	USCDI v1
Updated certification criteria and associated product	b2
Health IT Module CHPL ID	15.05.05.2959.ANKH.01.00.0.210111
Method used for standard update	Cures Update
Date of ONC ACB notification	11/21/2022
Date of customer notification (SVAP only)	N/A
Conformance measure	Measures B2.1, B2.2, B2.3
USCDI updated certification criteria (and USCDI version)	b2 - USCDI v1

Care Settings

All real-world data are observed from outpatient clinical oncology practices.

Metrics and Outcomes by measure

Below are utilization numbers by quarter for each metric. Conclusions and observations are listed below.

Metric	Q1 2023	Q2 2023	Q3 2023	Q4 2023
B2.1 CCDA	187	202	125	111
documents				
received				
B2.2 CCDA	100% (187)	100% (202)	100% (125)	100% (111)
documents				
successfully				
parsed				
B2.3 CCDA	95% (178)	89% (181)	91% (114)	87% (97)
documents				
imported				
C1.1 QRDA	None, sample	None, sample	None, sample	None, sample
documents	QRDA File	QRDA File	QRDA File	QRDA File
exported	Validated	Validated	Validated	Validated
F1.1	None, test	None, test	None, test	None, test
Immunization	Immunization case	Immunization case	Immunization case	Immunization case
cases reported	created	created	created	created
F4.1 Cancer	None, sample	None, sample	None, sample	None, sample
cases exported	Cancer case file	Cancer case file	Cancer case file	Cancer case file
	generated	generated	generated	generated

B2 observations and conclusions

Key Metrics:

- B2.1 CCDA documents received
- B2.2 CCDA documents successfully parsed
- B2.3 CCDA documents imported

Associated Criteria: B2

Relied upon software: None

Outcomes:

Customers are successfully receiving and incorporating CCDA R2.1 files via Direct messaging. In some cases, incoming Direct messages did not include CCDA files. Messaging including ONLY .RTF files, were not imported as they did not contain sufficient metadata to parse.

Further, in some cases, incoming files did not match new or existing patients and were archived.

No invalid CCDA files were encountered during RWT compilation.

Challenges: It was necessary to differentiate between Direct messages that did and did not contain CCDA files. All messages that did contain CCDA files were able to be parsed.

C1 Observations and Conclusions:

Key Metrics:

• C1.1 QRDA documents exported

Associated Criteria: C1

Relied upon software: None

Outcomes:

No practices manually exported QRDA files for the testing periods. Sample QRDA files were generated successfully.

Challenges: None

F1 Observations and Conclusions

Key Metrics:

• F1.1 Immunization cases reported

Associated criteria: F1

No practices submitted real-world immunization cases through Ankhos during the testing periods. A sample immunization case was created successfully.

F4 Observations and Conclusions

Key Metrics:

• F4.1 Cancer cases exported

Associated Criteria: F4

Relied upon software: None

Outcomes:

No practices submitted cancer case files during the testing periods. Test Cancer submission files were generated successfully.

Challenges: None

Key Milestones

Data were compiled for each calendar quarter in 2023. In all cases, the setting was in the context of outpatient medical oncology.