Our
Focus is
Your
Perfection



PHARMACOVIGILANCE RISK MANAGEMENT PLUS



Our Mission at PVRM+

Our mission at PVRM+ is to provide niche value-added Medical Communications Services to:

- Ensure Regulatory Compliance
- Enhance Efficiency and Productivity
- Reduce Costs Significantly.

We have a proven delivery model that enables companies to focus on their core business objectives.



Our Focus is You

North/South America Europe Middle East/Asia **USA** India Global Located in Hyderabad Headquarters in • Center of Excellence **New Jersey** • Executive for Delivery & Oversight Operations Project/Account Auditing Services • SME Consulting Management Our focus is your perfection...



Our Partnership Model

Domain Knowledge Expertise

Proven Delivery Model



Comprehensive & Flexible Solutions

Quality
Management
Systems Approach

Compliance to Regulatory Requirements



Our Comprehensive and Flexible Service Offerings

Medical Information Management

- Manage Medical Information Call Center
- Prepare New Global Standard Responses
- Maintain Global Standard Responses
- Provide Medical Information Staffing at Congresses

Medical Affairs Content Development

- Develop Disease State & Product Sales Training Materials
- Write Patient Education Content
- Create Congress Posters& Presentations
- Maintain CCDS & Product Monographs
- Write & Update REMS & Registries Materials

Medico-Marketing Content Creation

- Create Professional Materials for Sales Team
- Develop Content for HCP/KOL Presentations
- Prepare Materials for Display at Congresses
- Write Patient Product & Disease State Brochures
- Provide Patient Support Program Management
- Create Website Content

Clinical Writing Services

- Author & Maintain:
 - Protocols
 - ➤ IBs
 - ➤ IMPDs
- Write Clinical Study Reports, including Patient Narratives
- Prepare Clinical Evaluation Reports
- Develop CT Registry Summaries



Our Comprehensive and Flexible Compliance Service Offerings

QMS Management

- Write & Maintain Policies and SOPs
- Utilize Enterprise Learning Method for Training
- Implement and Manage CAPA System
- Conduct Root Cause Analysis

AE & PQC Compliance

- Capture of Spontaneous Adverse Event Reports
 Product Quality
 Complaints per
 Regulations
- Create & Maintain Safety Data Exchange Agreements
- Perform Monthly Reconciliations

Promotional Copy Compliance

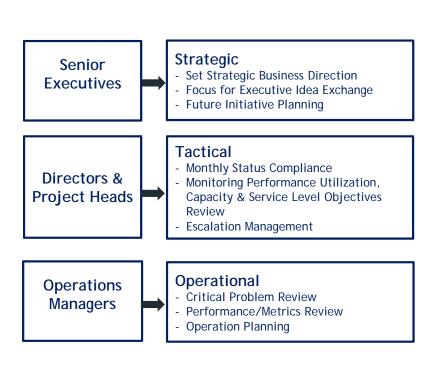
- Conduct Medical Review of Promotional Content
- Perform Referencing and Annotations
- Maintain Product References Master Log
- Attend Client Promotional Content Review Meetings

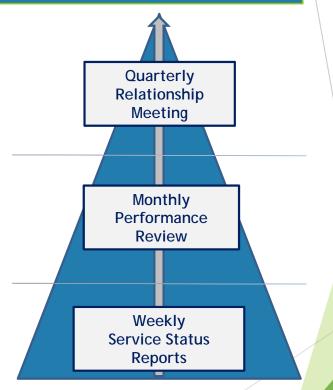
Audit and Inspection Support

- Conduct Internal Audits
- Perform Due Diligence External Audits
- Prepare Client Staff for Facing Audit and Regulatory Inspection
- Support Client Staff During External Audit and Regulatory Inspection



Our Proactive Project Governance





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Thank You!



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Contact Us At: kheston@pvrmplus.com