



## A Trusted Quality Solutions Partner

As healthcare shifts to value-based care, new federal mandates linking electronic reporting to reimbursement have accelerated the demand for new solutions and services for gathering and reporting the data used to measure value and performance metrics.

Starting in 2016, Quality Measures are shifting from chart-abstracted to electronic clinical quality measures (eCQMs) using Quality Reporting Document Architecture (QRDA) Category 1 format.

### Other Modules Available

-  Quality Measures
-  Meaningful Use Compliance
-  Readmissions Management
-  Sepsis Management
-  Administration



## See a clear picture of your clinical quality measures to improve quality scores and performance metrics.

If you are like most MEDITECH hospitals, you are concerned about the CMS mandate for electronically reporting your quality measures using the new QRDA 1 format. Some of the concerns about moving toward electronic submission include:

- Federally mandated deadlines such as the Inpatient Quality Reporting (IQR) program that requires 4 eCQMs submission in 2016, and 8 in 2017.
- Capturing and mapping the new quality data required for QRDA 1.
- Following the MEDITECH best practices to be prepared for tracking the new clinical quality data. Without the right value set mapping, your QRDA 1 files will have missing data.

To address these mandates, Iatric Systems has created QRDA Assist™ a proven professional service that will quickly and accurately give your hospital the ability to generate and provide the QRDA 1 files from MEDITECH to Iatric Systems Analytics on Demand™, or your existing quality vendor.



## QRDA Assist provides your hospital with:

- **Technical Readiness Assessment:**  
Our experienced clinical experts will evaluate your organization's process and information systems for capturing the data for eCQM in alignment with TJC, CMS, and other Quality Improvement registry requirements using the MEDITECH best practices specifications.
- **Gap Analysis, Data Integrity, and Remediation:**  
Determine the current state of your clinical data and value sets and ensure nomenclature mapping is up to date. Assist with data validation and integrity prior to submission and testing.
- **Assist with eCQMs Selection:**  
Assist your hospital in the selection of the eCQMs for the IQR and Meaningful Use programs, and ensure that you're ready to track the required value sets for 2016 and 2017 reporting.
- **ARRA Report Manager Assistance:**  
Provide instruction for installation of MEDITECH ARRA Report Manager (requires MEDITECH DR) and guidance for populating required parameters. Will provide expert assistance with troubleshooting, correction of errors, and running the QRDA reports..
- **Assist with Validation and Testing in PSVA:**  
QRDA assist will validate and test your data with the QualityNet Pre-Submission Validation Application (PSVA) tool.

Since the introduction of ARRA and Meaningful Use requirements, more than 200 hospitals have partnered with Iatric Systems to satisfy the ever-changing requirements for quality measures reporting. The Iatric Systems Professional Services team will bring their deep experience of healthcare data, workflow, and performance improvement to help your organization deliver electronic clinical quality measures.

Be prepared to electronically track your clinical quality measures. Remember, 2016 IQR reporting is due by 2/28/2017, and 2017 tracking for 8 eCQMs starts on January 1, 2017. Contact us now about the QRDA Assist program to help you get a quick start towards QRDA 1 efficiency, and to be prepared for IQR and MU eCQM reporting.