2024 REAL WORLD TEST PLAN - MDOFFICE 12.1

MDoffice 12.1

Lora Woltz

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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

Product Information

Product Information	
Plan Report ID Number: (ONC-ACB use only)	2024RWTP_MDOv12.1
Developer Name	MDoffice, LLC
Product Name	MDoffice
Version Number(s)	12.1
Certified Health IT	ONC Certification Criteria for Health IT
Product List (CHPL) ID(s)	15.04.04.2998.Mdof.12.01.1.221230
Developer Real World Test Page URL	https://sightview.com/about-sightview/onc-certification/

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
MDoffice	12.1	(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.

170.315(b)(1) Transitions of Care	
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.Mdof.12.01.1.221230
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

170.315(b)(2) Clinical Information Reconciliation and Incorporation	
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.Mdof.12.01.1.221230
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

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.

Summary of Test Method

This section describes the test methodology that will be 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.