OVERVIEW

MEDITECH Regulatory Reporting and Submission Services

Combines query, build, and optimization for reporting into a complete turn-key solution for your organization.

BENEFITS

- **Full-service regulatory submission services:** Includes query, build, monitoring, data validation, alerting/auditing, and trend analysis on all CMS major regulatory programs.
- **Deep experience:** Expertise with MEDITECH, regulatory data, and hospital operations, often resulting in increased quality measures year over year.
- **Proven success:** 100% success rate for on-time and complete submission services for clients.
- **Cost-effective:** Alternative to third-party reporting platforms by leveraging existing MEDITECH functionality.
- **Fills gaps left by other vendors:** Expert and complete services reduce the burden on in-house staff and fill gaps left by other reporting vendors.
- **Risk reduction:** Ensures system readiness and data flow for reports, minimizing the risk of penalties.

ADDITIONAL CAPABILITIES:

Enterprise-wide reporting:
Integration of additional systems and data sets in Microsoft PowerBI.

Vendor assistance:
Support with existing reporting vendor services to ensure complete submissions and data integrity.

Rise to New Regulatory Challenges with Simplification and Completeness

The most recent IPPS final rule introduces several new quality measures and reporting requirements that will impact hospital organizations. These changes raise the stakes for managing claims and disincentives higher each year, as well as the workload of clinical informaticists. New reporting requirements for new measures apply, as do disincentives hinging on new and/or validated data collection. As an organization, you must prepare for:

Understanding the rising stakes and avoiding financial disincentives

Submitting for new programs, additional measures, unique or more frequent submission periods

Ensuring system readiness for data flow for new and/or unique programs with specific requirements and reporting periods

Discover How CereCore Can Support Your Compliance Efforts

CereCore's comprehensive program with turn-key implementation, monitoring, validation, alerting/auditing, and trend analysis on all CMS and CDC major regulatory programs helps you adhere to the expectations for submission.

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Impact on Your Organization

SQL/NPR/RD Development: Receive reporting tools for each individual measure component per program, developed following certified workflows and functionality for attestation.

Reporting Presentation Layer (BCA/PowerBI/SSRS/Custom): Use industry-leading tools for ease-of-auditing and monitoring by site stakeholders.

HL7 Interface Development: Benefit from HL7 interfaces developed, implemented, validated, and monitored as required by both PI Public Health initiatives and other HL7 aspects of each regulatory program.

MEDITECH EHR Build: Receive MEDITECH-centric support for dictionary build, workflow build-out, education, and more.

On-going Support/SME Guidance: Receive guidance on current and future year regulatory expectations.

Submission Assistance: Expand your team's capacity by assigning data submittal to CereCore experts.

Core Measure Reporting: Direct third-party integrations and manual-based KART sampling, including SEP-1 – Chart Abstracted Measure and all chart abstracted measures.

Our Regulatory Reporting Success

100%
success rate: On-time and complete regulatory submission on behalf of our clients.

100%
renewal rate: Year over year for regulatory submission services.

Improved quality measures: Enhanced client quality measure scores after CereCore feedback addressing deficiencies in measure numerators or data flow.

Powerful Reporting and Submission Options

Microsoft SQL:

- Leverages MEDITECH and DR/SQL expertise for application configuration, QRDA curation, data integration, query build, dashboards, and more.
- Requires no additional software by utilizing data in the MEDITECH DR and the Microsoft SQL licenses you likely already own.

Microsoft PowerBI:

- Well-suited for comparing large quantities of data from disparate sources (e.g., MEDITECH alongside an external materials system).
- Requires no additional software by utilizing data in the MEDITECH DR and commonly owned Microsoft PowerBI licenses.

Watch the Video: How Regulatory Submissions Work at CereCore

Explore the chart: 2025+ Regulatory Programs

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ABOUT US

CereCore® provides IT services that make it easier for you to focus on supporting hospital operations and transforming healthcare through technology. With a heritage rooted in our nation's top-performing hospitals, we serve as leaders and experts in technology, operations, data security, and clinical applications. We partner with clients to become an extension of the team through comprehensive IT and application support, technical professional and managed services, IT advisory services, and EHR consulting, because we know firsthand the power that integrated technology has on patient care and communities.

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