

# **Regulatory Compliance**



Pre-Survey Prep | Post-Survey Response | Quality Improvement & Turn-Around

# **Resources to Navigate Regulatory Change**

The challenges facing the post-acute care leaders continue to escalate at a significant pace. The experts at Pathway Health are here to assist your facility develop new operational systems, or better utilize current processes, and provide an unbiased review of compliance performance for audit and survey preparedness.

# **Preparation is Key**

Pathway Health is your partner in readiness and success. We have a full complement of regulatory services and education available (virtual and on-site) that can be customized to meet your specific need.

Whether you need support in a focus area (infection prevention and control, abuse prevention, etc.) or a full Mock Survey, we are here to support your organization!



**Regulatory Review** - Mock Survey, review current regulatory status.



Targeted Regulatory Reviews -Customized to your needs. Single, Four and Five Focus Area options



Clinical Regulatory Reviews -Clinical documentation scan and Trend Snapshot to determine areas for improvement.



Regulatory Support Services (On-site or Off-site) - Regulatory Response Services; On-site Regulatory follow-up; POC/DPOC; virtual coaching; customized staff training.



**Customized Resources & Tools** - Assistance in customizing the policies, training, and tools for your organization.



**Staff Education & Training** - Customized training and, virtual, on-demand and on-site education to support the facility's improvement process.

Apply our expertise to your regulatory strategy. Contact us today.



# **Services Overview**



### **On-site Regulatory Review**

Mock Survey - Completed using CMS survey tools and following the federally-mandated survey process. A verbal exit report will be provided along with notes about initial resident-specific findings. A written report will follow containing the consultant's detailed findings and recommendations for improvements.

#### **On-site Targeted Regulatory Reviews**

A qualified consultant to complete a focused regulatory review using the Centers for Medicare and Medicaid Services (CMS) Critical Element Pathway Survey Tools and other CMS survey tools. Targeted Regulatory Reviews include:

- A verbal exit report, followed by a written report detailing the consultant's findings and a recommended Action Plan.
- One QuickPATH related to a focus area

Single Focus Area Includes: A sample

of records for five residents selected by

the customer; Policies and procedures

associated with the topic; Interviews of

residents and staff about the topic; and

Environmental observations related to

topic; Observation of staff practices

**Targeted Regulatory Review:** 

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Four Focus Areas (two-days on-site) Includes: A sample of records for five residents selected by the customer; and Competent Staffing; Incident and Accident Investigation Process; and one additional mandatory survey task or clinical topic review, as selected by the customer.

• 10% discount on Pathway Health products and tools (AAPACN®, Frailty, INTERACT®, LEAD and Restorative training are excluded)

#### **Targeted Regulatory Review:**

Five Focus Areas (three-days on-site) Includes: A sample of records for five residents selected by the customer; Infection Prevention and Control Program; Sufficient and Competent Staffing; Incident and Accident Investigation Process: Dining Observations; and one additional mandatory survey task or clinical topic review, as selected by the customer.

# **Off-site Regulatory Reviews**

A qualified consultant will complete an off-site review of the electronic clinical records. Includes a written report detailing findings and a recommended Action Plan. In addition, a follow-up call to review the report, answer questions and provide initial support for implementing the Action Plan. The electronic health record must contain the described documents for these reviews.

Regulatory Clinical Documentation Scan - Sample of residents from the Facility Survey Matrix (provided by the facility) for a clinical documentation review, focusing on regulatory vulnerabilities in the documentation. Includes:

- Assessments
- MDS

the topic.

- Care Plans
- Physician Orders
- Progress Notes
- Medical Diagnosis List

Regulatory Trend SnapShot – Focus on regulatory vulnerabilities in the clinical documentation related to publicly reported data. Includes:

- Health Inspection History
- Quality Measures
- Re-admissions

## Regulatory Support Services (On-site or Off-site)

#### Regulatory Response Services include the following:

Support for IDR/IIDR

- Draft Plan of Correction

Survey Recovery

- Immediate Jeopardy Removal
- POC Implementation

Directed POC

On-site Regulatory Follow-Up – A Pathway Health consultant re-evaluates progress on Action Plan implementation and reduction of regulatory vulnerabilities. The number of onsite days and focus will be established prior to the visit.

Virtual Coach for Action Plan Implementation - Web-based meeting with facility leaders to support implementation of the recommended Action Plan. An expert consultant will guide staff through the development of new or revised processes and offer suggestions and recommendations for putting the recommendations into action. Web-based meetings can be scheduled at the customer's desired frequency and schedule.

Keep your staff on the path to quality care and compliance. Contact Pathway Health.