## **REAL WORLD TESTING PLAN**

#### **GENERAL INFORMATION**

Plan Report ID Number: [For ONC-Authorized Certification Body use only]

**Developer Name:** Nuance Communications Inc.

Product Name(s): Nuance Electronic Quality Measures

Version Number(s): 21.1

Certified Health IT Product List (CHPL) ID(s): 15.04.04.2071.Nuan.21.03.1.211104

#### **Developer Real World Testing Page URL:**

https://www.nuance.com/content/dam/nuance/en\_us/collateral/healthcare/data-sheet/ds-nuance-quality-measures-real-world-text-cy2021.pdf

#### JUSTIFICATION FOR REAL WORLD TESTING APPROACH

Nuance seeks to comply with CMS requirements related to Real World Testing (RWT). Details in the Federal Register can be found here: <a href="https://www.federalregister.gov/documents/2020/05/01/2020-07419/21st-century-cures-act-interoperability-information-blocking-and-the-onc-health-it-certification#p-3580">https://www.federalregister.gov/documents/2020/05/01/2020-07419/21st-century-cures-act-interoperability-information-blocking-and-the-onc-health-it-certification#p-3580</a>. This is required since Nuance's ECQM has been certified compliant with 2015 Edition criteria in § 170.315, specifically (c)(1) through (c)(3).

The Nuance Electronic Quality Measures system is a self-contained, hosted solution, that is accessible with a browser and Internet connection. Customers may optionally access the software directly since it has been designed to 1) receive QRDA-1 formatted data from the client through automated, secure transmissions, 2) process it through algorithms to determine outcomes, and 3) export the data for submission to CMS/TJC.

As part of RWT, we'll identify at least two customers for monitoring the entire process, and confirming outcomes and reports are consistent with their expectations.

# STANDARDS UPDATES (INCLUDING STANDARDS VERSION ADVANCEMENT PROCESS (SVAP) AND UNITED STATES CORE DATA FOR INTEROPERABILITY (USCDI))

Standard (and version)	ICD10CM, ICD10PCS, LOINC, RXNORM, SNOMEDCT	
	All are updated/maintained separately by standards bodies such as CMS	
Updated certification criteria	N/A	
and associated product		
Health IT Module CHPL ID	N/A	
Method used for standard	N/A	
update		
Date of ONC-ACB notification	N/A	

Date of customer notification	N/A
(SVAP only)	
Conformance measure	N/A
USCDI-updated certification	N/A
criteria (and USCDI version)	

## MEASURES USED IN OVERALL APPROACH

For each measurement/metric, describe the elements below:

- ✓ Description of the measurement/metric
- ✓ Associated certification criteria
- ✓ Justification for selected measurement/metric
- ✓ Care setting(s) that is addressed
- ✓ Expected outcomes

## DESCRIPTION OF MEASUREMENT/METRIC

Measurement/Metric	Description	
170.315(c)(1) Clinical quality	Ability to record clinical quality data, and to create the QRD1 export file for	
measures (CQMs) – record and	the measures selected by a provider for a selected time period.	
export		
170.315(c)(2) Clinical quality	Ability to import a QRDA 1 file for the measures selected by a provider for a	
measures (CQMs) — import	selected time period, and to properly calculate each measure outcome.	
and calculate		
170.315(c)(3) Clinical quality	Enable a user to electronically create a data file for transmission of clinical	
measures (CQMs) — report	quality measurement data in the QRDA I or QRDA III formats for the facility or	
	an individual patient, in accordance with published standards	

#### JUSTIFICATION FOR MEASUREMENT/METRIC

Measurement/Metric	Justification			
170.315(c)(1) Clinical quality	The justification of this metric is to be compliant with the ability for the cli			
measures (CQMs) – record and	to A) "record all of the data that would be necessary to calculate each CQM",			
export	and B) "to export a data file at any time the user chooses and without			
	subsequent developer assistance to operate". Data to be recorded must be in			
	QRDA1 format. The client will have the option to export, in QRDA1 format,			
	an individual patient file or a file for the quarterly submission period.			
170.315(c)(2) Clinical quality	The justification of this metric is to be compliant with the ability for the client			
measures (CQMs) — import	to A) "import a data file in accordance with the standard specified in			
and calculate - Import	§170.205(h)(2) for one or multiple patients", and B) "calculate each and every			
	clinical quality measure for which it is presented for certification." The data			
	may be imported through the application (manually) or through the secure,			
	automated feed of compliant QRDA1 files.			
170.315(c)(3) Clinical quality	The justification of this metric is to be compliant with the ability for the client			
measures (CQMs) — report	to "electronically create a data file for transmission of clinical quality			
(Cures)	measurement data". These files can be used for submission to CMS/TJC.			

## CARE SETTING(S)

Care Setting	Justification	
Acute Care Facilities	The data that is used to calculate the outcomes for the Electronic Clinical Quality Measures is received from the participating Acute Care Facilities. The Electronic Clinical Quality Measures pertain to patient encounters and need to have that final bill data sent to us in the QRDA format for processing.	

### **EXPECTED OUTCOMES**

Measurement/Metric	Expected Outcomes	
170.315(c)(1) Clinical quality	Provider will capture all data elements from the QRDA 1 file for the selected	
measures (CQMs) – record and	measure and be able to export the data.	
export - Export Feature		
170.315(c)(2) Clinical quality	Provider will load (import) a QRDA1 file into the application to verify the	
measures (CQMs) — import	format of the file and calculate results of the outcome for the measures that	
and calculate - Import	were contained in the file.	
170.315(c)(3) Clinical quality	The expected outcome is to create valid QRDA I and QRDA III files for	
measures (CQMs) — report	submission to CMS at the facility level or at the individual patient level.	

# SCHEDULE OF KEY MILESTONES

Key Milestone	Care Setting	Date/Timeframe
Update of Electronic Clinical Quality Measures to new 2021 specifications.	Acute Care Facilities	July 10,2021
Data Collection and Review	Acute Care Facilities	Jul 2021 – Feb 2022
Real World Test Plan creation and submission	Acute Care Facilities	November 15, 2021
Conduct Real World Testing	Acute Care Facilities	Jan - Feb, 2022
Submit Real World Testing results	Acute Care Facilities	Mar 2022

### **ATTESTATION**

This Real World Testing plan is complete with all required elements, including measures that address all certification criteria and care settings. All information in this plan is up to date and fully addresses the health IT developer's Real World Testing requirements.

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Date: Nov 12, 2021