Driving Safety and Compliance Seamlessly

IQVIA's unique FSP synergy of pharmacovigilance and regulatory affairs

At IQVIA, pharmacovigilance and regulatory affairs work together to ensure the safety, efficacy, and compliance of medications. This collaboration offers significant benefits by streamlining processes, enhancing drug safety, and protecting public health.

IQVIA pharmacovigilance

- **Analyze adverse drug reaction** (ADR) data from multiple sources
- **Detect safety concerns** to minimize risks for patients
- Continuously **assess the risk-benefit** profile of medications, ensuring public safety

IQVIA regulatory affairs

- **Develop and submit regulatory filings** for product approvals
- Maintain documentation to **ensure compliance with standards**
- **Oversee post-marketing activities**, including labeling updates & safety reporting

Integrated pharmacovigilance and regulatory affairs

- Advanced analytics
- Global safety database
- E2E risk management
- 24/7 monitoring
- Accelerated approvals
- Regulatory intelligence
- Global regulatory network
- Seamless lifecycle management

IQVIA integrated pharmacovigilance and regulatory affairs

Best practices

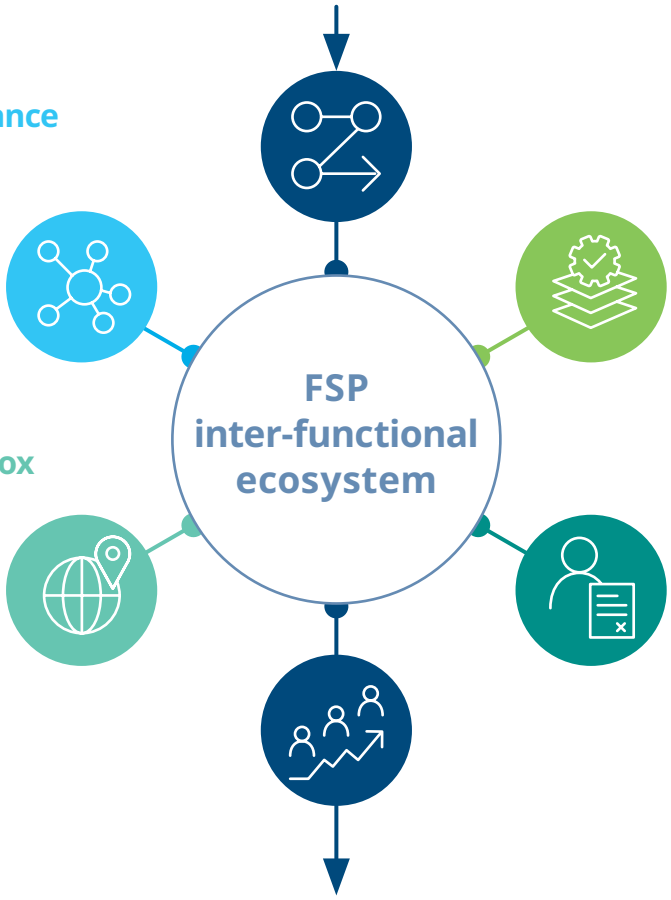
- Optimize cost, quality and efficiency across portfolio
- Continuous improvement
- Ongoing change management

Cross-functional governance

- One team culture
- Common goals and vision
- Holistic KPI review
- Knowledge retention

IQVIA FSP Country in a Box

- Rapid market entry
- Expert regulatory compliance support
- Scalable resources
- Localized expertise
- Effective cost management



IQVIA's coordination team

- 1 dedicated project manager
- Systematized collaboration
- Aligned tech investments for less system and data friction

Vendor consolidation

- Unified vendor management

Outcomes

Optimized budget efficiency

Reduced redundancies, lower vendor management costs, compliance expenses minimized

Unmatched quality and compliance

Continuous safety monitoring, regulatory expertise, risk management ensured

Faster risk detection and regulatory action

Real-time insights, seamless submissions, faster regulatory decisions

To discover more about how smoothly our pharmacovigilance and regulatory affairs experts join forces to offer you the best IQVIA experience, contact us: globalfspgtm@iqvia.com

IQVIA

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