Customized solution for effective patient safety monitoring.

Pharmnova is committed for holistic approach in all services covering entire product Life Cycle through various customized models based on the needs of Pharmaceutical organizations to safeguard patient's innate health while using client's products.

Flexible business model

As a full service provider, ad-hoc functional consultancy or resource support.

Dynamic changes in regulatory landscape require continuous monitoring across regions and implementation of changes effectively in the existing system to meet the latest regulatory requirements. At Pharmnova, we continuously monitor the latest regulations and guidelines, help our clients in establishment of effective Pharmacovigilance system.

Who We Are

ABOUT US

Pharmnova is a well-organized, innovative and knowledge driven contract research organization established by medical doctors, pharmacists, and other allied healthcare professionals.

Our experience in various business model like pharmaceutical organization, CRO, BPO and KPO brings in confidence and unique sustainable business model for our clients in maintaining seamless and robust pharmacovigilance system.

Along with our partners, we can provide services in various regions.

OUR SERVICES

Pharmacovigilance
Medical Writing
Medical Affairs
Quality Assurance
Medical Information
Consulting

CONTACT US

Info@Pharmnova.Com

+91 9870054879

www.pharmnova.com

Pharmnova Medical Research Pvt Ltd
Plot No 12, 6 George Avenue
Rajeswari Nagar, Chennai 600073

Pharmnova

Enhancing patient safety

Why Pharmnova

- ✓ Global expertise
- ✓ Proactive and integrated approach
- ✓ Partnership strategy and practices
- ✓ Data security and protection
- ✓ Cost effective

Certification

ISO 9001: 2015 Certified by IRQS

NABCB QM 006

"Quality is never an accident. It is always the result of intelligent effort."

- John Ruskin

Info@Pharmnova.Com
+91 9870054879
www.pharmnova.com

Contact us to know more about

Our Services

Literature surveillance - *Global Systematic Literature Surveillance managed by comprehensive review approach for case processing, aggregate analysis, risk and signal management, and by cutting down the cost and time for generating literature- origin safety information.*

ICSR management - *Automated and validated safety application enables optimized solution with an integrated cost-effective approach for case processing, signal management and literature review.*

Signal management - *Signal related decision-making by customized approach according to the product portfolio and market status together with integrated approach in safety application*

Risk management - *Our expertise facilitates swift decisions on risk minimization and communication to stakeholders promptly. We can also help our clients in representing REMS program and in establishing additional risk minimization activities according to RMP.*

Qualified Person for Pharmacovigilance - *With our network of experts in EU, UK and ROW countries, we support our clients in QPPV and local safety contact (local QP) requirements by minimizing operation cost.*

PSMF and XEVMPD services - *Establishment and maintenance of PSMF and XEVMPD information becomes seamless through our experienced team.*

SDEA management - *Our collaborative approach enhances our clients' relationship with their partners and distributors through representation of SDEA related activities.*

Aggregate report - *Our expertise in preparing all formats of aggregate reports boosts your confidence and enables cumulative analysis of product safety.*

Medical information and Call Centre - *24/7 support of medical information specialists in all time zones.*

Audit – *Our enriched knowledge on regulator's expectations and audit conduct mitigates the lacunae in the system and prepares the system for inspection readiness.*

PHARMACOVIGILANCE

- ☑ ICSR management
- ☑ Safety regulatory reporting and regulatory intelligence
- ☑ Risk management
- ☑ Pharmacoepidemiology
- ☑ Signal management
- ☑ Literature surveillance
- ☑ SDEA management
- ☑ Qualified Person for Pharmacovigilance (QPPV)
- ☑ PSMF preparation and maintenance

MEDICAL WRITING

- ☑ Aggregate reports
- ☑ Regulatory affairs documents
- ☑ Study documents
- ☑ Labeling and reference safety information (RSI)

MEDICAL AFFAIRS

- ☑ Medical monitoring in clinical trial
- ☑ Health hazard and risk assessment
- ☑ Clinical trial safety assessment
- ☑ Clinical data review

MEDICAL INFORMATION

- ☑ Medical inquiries, Product quality complaint and adverse event handling
- ☑ 24/7 Call Centre support

QUALITY ASSURANCE

- ☑ Due diligence audit
- ☑ Audit and inspection support

Pharmnova

Enhancing patient safety