Sightview software

2024 Real World Test Results
My Vision Express 22.0

Revision History

The Revision History table below provides a record of all revisions made to this document throughout its life cycle. Updates are tracked by date the revisions were made, the version number, a brief description of the changes made and reason, as well as the name of the reviser and the approver.

Effective Date	Version #	Change Description/ Reason	Created/Revised by	Reviewed by	Approved by
01/08/2025	2024	2024 Report	Lora Woltz	Lora Woltz	Lora Woltz

Sightview software

Table of Content

Product Information

Product Information	
Plan Report ID Number: (ONC-ACB use only)	2024RWTP_MVEv22.0
Developer Name	My Vision Express, LLC
Product Name	My Vision Express
Version Number(s)	22.0
Certified Health IT	ONC Certification Criteria for Health IT
Product List (CHPL) ID(s)	15.02.04.2998.A057.22.02.1.221220
Developer Real World Testing Page URL	https://sightview.com/about-sightview/onc-certification/

Sightview software

Introduction

This report describes the steps and results of Real World Testing for the following Sightview Software, LLC CEHRT software platform for calendar year 2024

Platform	Version	Criterion to be Tested
My Vision Express	22.0	(b)(1), (b)(2)

Note: 170.315(b)(1) Electronic Health Information Export was certified 12/20/2023 and therefore was not included in the 2024 test plan and was not subject to testing during the 2024 calendar year.

Standards Updates and USCDI

Both required and voluntary standards updates must be addressed in the Real-World Testing plan, including SVAP (Standards Version Advancement Process) and USCDI (United States Core Data for Interoperability). Real World Testing plans must include all certified health IT updated to newer versions of standards prior to August 31 of the year in which the updates were made.

[ ] Yes, I have products certified with voluntary SVAP or USCDI standards.

[X] No, none of my products include these voluntary standards.

Standard (and version)	All Standards included in C-CDA R2.1
Updated certification criteria and associated product	None
Health IT Module CHPL ID	15.02.04.2998.A057.22.02.1.221220
Health IT Module Product ID	Not Applicable
Method used for standard update	Minimum Standard Code Sets
Date of ONC-ACB notification	Not Applicable
Date of customer notification (SVAP only)	Not Applicable
Conformance measure	**170.315(b)(1) Transitions of Care**
USCDI-updated certification criteria (and USCDI version)	Version 1

Standard (and version)	All Standards included in C-CDA R2.1
Updated certification criteria and associated product	None
Health IT Module CHPL ID	15.02.04.2998.A057.22.02.1.221220
Health IT Module Product ID	Not Applicable
Method used for standard update	Minimum Standard Code Sets
Date of ONC-ACB notification	Not Applicable
Date of customer notification (SVAP only)	Not Applicable
Conformance measure	**170.315(b)(2) Clinical Information Reconciliation and Incorporation**
USCDI-updated certification criteria (and USCDI version)	Version 1

Sightview software

Care Settings

All Sightview Software, LLC CEHRT platforms are considered to be an **Ambulatory Specialty Care Practice** type care setting for use in ophthalmology practices.

Changes to Plan

This section documents any deviations from the previously submitted 2024 real world test plan on file.

- No Deviations from the submitted 2024 test plan are noted

Summary of Test Method

This section describes the test methodology used to complete real world testing. Test documentation types are anticipated to include:

- Test Plans Document
- Test Case Documents
- Test Results Matrix Template
- Non-conformity logs

Every requirement must have a minimum of one test case. It is likely that each requirement will have more than one test case in order to fully evaluate possible workflow variations and outcomes. Actual results will be listed as Pass or Fail. A description of events will be included when a test case fails. Use of synthetic data will be declared, where appropriate.

To retain the privacy of the participating clients, a generic identifier will be used to reference any test result information that may be specific to a client.

Sightview software

Test Results

170.315(b)(1) Transitions of Care

Health IT Module:	My Vision Express	Version:	22.0
Regulation Text Citation	170.315(b)(1)	Criterion Description:	**Transitions of Care**
Date Range From:	12/01/2024	**Date Range To:**	12/31/2024
Test Case Description	Send: Software can export a CCDA to the intended recipient Receive: Software can receive the transition of care in electronic format		
Relied Upon Software?	Yes – Updox Yes - Regulatory Compliance Platform	Relied Upon Software Role	170.315(h)(2) 170.315(b)(1)

Data Analysis – Number of times a user successfully sent a CCDA electronically to the intended recipient.						
Test source	**Client**	**Test Source Description**	**Total # of Sent Attempted**	**Total # of Sent Successful**	**Criteria**	**Synthetic Data Used?**
HISP Report	3 Practices	Review of HISP generated Report for Sent CCDA across 3 clients	0	0	(i)(A)	No
Data Analysis – Number of CCDA received by the EHR						
Test source	**Client**	**Test Source Description**	**Total # of Received Attempted**	**Total # of Received Successful**	**Criteria**	**Synthetic Data Used?**
HISP Report	3 Practices	Review of HISP generated Report for Received CCDA across 3 clients	9	9	(i)(B)	No

Non-Conformities

Non-Conformity Description	Expected Results	Mitigation Strategy	Retest Date	Retest Result (Pass / Fail)
DM traffic was stalled between October 17th and October 28th	Successful passage of DM for sent and received CCDA	Issue resolved via restart of RCP module	Performance restored 10/28/2024	Pass

Notes:
Message failures were reviewed with the following causes documented:

Not Dispatched

Functionality is not adopted by My Vision Express clients

Tested by:	HISP/LW	Approved By	LW	Date:	01/13/2025

Sightview software

170.315(b)(2) Clinical Information Reconciliation and Incorporation

Health IT Module:	MDoffice	Version:	12.1
Regulation Text Citation	170.315(b)(2)	Criterion Description:	**Clinical Information Reconciliation and Incorporation**
Date Range From:	12/01/2024	**Date Range To:**	12/31/2024
Test Case Description	Software can properly match a received Transition of Care/ Referral Summary to the correct patient; User can review, validate and incorporate a patient's medication list, allergies and problem list; software can create a C-CDA document that includes the reconciled and incorporated data		
Date Tested	12/15/2024 - 01/27/2025	Reviewed By:	Lora Woltz
Relied Upon Software?	Yes – Updox	Relied Upon Software Role	170.315(h)(2)

Data Analysis – Number of times a user reconciled the medication list from the electronically received and incorporated CCDA.

It is permissible to create synthetic patient data to emulate the action of Clinical Information Incorporation and Reconciliation if there is not enough naturally occurring referral activity for the chosen RWT practice.

Test Source	Client	Source Description	Total # of received referrals	Successful Incorporation	Successful Reconciliation	Pass / Fail	Synthetic Data Used?
Clinical Recon Workflow	Practice 1/ 2 providers	Review of users DM dash for Incoming CCDA to reconcile	6	0 attempted by Practice / 3 under test	Yes	Pass	Yes
Clinical Recon Workflow	Practice 2 / 1 Providers	Review of users DM dash for Incoming CCDA to reconcile	2	0 attempted by practice / 1 under test	Yes	Pass	Yes
Clinical Recon Workflow	Practice 3 / 1 provider	Review of users DM dash for Incoming CCDA to reconcile	1	0 attempted by practice / 1 under test	Yes	Pass	Yes

Non-Conformities

Non-Conformity Description	Expected Results	Mitigation Strategy	Retest Date	Retest Result (Pass / Fail)
non found				

Notes: Historically low usage of Clinical Reconciliation functionality. Functionality performs as expected.

Tested by:	HISP/LW	Approved By	LW	Date:	01/13/2025