

e-RegMonitor

eRegMonitor™ from IQVIA is the most complete, centralized repository of summarized, on- demand regulatory information affecting the life sciences industry. Powered by a team of dedicated, full-time research attorneys, eRegMonitor provides 24/7 web access and proactive e-mail notifications about critical legislative and regulatory issues in key Life Science areas.



COMPREHENSIVE

- Centralized repository of key life science-specific legislative and regulatory research
- 24/7 web access to high level summaries as well as detailed, citation-specific insight
- Full time, dedicated team of life sciences research attorneys

PROACTIVE

- Track pending laws throughout the legislative process
- Receive proactive notifications via e-mail and home page announcements when legislation is enacted

INTERACTIVE

- Community-based portal allows you to interact directly with our compliance experts
- Extensive keyword search capabilities allow you to easily find the info you need

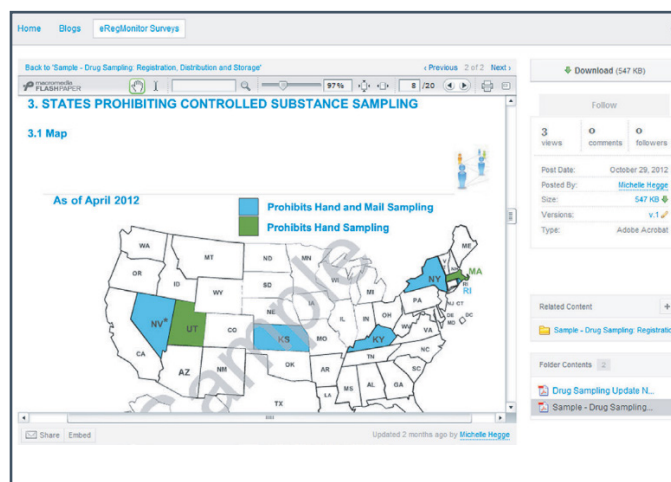
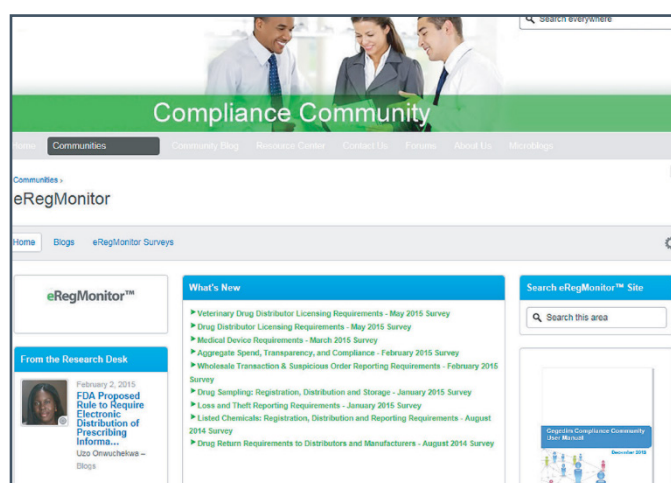
e-Regmonitor's user-friendly, web-based access makes it easy to share knowledge and encourages more informed decision making at every level of your compliance organization.

E-REGMONITOR INCLUDES 16 COMPREHENSIVE LIFE SCIENCE REGULATORY SURVEYS

- Advertising Restrictions
- Aggregate Spend, Transparency, and Compliance
- Drug Distributor Licensing Requirements
- Drug Quality Security Act Requirements and State Licensing
- Drug Return Requirements to Distributors and Manufacturers
- Drug Sampling: Registration, Distribution, and Storage
- Listed Chemicals: Registration, Distribution, and Reporting Requirements
- Loss and Theft Reporting Requirements
- Medical Devices Licensing Requirements by State
- Mid-level Practitioners: Sampling, Prescriptive, and Dispensing Authority
- Over-the-Counter Licensing Requirements Chart
- Prescription Drug Affordability Programs Survey
- Preprinted Prescription Pads: Content, Construction, and Usage
- Veterinary Drug Distributor Licensing Requirements
- Virtual Distributor and Virtual Manufacturer Licensing Requirements Survey
- Wholesale Transaction and Suspicious Order Reporting Requirements

WHAT MAKES US UNIQUE

Leveraging IQVIA's depth of knowledge in regulatory compliance enables you to make informed decisions and develop proactive strategies regarding the impact of current and future federal and state legislative and regulatory requirements on your operations - the key to achieving maximum efficiency and cost-effectiveness.



CONTACT US

1025 Boulders Parkway, Suite 301
Richmond, VA 23225
iqvia.com/contactus

