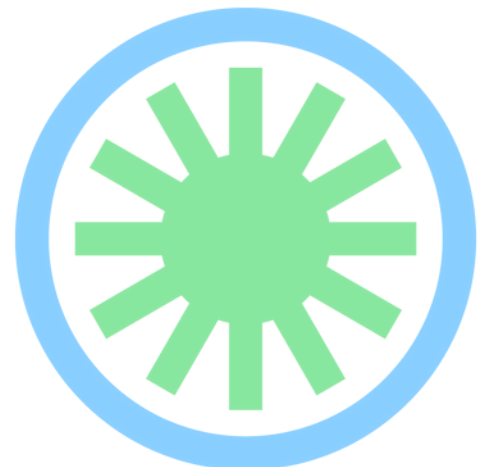




# **2024 Real World Test Results myCare Portal 4.0**



## 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.

<b>Effective Date</b>	<b>Version #</b>	<b>Change Description/ Reason</b>	<b>Created/Revised by</b>	<b>Reviewed by</b>	<b>Approved by</b>
01/08/2025	2024	2024 Report	<a href="#">Lora Woltz</a>	<a href="#">Lora Woltz</a>	<a href="#">Lora Woltz</a>
02/25/2025	2024	Revised to include num/denom metric	<a href="#">Lora Woltz</a>	<a href="#">Lora Woltz</a>	<a href="#">Lora Woltz</a>

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## Product Information

Product Information	
Plan Report ID Number: (ONC-ACB use only)	2024RWTP_MCPv4
Developer Name	Sightview EHR Holdings, LLC
Product Name	myCare Portal
Version Number(s)	4.0
Certified Health IT	ONC Certification Criteria for Health IT
Product List (CHPL) ID(s)	15.04.04.2998.myCa.04.02.1.221220
Developer Real World Testing Page URL	<a href="https://sightview.com/about-sightview/onc-certification/">https://sightview.com/about-sightview/onc-certification/</a>

## 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
myCare Portal	4.0	(e)(1), (g)(7), (g)(9)

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.

☒ No, none of my products include these voluntary standards

Standard (and version)	All Standards included in C-CDA R2.1
Health IT Module CHPL ID	15.04.04.2998.myCa.04.02.1.221220
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	<b>170.315(e)(1) View Download and Transmit</b>
USCDI-updated certification criteria (and USCDI version)	None

Standard (and version)	All Standards included in C-CDA R2.1
Updated certification criteria and associated product	None
Health IT Module CHPL ID	15.04.04.2998.myCa.04.02.1.221220
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	<b>170.315(g)(7) Application Access – Patient Selection</b>
USCDI-updated certification criteria (and USCDI version)	None

Standard (and version)	All Standards included in C-CDA R2.1
Updated certification criteria and associated product	None
Health IT Module CHPL ID	15.04.04.2998.myCa.04.02.1.221220
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	<b>170.315(g)(9) Application Access – All Data Request</b>
USCDI-updated certification criteria (and USCDI version)	None

## 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.

## Test Results

170.315(b)(1) Transitions of Care

Health IT Module:	myCare Portal	Version:	4.0
Regulation Text Citation	170.315(e)(1)	Criterion Description:	<b>View, Download and Transmit</b>
<b>Date Range From:</b>	01/01/2024	<b>Date Range To:</b>	12/31/2024
Test Case Description	CCDA is available in human readable format on the portal		
Relied Upon Software	Sightview Software EHR	Relied Upon Software Role	Provide Health Summary Data

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 CCDA to Portal	Total # of actual Viewed	Synthetic Data Used?
Global Portal Database	164 clients	Frequency of View Utilization	720752	8166	No

### Non-Conformities

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

Notes: Test report compiled for full CY 2024. Actual view rate is not in Sightview control as view action is voluntary to the patient. Performs as expected.

Tested by:	HISP/LW	Approved By	LW	Date:	01/13/2025
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## 170.315(b)(2) Clinical Information Reconciliation and Incorporation

Health IT Module:	myCare Portal	Version:	4.0
Regulation Text Citation	170.315(g)(7),(9)	Criterion Description:	Application Access
Date Range From:	01/01/2024	Date Range To:	12/31/2024
Test Case Description	Performance will pertain to token creation and the ability of the API to pull all category data for the chosen date or date range.		
Relied Upon Software	Sightview Software EHR	Relied Upon Software Role	Provide Health Summary Data

**Data Analysis – CEHRT is able to create an active token for a unique patient and then access the PHI data for all categories through the API.**  
**The API feature is not normally used in day-to-day practice. It is permissible to perform the workflow to initiate the API call for the purposes of demonstrating functionality for the use of RWT.**

Test source	Client	Test Source Description	Total # of API token calls (natural)	Total API calls tested (synthetic test data)	Total API calls successfully generated and ETT validated (synthetic test data)	Able to perform	Pass / Fail	Synthetic Data Used?
Review of System	N/A	Demonstrate ability to perform API token creation and All Data Request - EHR provided Test Patient data assigned portal credentials.	0	5	5	Yes – 5/5  Tested using synthetic patient data provided by EHR. Test UN/PW assigned to the demo patients. CCDA generated for each test instance and processed w/ ETT validator	Pass	yes

**Non-Conformities**

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

Notes:

API Functionality was not used by Sightview Software, LLC clients during the 2024 performance year.

Tested by:	Dev/LW	Approved By	LW	Date:	01/31/2025
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