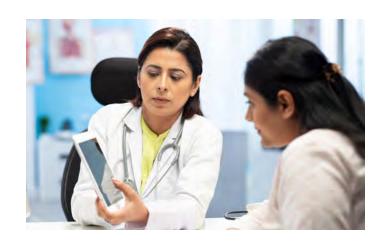


End-to-End Safety Solutions

Lifecycle safety expertise enhanced by AI/ML

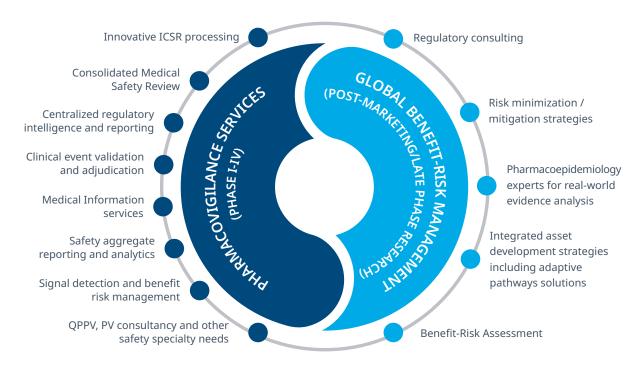
The challenges

In life sciences, product safety is an absolute imperative. As the importance of safety has increased, so too has the complexity of delivering safer products. In addition, the proliferation of new data sources is adding ever-increasing volume, workflows and cost.



Technology-enabled solutions across the product lifecycle

IQVIA can help. As one of the world's largest and most experienced safety and pharmacovigilance (PV) organizations, IQVIA brings extensive domain expertise and deep regulatory intelligence to every engagement. Using IQVIA Connected Intelligence™, our flexible, global delivery models employ leading-edge technology platforms, enabling predictable, quality delivery across complex multi-year programs. From early development to post-approval and beyond, let us help you increase safety, accuracy and compliance, while decreasing complexity and cost. Our services include:



IQVIA's safety teams are harnessing the power of automation, artificial intelligence (AI) and machine learning (ML) to deliver streamlined safety and surveillance processes including our award-winning Vigilance Platform.

VIGILANCE PLATFORM

Comprehensive Saas Safety/PV solution employing automation and Al to simplify PV processes and streamline operations

SAFETY MONITORING SERVICES WORKBENCH

Harmonizing a fragmented safety processing environment through proactive workflow management

ROBOTIC PROCESS AUTOMATION (RPA)

Workflow automation including data entry, QC, narratives, case migration and document comparison

QUALITY CONTROL DATABASE

Providing real-time case quality visualizations, validated trend analysis and automatic notifications

REGULATORY INTELLIGENCE DATABASE

Enabling regulatory compliance with access to a single source for comprehensive requirements

IQVIA safety – 2022 delivering excellence



Case ID & intake, case processing

>2M ICSRs processed 99.83% TAT compliance 99.05% end of line quality



Benefit risk management

9 projects supported 70 risk management plans



Audits and inspections

157 audits/inspections supported 80 customer audits **77** inspections



Aggregate reporting

>1.3K aggregate safety reports across 131 clients 99.89% Aggregate report compliance 99.44% Aggregate reports end of line quality



PV platform hosting and support

Hosted and managed 389 safety database enterprises 52 IRMS divisions 15K helpdesk tickets closed



Medical safety advisors

155 SOP updates

769K medical review of ICSRs **107** signal runs, **1K** PBRERs 2K DSURs, 132 RMPs, and 2K PADERs 98% TAT compliance 99.5% medical peer review quality



QPPV and **PV** agreements services

30 countries supported with **QPPV** requirements

18 PSMF updates with 96% compliance 106 PVA updates

5 PVA svs provided to 3 top pharma with 98.12% compliance



Medical Information

31 medical Information projects 8 AE intake/LAPs projects

147K inquires, 267K adverse events, 11K product complaints

99.97% AE/PQC compliance, 16 second average call wait time



Literature screening

95 literature customers with 1,500 products

>425K biomedical literature abstracts reviewed

>27K full text articles ordered

>99% compliance EoL quality >99% literature absract compliance



Regulatory reporting and compliance

1.4M reg submissions 1.7M investigator alert letter

submissions delivered 99.65% overall EC/RA submission compliance



Signal management

1,026 signal detection runs performed for 419 distinct products 85 signal validations and **63** signal evaluation reports authored covering 38 products 100% compliance



World-class cross functional teams deliver comprehensive patient safety solutions across the product life cycle



Better patient outcomes across product lifecycle



customers



>1.400 projects